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## Utilisation of Off-River Habitats

## by Lowland River Fishes

## Carolyn M. Knight

Thesis submitted for the degree of Doctor of Philosophy

University of Durham School of Biological and Biomedical Sciences

May 2006


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## Abstract

This study assessed the importance for fish of lateral connectivity of river side-channels to the River Frome, a lowland chalk stream in Southern England. Lateral connectivity and habitat complexity is under threat in many floodplain river systems due to human disturbance. A holistic approach was used to investigate fish communities in seven sidechannels (including drainage ditches, natural streams and a millstream) and to assess the functionality provided by these habitats to fish species. Seasonal electric fishing over three years was used to monitor fish assemblages in each channel in relation to biotic and abiotic variables. Fish movements between the main channel and lateral habitats were monitored continuously in five locations with PIT (passive integrated transponder) telemetry. In total over one hundred pike (Esox lucius) and dace (Leuciscus leuciscus) were radio tracked to monitor the movements of individual fish with increased sensitivity.

Each side-channel provided a distinct habitat and supported a different fish community. Flow was the main discriminating factor between channels and their assemblages. General linear models of biotic and abiotic variables did not predict abundances of individual species effectively. In contrast, habitat stability was a good indicator of species diversity and may prove a useful management tool. Side-channels were used for different functions by different species. A single fish species used a range of habitats within the river system, each for different functions with functional differences in use between seasons.

The River Frome dace population exhibited a structured population with individuals exhibiting different uses of the main river and side-channels. The structure of the population is based upon the availability of both lateral and longitudinal connectivity, with dace moving between the main river and side-channels and also making excursions over 10 km along the main river channel. Drainage ditches are a particularly important pike spawning location, with males arriving earlier onto this distinct habitat. Protandry (early arrival strategy of some males) was exhibited by slow-growing male pike which arrived as early as December. Growth was also related to home range size in males, with faster growing males inhabiting larger, home ranges.

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## Chapter 1

## General Introduction

# 1.1 River Floodplain Habitats: Functionality of a Declining Fish Habitat 

Some of the first conceptual models of the principles of river systems focussed primarily on longitudinal (main river channel) processes and functions (Chorley 1962, Cummins 1974, Huet 1954, Leopold and Maddock 1953, Vannote et al. 1980). Subsequent work soon showed that focussing only on the longitudinal main river corridor was too limited for understanding the ecology of lowland freshwater fishes and that the ecological integrity of floodplain rivers depends on the ecological connectivity acting across their floodplains (Amoros and Roux 1988, Ward and Stanford 1995). In the four-dimensional concept of running water systems (Ward 1989) the four dimensions are the longitudinal dimension, latitudinal dimension, the vertical dimension consisting of interactions between the channel and contiguous groundwater and finally the temporal scale. In this model the ecological integrity of temperate floodplain rivers is largely due to hydrological features, such as the flood pulse, which seasonally forms a moving littoral traversing the floodplain surface across the aquatic/terrestrial transition zone (ATTZ) (Junk et al. 1989). Temperate river systems class into two types; montane snow-fed systems and oceanic nonmontane rain-fed systems, with snow-fed systems tending to be more susceptible to flash flooding due to snowmelt. Chalk rivers in particular tend to respond relatively slowly to rainfall due to the buffering effect of aquifers. Thus hydrology plays a primary role, inducing disturbances, controlling dynamic processes and connections/disconnections, differentiating and structuring habitats, which ultimately provides a variety of habitats for fish communities (Niaman and Decamps 1990, Ward and Stanford 1995). Such floodplain habitats are important factors in the ecology of lowland river fishes.

For fishes, spatial patterns of assemblages and longitudinal zonations, as well as lateral movements, are strongly influenced by connectivity features. Fish species through their life-stage specific requirements, exploit the longitudinal and lateral dimension of the river over different time scales. Good longitudinal and lateral
connectivity is important for the ecology of many fish communities (Bayley 1995, Gore and Shields 1995, Ross and Baker 1983).

Lateral floodplain habitats include small natural flowing side-streams (small tributaries), agricultural drainage channels which tend to be slow-flowing and highly vegetated, oxbows which are also low-flow vegetated habitats and the inundated floodplain itself. They provide an important role in the ecological functioning of lowland rivers by providing a mosaic of habitats used by fish for feeding, spawning, sheltering and as a residential location (Gore and Shields 1995). Many organisms living in or near the floodplain exhibit characteristic adaptations to enable them to exploit this resource. Floodplain habitats are important for some species of fish, while providing supplementary habitat for other fish (Bayley 1995, Borcherding et al. 2002, Ross and Baker 1983). They support major concentrations of fish (Fraser et al. 1999) because they offer important habitat for a wide variety of activities such as feeding, shelter and spawning (Bell et al. 2001, Neumann et al. 1996, Sommer et al. 2001). Lateral connectivity diversifies the species composition of the whole river system. Ross and Baker (1983) caught 33 species of fish from the Black Creek Stream, Mississippi and its floodplain during spring floods. Of those 33 species, 26 were caught on the floodplain, 17 of which were caught only on the floodplain, illustrating the importance of the floodplain to the fish community in the Mississippi River. It is also true for small floodplain systems such as the River Garonne (Gozlan et al. 1998)

Floodplain systems have been shown to support greater numbers of fish than those without lateral aquatic habitats (Bayley 1995). In an investigation into the productivity of floodplains Bayley (1995) compared productivity of river-floodplain systems against that of equivalent stable water-level bodies. He found that all numeric fish yields and biomasses (for both tropical and temperate regions) were higher in floodplain systems than stable systems. Disrupting the complex connectivity system by isolating the main channel from its adjacent floodplains or by blocking longitudinal fish paths results in corresponding declines of fish populations. This ranges from lower catches to the total collapse of a particular species (Jungwirth 1998, Northcote 1998).

In many temperate floodplain systems much of the lateral, as well as longitudinal, connectivity has been lost as a result of human activities. In some systems, such as the Danube, only certain aspects such as the backwaters remain in a few areas, elevating their importance even further. Lateral side-channel and marginal habitats have been advocated as relatively more important for rehabilitation than in-stream structures in low gradient rivers (Bayley et al. 2000). The importance of off-river habitat creation and restoration is highlighted by Pretty et al., (2003) who recommend the creation of off-channel, marginal and floodplain habitats as a better strategy for rehabilitating lowland rivers, than restoration attempts in the main river channel.

In unaltered areas a rich mosaic of habitats remains on the floodplain, although little work has yet been carried out into the significance and functioning of a range of these habitats within a single catchment. With increasing threats to floodplain habitats from anthropogenic sources it is essential to understand the value of these habitats for river communities both in terms of specific individual habitat functions and as lateral habitat more generally.

### 1.2 Aims and Structure of Thesis

This study set out to generate an overview of the fish communities of lateral sidechannel habitats and the functionality of these habitats within the river system. With lateral off-river habitats increasingly threatened (Gore and Shields 1995, Olson and Dinerstein 1998) a scientific basis for management and conservation of lateral connectivity is necessary. The aim of this study was to provide this information in terms of:

- the diversity of side-channels in both their habitat and fish assemblages,
- their functionality for fish populations,
- their contribution to the recruitment of fish populations, so that recommendations could be made to managers and conservation bodies.
The River Frome in Dorset, Southern England, is a relatively unaltered river system, although like many British rivers it suffers from relatively high water abstraction.

Still, it provided a semi-natural system within which to investigate the importance of lateral connectivity; something that is increasingly rare nowadays in temperate locations.

The overall aim of this thesis is to investigate whether:
"lateral habitats significantly contribute to the fish populations of the river system." Specific aims are to examine whether:

- off-river habitats can be classified into types such as natural flowing streams, flowing millstreams and non-flowing drainage ditches by their habitats and fish species assemblages.
- Fish use side-channel habitats for a variety of different functions such as spawning, sheltering and feeding.
- Fish populations are adapted to the presence of off-river habitats in the river system.

A holistic approach was used to achieve these aims and test the hypotheses, which allowed for the investigation of the side-channel habitats at several levels. First, an overview of the habitat characteristics of each side-channel is presented with defining features highlighted (Chapter 4). The assemblages of different channels were characterised and attempts made to predict them according to habitat variables, a potentially useful technique if successful for designing and prioritising floodplain habitat management (Chapter 4). Deeper investigation into the functionality of sidechannels for fishes, and the specific functions that individual species use sidechannels for, was made by detailed analysis of movements between main river and lateral habitats (Chapter 5). Two species were then considered in detail as case studies. First, connectivity is discussed in respect to a rheophilic prey species, the dace (Leuciscus leuciscus), (Chapter 6). A methodological chapter (Chapter 7) then precedes the second species specific case study, introducing and examining home range estimation in riverine environments as this technique formed an important component of the following chapter. Then the importance of longitudinal and lateral connectivity within different life history strategies were considered with respect to a limnophilic predator, the pike (Esox lucius) (Chapter 8). These examples were selected to explore further the role that connectivity plays for various processes acting at different scales and levels. Finally the wider implications of this work, particularly
in terms of management and maintenance of lateral connectivity are discussed (Chapter 9).

It had been intended to also consider fish response to flooding and use of the inundated floodplain. Unfortunately a particularly dry period resulted in virtually no over-bank flooding during the course of the study. This issue was addressed to some extent by a large scale high flow experiment included in Appendix 14.

This thesis was carried out as part of a larger Natural Environment Research Council (NERC) thematic LOCAR (Lowland River Catchment) project, number NER/T/S/2002/00229. All radio- and PIT- tagging was carried out under UK Home Office licence.

## Chapter 2

Study Area

# 2.1 Characteristics of the River Frome Catchment 

### 2.1.1 GEOLOGY AND HYDROLOGY

The River Frome rises near Evershot (NGR ST047576), and flows south east through Dorset in the South of England for approximately 65 km , entering the English Channel at Poole Harbour. The Frome catchment covers $300 \mathrm{~km}^{2}$ and is comprised mainly of agricultural land (Casey 1969). The catchment is underlain throughout by chalk and this is exposed at the surface over the majority of the catchment. The river is described as a typical medium-sized chalk stream (Crisp et al. 1982).

Chalk is a soft, permeable, calcareous rock that readily absorbs and transmits water. Rainwater rapidly percolates into the chalk and accumulates in aquifers, which may be up to 80 m deep and may retain water for 20 years or more (Ladle and Westlake 1995). Aquifers help to regulate flow; damping the effects of heavy rainfall and releasing water continuously even during spells without precipitation. Water temperature is also buffered with groundwater being cool in summer and warm in winter, relative to air temperatures. Mean monthly water temperatures in the Frome range from $6.5^{\circ} \mathrm{C}$ in January to $17.4^{\circ} \mathrm{C}$ in July (Crisp et al. 1982). Similarly, the chemical composition of the water is more consistent than on rivers with a lower input from aquifers. The pH of the River Frome is slightly alkaline, between 7.5 and 8.5 , due to the influence of the chalk (Anonymous 2005).

Discharge in the River Frome at East Stoke was much lower during the study period than usual (Figure 2.1 shows discharge during the study period and the 3 years previously which represent typical discharge over a much longer period). This was due to a very dry spell during the study with very little rainfall. Groundwater accounts for a large proportion of the river discharge, but this may also have been drained due to the long period without rain. Seventy four percent of the total River Frome discharge at East Stoke was found to have originated as groundwater by Paolillo (1969). The mean annual current velocity upstream of East Stoke was estimated to be
$25 \mathrm{~cm} \mathrm{~s}^{-1}$ by Crisp et al (1982). However, a high local variability may be observed with mean velocities of $4-15 \mathrm{~cm} \mathrm{~s}^{-1}$ inside dense weed beds, compared to velocities in excess of $50 \mathrm{~cm} \mathrm{~s}^{-1}$ between them (Ladle and Westlake 1995).


Figure 2.1 Mean monthly flow over East Stoke gauging weir from 2000 to 2005 showing the much lower flows during this study in 2004 and 2005. Error bars represent monthly standard deviation (Source CEH unpublished data). No flow data was available for November 2004.

### 2.1.2 WATER QUALITY

The overall water quality of the River Frome is good with over $97 \%$ of the catchment demonstrating good or very good water quality (River Ecosystem classification 1 or 2) (Anonymous 2005). The river is predominantly calcium bicarbonate bearing, the source of which is weathering processes within the chalk aquifer (Howden 2004). Elevated levels of phosphate (predominantly from point source sewage inputs) are found in the river and its major tributaries the Tadnoll Brook and the River Cerne (Howden 2004) reflecting a eutrophic watercourse.

The EC Nitrates Directive ( $91 / 676 / E E C$ ) requires Nitrate Vulnerable Zones (NVZs) to be established in catchments where high or rising levels of nitrate have been identified and measures be implemented to reduce nitrate pollution. Approximately $90 \%$ of the Frome Catchment is designated as an NVZ. Nitrogen sources are both diffuse agricultural runoff and point sources of sewage effluent.

### 2.1.3 ABSTRACTIONS

There are currently 308 abstraction licences held within the Frome Catchment and a total of $58 \mathrm{Ml} \mathrm{day}^{-1}$ of water is abstracted from the Frome (Anonymous 2005). While these abstractions are spread throughout the whole area, large groundwater abstractions are concentrated on the chalk aquifer. The end use of the water abstracted determines the impact the abstraction has on the river. For example, water abstracted for use in fish farms is eventually returned to the river causing a lesser effect on water volume than water abstracted for spray irrigation, which results in a total loss to the system. However, water that has passed through fish farms is more prone to be of lower quality with high nutrient loading. The largest abstractor by volume on the Frome is aquaculture for fish farming which removes $56.5 \%$ of the total volume abstracted. Water abstracted for agriculture is only $1.4 \%$ of the total abstraction.

### 2.1.4 LANDSCAPE AND LAND USE

Dorset is predominately rural and relatively free from heavy industry. Existing industry on the Frome catchment is light and mostly related to agriculture and the two
major towns of Dorchester and Wareham, which have populations of 15,000 and 8,000 residents respectively. The majority of the countryside is used for animal pasture or arable farming, interspersed with small villages, woodlands and heathlands towards the south. Aquaculture for fish farming and watercress beds feature in some areas of the Frome. The floodplain of the River Frome features extensive historic water meadow systems. River management includes an annual mechanized weed-cut by the Environment Agency to improve summer water drainage. Excessive sedimentation is also an issue in the Upper Frome catchment. This is believed to be due to poor land management practices, causing runoff of soil into the river (Heywood and Walling 2003).

### 2.1.5 CONSERVATION DESIGNATIONS

The Frome catchment area contains one of the highest concentrations of designated areas for nature conservation in England (Anonymous, 2005). The River Frome is a UK Biodiversity Action Plan (BAP) chalk stream priority habitat. The Dorset Area of Outstanding Natural Beauty (AONB) covers a large proportion of the River Frome. The Environment Agency has a duty to protect and enhance the natural beauty of the AONB under the Countryside and Rights of Way Act.

The Frome and its water meadows downstream of Dorchester are designated as a Site of Special Scientific Interest (SSSI). Between the tidal limit of the Frome to Dorchester is a target area for restoration of floodplain grazing marsh. A Water Level Management Plan will be developed in the future for the Frome SSSI (Anonymous 2005).

### 2.1.6 RECREATIONAL USE

Angling for both salmonids and coarse fish takes place throughout the catchment, although coarse angling is focused in the lower reaches. There are few places with a public right of access to the river. The main public access to the river is at the tidal reach in Wareham. Here, there is a public right of navigation within the tidal limit, which extends just upstream of Wareham.

### 2.2 Description of Study Section of River

## Frome

The river downstream of East Stoke (Figures 2.2, 2.3, and 2.4) is able, under high flow conditions, to over-top its banks and flow freely onto the floodplain with no flood management in place. Winter floods are frequent but summer floods, although common 20 years ago, are now very rare. In the lower reaches of the river there is very little riparian tree cover but there are rich beds of submerged macrophytes, especially Ranuпculus species.


Figure 2.2 Winter (left panel) and summer (right panel) views of the River Frome at East Stoke


Figure 2.3 Upstream (left panel) and downstream (right panel) views of the tidal River Frome at Wareham.


Figure 2.4 Map of the full study area in the Frome catchment and the location of the catchment in the UK. The main river channel is marked in blue and side-channels are coloured green.

The East Stoke reach of the River Frome has a network of river channels (UK national grid references SY867863 to SY898862) (Figure 2.5). Some of these are flood relief channels, man-made millstreams, relics of historic meadow systems and natural
streams. The streams vary in surface area from Millstream the largest to Goldsacs the smallest (Table 2.1) In its lowest reach, the floodplain of the Frome widens out into pasture, marsh and some acidic heathland.


Figure 2.5 Map of the East Stoke study area of the Frome. The main river channel is marked in blue and side-channels are coloured green.

Table 2.1 Surface areas of each of the seven side-channels sampled.

|  | Surface area of 200 m section $\left(\mathrm{m}^{2}\right)$ |
| :--- | :---: |
| Railway | 573 |
| Millstream | 1200 |
| Rushton | 613 |
| Luckford | 1060 |
| Goldsacs | 453 |
| Holme Bridge | 820 |
| Flood Relief | 633 |

### 2.2.1 EAST STOKE MILLSTREAM

The millstream at East Stoke separates from the river and runs for 1.4 km before rejoining (Figure 2.5). Five hundred metres after leaving the main river the millstream runs through a fluvarium, built on the site of the old mill. Fifty metres downstream of the fluvarium is a small ( 0.5 m high) Environment Agency flow gauging weir. Upstream of the fluvarium the stream is referred to as the Millhead (Figure 2.6), and downstream the Millstream (Figure 2.7). As the Millhead leaves the main river fish will move downstream into the Millhead with the flow of water, unlike all the other side-channels sampled. The Millhead is typically 1-2 m deep with dense patches of emergent vegetation in some areas. The substrate in the Millhead is mostly silt overlying gravel and clay with gravel exposed in faster flowing areas. The millstream is shallower, reaching 1 m depth in only the deepest pools. It flows faster than the millhead and so the substrate is generally coarser, mostly sand and gravel. Discharge through the millstream can be controlled by a series of hatches both in the fluvarium and at the start of three smaller channels along the length of the millstream, which rejoin it further downstream.


Figure 2.6 Winter (left panel) and summer (right panel) views of the East Stoke Millhead.


Figure 2.7 Winter (left panel) and summer (right panel) views of the East Stoke Millstream.

### 2.2.2 NATURAL STREAMS

Natural streams are varied in type and length along the Frome. Two small tributaries of the Frome in the East Stoke area were included in this study: Goldsacs stream and Luckford Lake (Figure 2.5). Goldsacs stream is the most downstream side-channel that was sampled in this study (Figure 2.8). It is 5.5 km in length and flows across the Bovington army ranges (heathland) before joining the Frome. As a result its predominant substrate is sand. Luckford Lake stream originates from Luckford Lake on Coombe Heath nature reserve and flows for 4.5 km before joining the Frome. It is slow flowing and consequently its dominant substrate is silt (Figure 2.9).


Figure 2.8 Winter (left panel) and summer (right panel) views of Goldsacs stream.


Figure 2.9 Winter (left panel) and summer (right panel) views of Luckford Lake stream.

### 2.2.3 DRAINAGE DITCHES AND FLOOD RELIEF CHANNELS

Four agricultural drainage ditches and one flood relief channel were sampled during the project. These habitats are known to support a rich invertebrate diversity due to their lower flow and chemically different environment to the main river (Armitage et al. 2003).

Holme Bridge ditch is the furthest downstream ditch in the study area (Figures 2.5 and 2.10). It is shallow (up to 20 cm ) in its lower reach with heavy tree cover, but opens up and deepens 150 m upstream of the entrance to the Frome. Its main substrate is silt and it has submerged, emergent and floating vegetation in most unshaded areas. Moving upstream, Rushton ditch is the next agricultural drainage ditch (Figures 2.5 and 2.11). It is 800 m long and flows across grazed fields before joining the river. It is mostly over 1 m deep in the centre, generally heavily silted with some patches of gravel and supports large amounts of emergent vegetation. The furthest upstream agricultural ditch sampled is the Railway ditch (Figures 2.5 and 2.12). This ditch is 1200 m long and runs alongside a rail track for the majority of its length. It is mostly shallow (up to 50 cm ) and heavily vegetated along its entire length with emergent vegetation and some tree cover. While most of the substrate is silt there are patches of gravel originating from the railway.


Figure 2.10 Winter and summer views of Holme Bridge ditch.


Figure 2.11 Two summer views of Rushton ditch.


Figure 2.12 Winter (left panel) and summer (right panel) views of Railway ditch.

The flood relief channel (Figure. 2.5 and 2.13) is the shortest of the channels at 200 m . It is the only blind-ending channel, with a sluice gate blocking flows at the
top. In its lower stretch it reaches 1.5 m deep and is heavily vegetated, mostly with submerged vegetation. Further upstream it becomes very shallow down to only 10 cm and is heavily shaded, with little in-stream vegetation.

At the start of the study the Flood Relief was very overgrown and heavily silted and had been cut off from the main channel except under very high flows (Figure 2.14). In October 2003 it was reopened and the first 100 m was cleared of all aquatic vegetation to enhance fish access (Figure 2.13).


Figure 2.13 Winter (left panel) and summer (right panel) views of the Flood Relief channel after reopening.


Figure 2.14 Flood relief channel prior to reopening and clearance. Left panel shows the dense emergent vegetation in the flood relief channel and right panel shows the blocked connection with the main river.

## Chapter 3

## Materials and Methods

### 3.1 Habitat Monitoring

Seasonal sampling was carried out quarterly in March, June, September and December during 2003, 2004 and 2005. Seven side-streams were sampled including drainage ditches (Railway, Rushton, Flood Relief and Holme Bridge ditches), natural flowing streams (Goldsacs and Luckford streams) and a millstream (see Figure 2.5). The first 200 m of each stream was sampled in 50 m sections, separated by stop-nets, as it was thought this distance would be representative of the habitat and species assemblages throughout each channel.

Habitat surveys were carried out for every 25 m of the 200 m sections to relate to patterns of use by fishes. Dissolved oxygen ( $\mathrm{mg} \mathrm{l}^{-1}$ ) and temperature $\left({ }^{\circ} \mathrm{C}\right)$ were recorded at 25 m intervals before the water and sediment were disturbed during fishing. At these locations depth (m) was measured at the centre and $25 \%$ and $75 \%$ across the channel. The width was also measured as the wetted width (m) at each 25 m interval. Flow type (slack, glide and turbulent), vegetation (submerged, emergent and overhanging), in-stream structures (boulders, tree roots, woody debris) and substrate (silt, clay, sand and gravel) were estimated as a percentage of the 25 m stream area to $5 \%$ accuracy for each category.

A hydrolab minisonde (Hydrolab; Colorado, USA) was used to monitor dissolved oxygen, temperature and pH every fifteen minutes for fortnightly periods in sidechannels monitored with PIT detectors (see Section 3.6). Every fortnight the hydrolab was collected from the logging location, recalibrated and fitted with a new oxygen membrane and the data downloaded.

### 3.2 Seasonal Sampling of Assemblages

Fish were sampled by electric fishing each section using 50 MHz pulsed DC at 1-2 Amps with a generator supply (Electracatch International, Wolverhampton, Uk). The 50 m sections were fished once only, except one section per side-stream that was double fished in order to calculate fishing efficiency and population estimates. Triple fishing all sections would have allowed greater accuracy in the results with less variability due to variation in fishing conditions between channels and seasons. However the high silt composition of many of the channel beds would not have permitted three fishing passes to be undertaken during the same day.

Fish were identified to species and the fork length of each fish captured was measured, up to a total of 30 randomly sub-sampled individuals per species per stream section, after which individuals were counted. Where possible the sex of each fish was determined. Scale samples were taken from all pike captured (see section 3.11 for further detail). Fish were returned immediately after they were processed, unless the section was being double-shocked in which case they were held until after all fishing of that section was finished.

### 3.3 Main River Sampling

The main river from the East Stoke weir to Rushton ditch (Figure 2.5) was sampled with electric fishing once annually in March (2003-2005). It had been hoped that sampling would be carried out during more seasons but it was not possible to obtain a licence from the Environment Agency to do so. As a result, fishing was mainly carried out in order to radio and PIT tag fish and not directed at sampling the full species assemblage.

The fishing was conducted with a boom boat powered by a small diesel engine (designed and constructed by CEH). A series of cathodes trailed from the back of the boat and two circular anodes with droppers hung from the side of the boat. One person stood at each side of the boat ready to net fish and put them in the holding bin in the centre of the boat. Electric fishing was carried out with 50 MHz pulsed DC at approximately 2 Amps. The river was fished in sections with several downstream passes per section in order to maximise fish capture. Fish were processed in the same manner on the river bank as for side-stream sampling and were released near to their capture site.

The main river in the tidal reach was sampled between Ridge and Wareham Pool (Figure 2.4) in January and June during 2004 and 2005 also using a boom boat. This fishing targeted dace and roach large enough to PIT tag in order to monitor upstream migration. Several downstream passes were made and fish were processed (anaesthetised, identified, measured and PIT tagged) in the boat (away from public interference) before being released near to their capture site.

### 3.4 PIT Tagging

The minimum fish size used for tagging varied between species. Table 3.1 gives minimum tagging sizes. Those individuals large enough to receive a PIT tag were lightly anaesthetised in 2-phenoxyethanol (1:1000) in river water. Scales were removed from above the lateral line just by the dorsal fin in order to age the fish. Fish were then weighed and tagged. To insert the tag a small 4 mm incision was made on the mid-ventral line anterior to the pelvic girdle deep enough to penetrate the peritoneum. The PIT tag was inserted by gently pushing it into the body cavity. PIT tags used were either 23.1 mm long, 3.9 mm in diameter and weighing 0.6 g in air (Texas Instruments, TIRIS) or 12 mm long, 2.2 mm in diameter and weighing 0.1 g in air (UKID systems, Preston, UK) (Figure 3.1). The incision was closed with commercial grade acrylamide gel and the fish was held in a bin for recovery prior to release at the catch location. This procedure is demonstrated in Figure 3.2.

Table 3.1. Minimum size of fish PIT tagged for two different tags. Sources: (A. Pinder and M. Lucas Pers. Comm.) (Skov et al. 2005, Jepsen et al. 2001).

|  | Minimum fish size $(\mathrm{cm})$ |  |
| :--- | :---: | :---: |
| Species | TIRIS tags | UKID tags |
| Pike | 12 | 7 |
| Roach | 15 | 12 |
| Dace | 12 | 10 |
| Eel | 30 | $\mathrm{~N} / \mathrm{A}$ |
| Gudgeon | 8 | $\mathrm{~N} / \mathrm{A}$ |
| Grayling | 15 | $\mathrm{~N} / \mathrm{A}$ |
| Trout | 15 | $\mathrm{~N} / \mathrm{A}$ |



Figure 3.1 PIT tags used in the study, 23 mm Texas Instruments (TIRIS) tag on left and 12 mm UKID tag on right.


Figure 3.2 PIT tagging of a small pike. The tag is inserted through a small incision in the pike, which is then sealed with acrylamide gel.

Initially fish large enough were tagged with both sorts of tags in order to enable detection by the two types of reader used in the main river and side-channels (a UKID detector spanned the main river at East Stoke weir, while TIRIS detectors spanned the side-channel entrances, see section 3.6 for more detail). However, the tags were found to interfere with each other and subsequently fish were tagged with only one. In most cases fish reaching the minimum size were tagged with 23 mm TIRIS tags and only the smallest individuals were tagged with 12 mm UKID tags. However, half the fish caught in the tidal reach were tagged with UKID tags and the other half with TIRIS tags so that both main river movements (detected by the UKID detector at East Stoke weir) and passage into side-channels (detected by TIRIS detectors) following upstream migration could be monitored.

Recaptured fish were identified by scanning with a handheld PIT tag reader (Allflex portable RFID reader and Casper UKID reader). If a tag was recorded the number was noted and the fish weighed and measured. A note of its new catch location was made.

### 3.5 Management of Assemblage Data

With the large amount of data collected it was deemed necessary to develop an appropriate database. This was achieved through the development of an MS Access database that separated tagged and untagged fish into two separate tables. Data on tagged fish measurements and catch data were separated into two further tables. Two forms were created to enable simple and error free data entry for tagged and untagged fish as illustrated in Figures 3.3 and 3.4. Storing data in MS Access enabled the data to be interrogated more simply and to greater depth.


Figure 3.3 Data entry form for PIT- and radio-tagged fish demonstrating easy entry boxes for details of tags and two tables for (1) recording measurements of individual fish and (2) for recording details on capture and recapture. Many fields contained controls on data entered to minimise the possibility for entering incorrect values and so that PIT tag details could not be repeated.


Figure 3.4 Data entry form for non-tagged fish. This form enabled either fish measurements or counts of a species to be entered. Controls were set in fields to limit incorrect data entry.

### 3.6 PIT Telemetry

PIT telemetry was used in order to continuously monitor movements of fish between side-channels and the main river. It gave finer scale temporal information on fish movements but was more restricted in terms of the species and fish sizes tagged than electric fishing. Five pairs of PIT detectors were installed between 10 m to 20 m upstream of the entrance (so as to be sure fish detected were entering the channel and not just sheltering in the entrance) of five side-channels (Figure 3.5). These detectors were installed at Railway (Figure 3.6a) and Rushton ditches (Figure 3.6b), at Luckford Lake stream (Figure 3.7) and at the in-flowing entrance (Millhead) (Figure 3.8a) and out-flowing exit (Millstream) (Figure 3.8b) of the East Stoke millstream. These sites were selected because all were within reasonably local proximity and provided a selection of different habitats for fish to access. A single UKID PIT detector was installed on the main river (Figure 3.9) as part of a separate study that
was also utilised in this study. This main river detector did not give the direction of fish movement and worked only for fish tagged with UKID tags.


Figure 3.5 Map of the location of PIT detectors in the study area


Figure 3.6 TIRIS PIT detectors at (a) Railway ditch and (b) Rushton ditch. The capacitor array was mounted on the wooden stakes and protected from rain with tape and plastic bags


Figure 3.7 TIRIS PIT detector at Luckford Lake showing (a) the protective box holding batteries and detector electronics on the bank (the stream channel runs parallel behind the main river channel in view) and (b) the two antenna.


Figure 3.8 TIRIS PIT detectors at the (a) Millhead and (b) Millstream. The solar panel supplementing the two 110 Ah 12 V batteries can be seen by the Millstream detector. The bank of tuning capacitors in protective bags can be seen in both.


Figure 3.9 UKID PIT detector on the main River Frome at East Stoke, comprising 13 multiplexed antennae attached to hydraulic vanes. The vanes lift every four hours to clear any debris caught on them.

Side-channel PIT detectors were constructed using a commercially available radio frequency identification system (Texas Instruments TIRIS S-2000) and were based on the methodology of Zydlewski et al., (2001). The system consisted of a half-duplex (HDX) reader module (TIRIS RI-RFM-008) operating at 134.2 kHz , connected to a control module (TIRIS RI-CTL_MB2A). Power draw was 2-3 Amps depending on the tuning and size of the detector. Power source for the reader/control modules varied. Two units (Railway and Millhead) were powered by mains electricity, one supplied locally and the other through 450 m of armoured cable. Each was downregulated using a $240 \mathrm{~V}-15 \mathrm{~V}$ low noise step down transformer. The three remaining detectors (Millstream, Rushton and Luckford) were powered by two 110Ah 12V DC lead-acid batteries connected in parallel. Batteries lasted two weeks on a single charge. The millstream was supplemented by a solar panel ( 150 cm by 100 cm ), which supplemented the batteries of this the largest (width and depth) PIT detector. Data was logged onto a Flinka Fiskar logger (Flinka Fiskar, Orkellunga, Sweden) that wrote onto 32 MB compact flash cards.

Reader modules and data logger were housed in a single plastic box 25 cm long x 25 cm wide $\times 15 \mathrm{~cm}$ deep (Figure 3.10). The reader module was connected to an open loop inductor antenna that both generated an electromagnetic field to energise the tag and received transmitted signals from the tag. The antenna was constructed using 12gauge insulated THHN multi-strand wire. One or two loops were wound to form an inductor coil around the banks and bed of the stream, leaving sufficient distance at the surface of the stream for it still to be enclosed in the antenna when water levels rose. Depth and width of the antenna coils varied but reached maxima of 6.5 m wide and up to 80 cm deep. Antenna coils were supported at each bank by a wooden post, across the surface of the stream by a length of rope and were pegged to the stream bed with plastic pegs hammered in at 40 cm intervals. The two ends of the antenna were connected to the reading module through the bank of tuning capacitors (TIRIS RI-ACC-008). Selection of combinations of capacitors allowed the antenna circuit to be tuned to the resonant frequency, with fine tuning enabled using a potentiometer.


Figure 3.10 (a) Waterproof box with two units (master and slave), each containing a reader module and a control module, on the left and logger box on the right (b) Battery powered detector showing two 110Ah 12 V batteries and reader/control box all housed in a protective wooden box (c) Wooden housing of PIT equipment for mains powered reader showing mains power supply, step-down transformer and reader/control box

Each PIT array at each site had two antennae (or gates), each connected to a reader unit (labelled gate 1 and gate 2) to record the direction of travel of fish passing through. Thus, there were two antenna, spaced between 2 m and 5 m apart, each with its own bank of tuning capacitors and two reader/control modules. The reader/control modules were however, housed together in one protective plastic box and both recorded to only one data logger. One detector was assigned as the "master" and the other the "slave" in order that transmit and receive phases were coordinated, to avoid close range interference.

The detection range, measured as the distance between the plane of the antenna loop and the tag, varied with tag orientation. Maximum detection distance of 90 cm was achieved when the tag was horizontal. Minimum detection distance occurred at the centre of the antenna where the field was weakest. Minimum detection distance did not fall below 20 cm on any detector. Detectors scanned four times per second so any transponder entering the antenna field for less than a second was recorded. Such a low read-rate was possible because fish were relatively slow moving in slow flows; this compromise meant that power consumption was lower and enabled batteries to last 2 weeks. The functioning of each PIT detector was tested once a week when flash cards were changed (or more often when necessary) by passing a PIT tag through each gate
and noting the time. This tested whether the detector was detecting efficiently and that the logger was recording the correct time.

As a developing technology, problems were experienced with maintaining the continuous running of the PIT detectors. Problems ranged from the equipment itself, such as failed batteries and malfunctioning loggers not logging or producing unreadable files, to natural hazards such as floating trees destroying detectors during floods and damage by cattle to cables or equipment on a number of occasions. As a result there were periods when PIT detectors were not working and data was not collected (Appendix 2).

Problems were encountered if a tagged fish remained within the reader field for an extended period. In this case other tag records could be masked with the reader only capable of detecting one tag at a time, and well in excess of 10000 records could be recorded. The latter proved to be the greatest difficulty with the PIT system as logger files and memory could fill in a matter or days or even hours if a fish was continuously present. The issue of possible masking of other tags by a 'sitter' fish is common to PIT systems and software has been developed to reduce this problem. Such software records a single tag for a short time and then re-records it only at intervals if it is still present. This was not available with the Flinka Fiskar loggers used in this study. Placement of detectors in positions in the channel where fish are unlikely to linger (i.e. with faster flow and little cover) is one possibility to minimise the effect of 'sitters' and this was attempted in this study (although the flexibility was limited by the need to be near the channel mouth in sufficiently shallow water for the reader to be installed).

The main river PIT detector consisted of thirteen antennae, each housed within a fibreglass fin 3.5 cm wide, 48 cm long and 200 cm high across the 4.5 m wide middle section of the East Stoke weir (Ibbotson et al. 2004). Each fin was spaced at 35 cm intervals across the river in staggered formation to reduce the effect of obstruction to downstream river flow. Each antenna was connected to a 24 V DC single point decoder that continually scanned for PIT tags and stored any detections with a date and time stamp on a computer. This detector was of a full-duplex reader operating at

134 kHz for the exciter and 67 kHz for the signal back from the tags. It scanned 5 times per second.

To reduce the accumulation of debris on the equipment due to the river flow each fin was connected to a galvanised steel support structure that housed a pneumatic system. The pressure in the pneumatic system was maintained with a compressor which held the fins in a vertical position using low pressure (138-207 $\mathrm{Nm}^{-2}$ ). Once debris accumulated on any fin the force was detected by a sensor which triggered the fins to be raised clear of the water under high pressure $\left(689 \mathrm{Nm}^{-2}\right)$ allowing debris to be washed downstream. The force required to trigger a lift could be adjusted as necessary. Once a lift was completed the fins were lowered back into position. The whole process took approximately 20 seconds. A timer operated system ensured an automatic lift every 4 hours.

The active PIT detector within the fins was 150 cm deep and 40 cm long. The maximum range of the PIT tags was 20 cm so the staggered array of PIT detectors 35 cm apart provided detection across the entire central channel. Two side-channels, either side of the central channel were not instrumented, but flow only passes through these side-channels at very high flows reducing the likelihood of missing fish. The detector was set up on the downstream face of the gauging weir where the bed was uniform and flow was fast ( $\sim 2 \mathrm{~m} \mathrm{~s}^{-1}$ ). This discouraged the occurrence of 'sitters'.

### 3.7 Management of PIT data

As each TIRIS detector interrogated the antennae and potentially stored data four times per second it was possible for a huge amount of data to be created. The logger created a text file, which was converted to Excel format. Macros written in Visual Basic for Applications (VBA) were utilised to compress and analyse the data files produced by PIT detectors (Peter Knight, www.PITsoftware.co.uk). A number of macros were run. The first, "Standardise", removed the hourly time marker records and sorted the file into fish order. It merged repetitive records, to leave only one
record for each occasion that a fish was detected at one antenna, giving the arrival and departure times of the fish from the antenna. This reduced the number of records for the file by a factor of between 10 and 100 . Finally it looked up species codes in the Access database and recorded the species of each PIT tagged fish. A second macro, "Join", was then used to join consecutive files from the same PIT reader together to form a continuous dataset where applicable.

Three further macros were used to provide preliminary analysis. The first, "Movement", reported the transits of each fish through the detector and the direction in which it passed. The second, "Location", gave the location of each fish detected, whether it was on the main river or side-channel side of the detector. These macros also highlighted when a fish apparently spent an extended time between the two antennae, which would suggest that they had passed undetected through the detector. The third macro, "Count", provided (for one or more specified species) the number, over time, of fish that had been detected and were known to be in the side-channel. These three macros were only carried out on continuous data sets.

Another macro, "Overview", was used to give a pictorial representation of fish movements, based on the Combined Positions Record. In the Overview spreadsheet, each fish in the data set was given one line representing each PIT tag reader. Within each such line, each character represented one day. For each fish, each reader and each day, if the fish was detected at the reader or if its position relative to the reader was known (for continuous data runs), this was represented by an appropriate symbol. The total history of the interaction of each fish with each reader was thus shown in one line of symbols, and each fish's activity throughout the project was shown in its group of lines, one for each reader. This pictorial representation simplified events within any one day, showing the position of fish at the end of each day. It thus contained less information than the Combined Positions Report. However the overview was used for looking at initial patterns.

A final macro added and organised each new batch of PIT data into the "Combined Positions Record" (CPR). This was a fish centred file which brought together, for each fish, every movement through all of the five TIRIS readers. This large Excel file was the main TIRIS PIT data library, and the starting point for any analysis of
particular fish. The complete CPR contained 32753 individual location records of 1808 individual fish.

Examples of the standardised PIT data, the Combined Positions Report and the Overview are given in Appendices 3, 4 and 5.

### 3.8 Radio Tagging

Radio tagging was carried out on the river bank under as aseptic, sterilised conditions as possible and using sterilised instruments. Large pike ( $>55 \mathrm{~cm}$ ) were anaesthetised in 2-phenoxyethanol (1:1000 dilution in river water) to a level at which they no longer responded to external stimuli and did not attempt to maintain an upright position in the water. Throughout the procedure a constant flow of water containing 1:5000 2phenoxyethanol was washed over the gills to maintain oxygen and anaesthesia levels. A gas sterilised Biotrack TW-5 radio tag (Biotrack Ltd., Wareham, UK) of dimensions: 8.0 cm long, 1.6 cm diameter and 22 g weight in air, with an internal coil antenna was used (Figure 3.11). As only pike over 50 cm were tagged the tag constituted a very small percentage of the fish's weight (range 0.08-0.5 \%).

The tag was interfaced to a micro-controller to increase pulse rate when tilted and thus indicate periods of activity. The tag was implanted into the body cavity through an incision anterior to and just above the pelvic fin. The incision was closed with soluble Vicryl 2 mm sutures (Johnson \& Johnson Intl, Brussels, Belgium) and covered with commercial grade cryanoacrylate adhesive. A prophylactic intramuscular injection of Baytril (Bayer plc, Bury St. Edmonds, UK) ( 1 mg per 5 kg of fish) a broad spectrum antibiotic was administered to each fish before release. Pike were transferred to recovery tanks immediately after tag implantation and remained there until they had regained full consciousness (approximately 15 minutes). They were released soon after recovery from anaesthesia, as it was considered that a swift release provided the fish with the least stressful form of recovery (Crossman 1977).


Figure 3.11 Example of radio tags used during the study. From left TW-5, TW-4 and PIP 3 tags.

Small pike ( $25<\mathrm{FL}<55 \mathrm{~cm}$ ) were anaesthetised in 2-phenoxyethanol (1:1000 dilution in river water) to a level at which they no longer responded to external stimuli and did not attempt to maintain an upright position in the water. External tags were used for these fish as maximum battery life was 3 months due to the smaller weight these fish could carry. By attaching tags externally with dissolvable sutures the tag could be removed or would drop off at the end of the study period. A Biotrack TW-4 radio tag of dimensions $10 \times 8 \times 6 \mathrm{~mm}$ and 0.7 g in air with a 10 cm whip antenna and battery life of 2-3 months was used for pike over 25 cm long (percent of body weight range 0.04-0.1 \%) (Figure 3.11). Two sutures attached to the tag were passed through the body posterior to the anal fin and a backing plate on the other side of the fish and then tied. Sutures were left loose in order to minimise damage to the fish caused by the tag rubbing (Figure 3.12). A Biotrack PIP 3 radio tag of dimensions $7 \times 7 \times 4 \mathrm{~mm}$ and 0.4 g in air with a 15 cm whip antenna and battery life of $3-4$ weeks was used for pike between 10 and 25 cm long (percent of body weight range $0.4-5 \%$ ) (Figure 3.11). These were attached with a single loop passed through the body just posterior to the dorsal fin (Figure 3.13). Pike were transferred to recovery tanks until they regained consciousness and were released soon after. This method of tag attachment was preferred because it minimised contact with the body and resulted in no damage to the fish. However, it was only possible with the very smallest tags (with short battery lives) as larger tags would have come free too quickly. Tank investigations
showed that 10 to 15 cm pike tagged in this way were able to move freely through vegetation and feed and abrasion effects from tag attachment were minor.


Figure 3.12 Demonstration of the tag attachment of TW-4 tags to pike between 25 and 55 cm .


Figure 3.13 Demonstration of the tag attachment of PIP3 tags to pike between 10 and 25 cm .

Adult dace ( $>18 \mathrm{~cm}$ ) were anaesthetised in the same manner as small pike (percent of body weight range $0.2-0.6 \%$ ). A Biotrack TW-4 radio tag was attached externally just below and posterior to the dorsal fin (Figure 3.14). Two sutures attached to the tag were passed through the fish and a backing plate on the other side of the fish and then tied. Sutures were left slightly loose in order to minimise damage to the fish caused by the tag rubbing. This improvement to the method of Beaumont et al. (1996) was found to cause little damage to the fish. Dace were transferred to recovery tanks until they regained consciousness and were released soon after to minimise stress.


Figure 3.14 Demonstration of tag attachment of TW-4 tags to dace.

### 3.9 Radio Tracking

Tagged fish were located by walking the river bank whilst listening to tag frequencies until a tag was detected (Figure 3.15). A hand-held radio receiver (Sika, Biotrack Ltd.) connected to a 3-element Yagi antenna was used to locate fish. Frequencies were stored in the receiver and scanned through at 4 second intervals until a tag was located. Tag pulse rates were $0.5-1 \mathrm{~Hz}$ so the 4 second interval provided an effective compromise in scanning through the necessary range of frequencies with minimal chance of missing a tag. The location of the fish along the length of the river could be ascertained by moving to the position along the river bank receiving the strongest signal. As the distance to the tag reduced, the gain was adjusted to reduce the arc over which the signal could be detected. Once the position longitudinally along the river was found, the tracker moved five or ten metres up- or downstream in order to triangulate the position across the river. Each time a fish location, referred to as a fix, was determined, a standardised set of data were recorded on a form. Data recorded include date and time (in GMT).


Figure 3.15 The author radio tracking

The fish position was recorded as coordinates by referring to a GIS map of the river. The fix was stated as a position relative to marker posts along the bank in addition, for example 8 m upstream of T 2 (post T 2 in the main river). The exact locations of these marker posts were known, allowing the coordinates recorded in the field to be checked against those calculated from the written description of location. For each fix, the fieldworker recorded an estimate of the accuracy of that fix, based on signal strength, longitudinally and laterally. In general precision was between 0 and 2 m (in some cases the pike was seen) in both directions. In wider sections of river, or with weaker tags and when fish were in dense vegetation weakening the signal precision was lower. At worst it was 8 m .

Sixty eight pike (mean 49.2 cm ; range $9.2-101 \mathrm{~cm}$ ) and forty dace (mean 22.7 cm ; range $18.9-27.1 \mathrm{~cm}$ ) were tracked during the course of the study. Appendices 6 and 7 give the tracking duration and size of individual pike and dace respectively. Twenty five pike had been tagged during a previous study from 2000 to 2003 and continued to be monitored during this study. Tracking intensity of pike varied according to the specific objectives and is described in later chapters. For instance, pike were tracked intensively once or twice a day during the spawning period (February to June) but weekly to maintain contact during the summer. It was necessary to track dace daily due to their high levels of mobility. A number of tags failed or their frequencies
drifted significantly during the study, particularly in spring 2004 causing the tracking duration of some pike and dace to be shortened to up to only a week in some cases.

# 3.10 Analysis of Animal Movements from Telemetry Data 

The most appropriate home range estimation technique for the study was determined during an investigation using data from the radio tracked pike. A summary is given below and details of the optimal method and reasoning behind this conclusion are given in Chapter 7.

Methods used to estimate home ranges in restricted environments (e.g. rivers) are liable to overestimate home range area through the inclusion of unused habitats (e.g. river bank). This reduces the accuracy of management decisions such as habitat management which can be made from the data. Location data from 23 radio-tagged pike were used to examine the efficiency of standard home range estimators, when clipped to include only the area of home range within the river, for determining seasonal changes in habitat use. Cluster analysis demonstrated changes in core area most clearly between seasons, whilst kernel analysis most effectively showed seasonal variation in excursive activity. Range span did not statistically demonstrate any seasonal variation. Cluster analyses, kernel contour estimators and convex polygons round all locations gave progressively greater technique-based variation. Area clipping reduced out-of-river error and hence bias generated by this error, aiding detection of seasonality.

### 3.11 Scalimetry and Growth

Inspection of the hard structures of a fish can reveal the age and growth rate of that individual. The alternation between rapid and slow growth during the fish's life may be reflected in the scale structure by the spaces between the daily growth rings (Bagenal 1978). This pattern is particularly clear in cold and temperate regions with the slowest growth occurring in the winter. The number of annuli gives the age of the fish and the spaces between them reflect the growth rate.

Scales removed from pike were used to measure pike age and growth. Scale samples were always taken from the same place on the fish's body to ensure that the earliest formed scales were used for analysis. For pike, some of the earliest scales to develop are those just below the dorsal fin and above the lateral line (Figure 3.16). All scales used in this study were taken from this location.


Figure 3.16 Location of scale removal on pike

Approximately ten scales were sampled from each fish using tweezers. Unflawed scales (those without regenerated centres or deformations) were selected and placed in a petri dish containing $4 \%$ sodium hydroxide for 5 minutes. Scales were then washed in water and mounted between two microscope slides.

The slides were examined under a Projectina microscope at x10 magnification. The distance from the scale centre to each annulus and the scale edge was measured along
the axis demonstrated in Figure 3.17. The clearest three scales of each fish were measured in this way.


Figure 3.17 Pike scale demonstrating three annuli and the position of measurements of each annulus and total scale length.

### 3.12 Back-calculation and Growth Estimation

Back-calculation enables the estimation of an individual's growth history from annual growth patterns measured from hard structures (in the case of this study fish scales). It is based on an approximately proportional relationship between the linear growth of the scale and the length of the fish. The length of the fish when captured, the length of the full scale radius and the lengths of the radii at each age permit the length of the
fish at each age to be estimated. The linear regression model (Fraser 1916) was used to back-calculate fish length at age as it is a frequently used model (Dauba and Biro 1992, Mann 1974, 1976a) and gave good fit to the data set ( $R^{2}=90 \%$ ). The linear regression equation:

$$
\begin{equation*}
R_{n}=a+b L_{n} \tag{1916}
\end{equation*}
$$

Where:

$$
\mathrm{a}=\text { intercept },
$$

b = slope,
$\mathrm{R}_{\mathrm{n}}=$ scale radius at age,
$L_{n}=$ length at age

The Von Bertalanffy growth model (1938) was used to calculate individual growth of pike because it is well adapted to the logarithmic type of growth and has been widely used, allowing for easy comparison.

$$
L_{t}=L_{\infty}\left[1-e^{-K\left(t-t_{0}\right)}\right] \quad \text { Von Bertalanffy (1938) }
$$

Where:
$L_{t}=$ Length at time $t$
$\mathrm{L}_{\infty}=$ Theoretical maximum size when growth would be nil
$K=$ Constant that characterises growth speed to reach $L_{\infty}$
$\mathfrak{t}_{0}=$ Theoretical time when $L=0$ (although size at birth is often $>0$ )

Thus by using back-calculation it was possible to calculate an individual's growth from birth to capture. The parameters of the Von Bertalanffy equation can be calculated using the Ford-Walford equation (Ford 1933, Walford 1946). Size at $\mathrm{L}_{\mathrm{t}+1}$ must be plotted on a graph against size at $\mathrm{L}_{\mathrm{t}}$ to give the Ford-Walford equation:

$$
\begin{equation*}
L_{t+1}=a+b L_{t} \tag{Ford1933,Walford1946}
\end{equation*}
$$

Where:
$\mathrm{a}=$ intercept
b = slope
$L_{t}=$ length at age $t$

The output of the Ford-Walford equation was then used to calculate the parameters of the Von Bertalanffy equation:

$$
\begin{aligned}
L_{x} & =a /(1-b) \\
K & =-\ln (b)
\end{aligned}
$$

## Chapter 4

Side-Channel Characteristics and
Species Assemblages

### 4.1 Introduction

In rivers interaction between the abiotic and biotic factors affects the distribution and abundance patterns of species and thus community structure (Brown 1984). Both abiotic variability (e.g. dissolved oxygen, temperature, flow) and biotic processes (e.g. predation, competition) affect fish assemblages. Poff (1997) compared abiotic factors to multiple-scale 'filters' that control local occurrence or abundance of species. Assemblages locally reflect species' functional ability to cope with ecological conditions. Within sites, fish abundances vary with the stability or persistence of suitable habitats as well as the interaction between species (Freeman et al. 2001).

A broad means of assessing the value of a particular habitat to the entire aquatic community is to study the species assemblages of organisms using that habitat. Monitoring the community as a whole enables subtle changes in the assemblage diversity and composition to be detected. The suitability of a habitat to a particular species or age-group may vary as the conditions in the habitat fluctuate (Pessanha et al. 2003, Rueda and Defeo 2003). Long term change in species composition may also be symptomatic of anthropogenic alterations such as flood control measures (Bayley 1995, Bunn and Arthington 2002). Thus analysis of species assemblages provides a powerful tool for assessing the significance of a particular habitat within the ecosystem, as well as the functioning of the habitat supporting the community throughout the year.

Fish assemblages of aquatic floodplain systems have great potential to reshuffle during flood events and to vary according to season (Bayley 1995, Hoeinghaus et al. 2003). The underlying objective of this chapter is to compare and contrast the various different off-river habitats available in a typical chalk stream and the fish species assemblages that use them throughout the year. Thus, the importance of both a single side-channel and a network of contrasting off-river habitats within a floodplain river system are addressed. It is hypothesised that although the side-channels of a chalk stream may appear the same at the macro-scale, subtle variation in the habitat and fish community differentiates their contribution to the river system.

### 4.2 Materials and Methods

### 4.2.1 SAMPLING

Seven side-channels (Chapter 2) were sampled with electric fishing four times a year during 2003-2005. Side-channel temperature, pH and dissolved oxygen was measured with a hydrolab. Chapter 3 gives further detail on the procedures used.

### 4.2.2 ANALYSIS

Detrended correspondence analysis (DCA) with detrending by linear segments and nonlinear rescaling of axes was undertaken to confirm that the data were linear before carrying out a principle components analysis (PCA) (ter Braak and Prentice 1988). A partial PCA with year and season controlled for as covariables was completed. Environmental variables were centred and standardised to allow them to be directly comparable.

Population size within a single section of side-channel was estimated for a number of depletion sampled (double or triple fished) sections using the weighted maximum likelihood method (Carle and Strub 1978). While these estimates were unlikely to be accurate measures of population size (especially outside the 50 m section to which they applied) they did provide an indication of fishing efficiency. This allowed the accuracy of abundance and density estimates calculated from single pass fish catches to be gauged and the level of representation of these estimates of population size to be judged.

Two diversity indices were used to investigate the side-channel assemblages. Species richness $S$, the total number of species present, was a simple measure of diversity. Shannon's diversity index $H^{\prime}$ which is influenced by the total number of species and by the evenness of distribution of catch between those species was calculated with the equation:

$$
H^{\prime}=-\sum_{i=1}^{s} p_{i} \ln \left(p_{i}\right)
$$

Where $\mathrm{p}_{i}=$ the proportion of the $i$ th species in a sample

Diversity indices were calculated for the total assemblage at each location, seasonal assemblages at each location and also for each sample.

Canonical correspondence analysis was used to investigate the influence of environmental variables on species distribution after DCA. A partial CCA was undertaken with year and season controlled for as covariables so that only the interaction between species and environment factors was presented. To remove the effects of collinearity (the amount of redundancy within the environmental information) in the CCA, only those variables were selected that explained a significantly additional proportion of the variance independent of the other variables. Forward selection was used to identify the smallest subset of variables explaining the most variation in the assemblage data. Each variable was tested through Monte Carlo permutation tests (with 999 unrestricted permutations), with variables added to the model according to Bonferroni corrected $P$-values. Rare species were down-weighted in the CCA ordinations. The first two ordination axes were presented as a triplot of samples, species and environmental variables. Then simple scatterplots of the first two axes with only species on an enlarged scale were made, with each species coloured according to its hydraulic preference.

General linear models were used to predict species diversity and catch from environmental variables. The data set was split randomly into two groups, a first group of 236 samples used to create the model and a second group of 100 samples used to test the model. Variables with an $\alpha$-value $<0.05$ in the full general linear model were selected to include in the final model. Equations were then constructed from the coefficients provided and were used to test on the second data set.

A habitat stability index was created and calculated for each stream. This index used the variability (standard deviation) to give an indication of the stability of a particular side-channel relative to all side-channels. The equation used was:

$$
\text { Stability }=\sum_{\text {hab var }} S D_{\text {loc }} / S D_{T}
$$

Where $S D_{l o c}=$ standard deviation of each habitat variable for individual side-channels
$S D_{T}=$ standard deviation of each habitat variable for all locations

The relationship between this habitat stability index and species diversity was then investigated with a Pearson correlation coefficient. This method was used because of the small sample size.

Ordination methods were carried out in CANOCO 4.5 with ordination figures constructed in CanoDraw. All other graphs were made in SigmaPlot 2000. Statistical analyses were carried out in Minitab 14.

### 4.3 Results

### 4.3.1 SIDE-CHANNEL HABITAT CHARACTERISTICS

### 4.3.1.1 Temporal variation

As with most ecosystems the side-channels were found to be dynamic habitats, showing daily, seasonal and to some extent annual variation in habitat characteristics. Seasonal and annual variations in abiotic factors measured during seasonal sampling are presented in Appendix 8 and Appendix 9. Temperature, dissolved oxygen and vegetation showed the highest amount of seasonal variation. Interannual variation was apparent but at a low level. Daily fluctuations were apparent in side-channel temperature, dissolved oxygen and pH (Figures 4.1 and 4.2). Daily fluctuations in all three variables were stronger in summer than winter. Daily fluctuations in dissolved oxygen were more extreme in Railway than Luckford due to the large amount of macrophytes and very little shade in Railway.

Temperature varied seasonally, peaking in August in most side-channels and generally following the same pattern as the main river (Figure 4.3). Millstream temperatures were similar to the main river as could be expected because Millstream leaves the river and rejoins only 1.4 km downstream. The other side-channels exhibited buffered temperatures compared to the main river (Figure 4.3), suggesting that they are all in part ground water fed. Temperatures peaked at a lower temperature in the summer, were similar to the river in autumn and were slightly lower in winter and spring. There were no significant differences in year-round monthly temperatures between locations, but summer (June, July and August) temperatures were significantly higher in the main river and Millstream than in the other side-channels $(A N O V A$, Tukey test; $F=7.46, P<0.01, d f=6)$.


Figure 4.1 Daily variation over 7 days in temperature, pH and dissolved oxygen in the Railway ditch during winter (January) 2005 and summer (June) 2005.


Figure 4.2 Daily variation over 7 days in temperature, pH and dissolved oxygen in Luckford during winter (February) 2005 and summer (June) 2005.


Figure 4.3 Average monthly temperatures in 2004 for each side stream (black line) and for the main river (red line). Error bars represent standard deviation. Temperature data were collected with Tinytalk temperature loggers every 15 minutes. Temperature data for Goldsacs stream was not available. (Source CEH and University of Exeter unpublished data).

### 4.3.1.2 Side-channel variation

The gradient length of the habitat characteristics was 1.51 standard deviation units, so the data showed a linear response suggesting that principle components analysis (PCA) was appropriate (ter Braak and Prentice 1988). Principle components analysis of the habitat characteristics measured seasonally demonstrated the distribution of
environmental variables (Figure 4.4). Principle component axis (PC) 1 was dominated by flow while PC2 was characterised by woody debris and macrophyte vegetation. When describing the side-channels by their environmental characteristics the clearest separation was by flow, with Goldsacs and Millstream faster flowing than the other channels (Figure 4.4). Those slower flowing or slack channels separated according to the importance of either macrophytes and clay substrate or woody debris and silt substrate in each stream.


Figure 4.4 Principle components analysis of 18 environmental variables measured during sampling in March, June, September and December 2003 to 2005 with (a) the four 50 m sections for each side-channel grouped and (b) the correlations for variables.

### 4.3.2 SIDE-CHANNEL SPECIES ASSEMBLAGES

### 4.3.2.1 Species abundance, richness \& diversity

A total of 17 fish species were caught during the three year sampling period (Table 4.1). Minnows were significantly more abundant than any other species (ANOVA, Tukey test; $F=3.07, P<0.014, d f=6$ ) and were often estimated due to the very large numbers. Species composition varied between side-channels with some streams supporting a larger range or different assemblage of species than others.

The estimated population after two fishing passes was generally close to the number of fish caught, except in some cases for minnow and dace (Table 4.2). Confidence intervals for the population estimates were generally small except for three occasions (two bullhead and one dace) when the catch from the second pass was still high. Fish catch from the first pass was always lower than the population estimate and was often just outside the $95 \%$ confidence limit, suggesting that calculation of fish abundance made from single pass fishing often underestimated actual population size. The fact that this was the case for all species means that fish catch recorded during sampling was comparable between species in this study.

Species density as the number of individuals per $\mathrm{m}^{2}$ in each side-channel differed between locations suggesting that some streams were more heavily utilised than others (Table 4.3). Relative dominance of each species within the community also varied between side-channels (see relative catch Table 4.3), although minnows were the dominant species in all side-channels.

Table 4.1 Latin and common names and total number of fish sampled in the sidechannels over three years (2003-2005) by individual side-channel (RW, Railway; MS, Millstream; RU, Rushton; LL, Luckford; GS, Goldsacs; HB, Holme Bridge; FR Flood Relief).

| Latin name | Common | Number of fish |  |  |  |  |  |  |
| ---: | ---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | name | RW | MS | RU | LL | GS | HB | FR |
| Cottus gobio | Bullhead | 1 | 970 | 17 | 16 | 331 | 12 | 13 |
| Leuciscus leuciscus | Dace | 8 | 564 | 1 | 547 | 123 | 13 | 51 |
| Anguilla anguilla | Eel | 99 | 551 | 102 | 96 | 150 | 201 | 83 |
| Platichthys flesus | Flounder | - | 116 | - | - | 269 | - | - |
| Thymallus thymallus | Grayling | - | 34 | - | - | 2 | - | - |
| Gobio gobio | Gudgeon | - | 45 | 3 | 14 | - | 9 | 1 |
| Lampetra planeri | Brook | 4 | 24 | 27 | 16 | 256 | 2 | 2 |
| Phoxinus phoxinus | lamprey | Minnow | 6201 | 7491 | 3138 | 50979 | 1942 | 16340 |
| 126140 |  |  |  |  |  |  |  |  |
| Perca fluviatilis | Perch | - | - | 4 | 2 | - | 1 | 2 |
| Esox lucius | Pike | 153 | 52 | 83 | 52 | 2 | 58 | 73 |
| Oncorhynchus mykiss | Rainbow | - | - | - | - | 1 | - | - |
| Rutilus rutilus | trout | - |  |  |  |  |  |  |
| Salmo salar | Salmon | - | 99 | 6 | 385 | 16 | 78 | 15 |
| Gasterosteus | 3 spine | - | 499 | - | 13 | 13 | 1 | - |
| aculeatus | stickleback | 4 | 11 | 3 | 1 | 13 | 2 | 17 |
| Barbatula barbatula | Stone loach | 6 | 963 | 25 | 84 | 221 | 23 | 18 |
| Leucaspius delineatus | Sunbleak | - | - | 1 | - | - | - | - |
| Salmo trutta | Brown trout | - | 174 | - | 13 | 482 | 1 | - |

Table 4.2 Population estimates in 50 m side-channel sections calculated from depletion sampling using the weighted maximum likelihood method.

| Species | Location | Date | Catch |  |  |  | Estimated <br> Population | $\begin{gathered} 95 \% \\ \text { Confidence } \\ \text { interval } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | 1 | 2 | 3 | Total |  |  |
| Bullhead | Goldsacs | Jun 04 | 5 | 4 | - | 9 | 11 | 5.2 |
|  |  | Mar 05 | 15 | 8 | - | 23 | 27 | 7.8 |
|  |  | Jun 05 | 8 | 2 | - | 10 | 10 | 0 |
|  |  | Sep 05 | 18 | 16 | - | 34 | 60 | 42 |
|  | Millstream | Sep 04 | 26 | 13 | - | 39 | 47 | 12.7 |
|  |  | Dec 04 | 11 | 5 | - | 16 | 17 | 2.7 |
|  |  | Mar 05 | 7 | 1 | - | 8 | 8 | 0 |
|  |  | Sep 05 | 19 | 6 | 0 | 25 | 25 | 0 |
| Dace | Flood Relief | Jun 05 | 14 | 4 | - | 18 | 18 | 0 |
|  | Luckford | Dec 04 | 22 | 2 | - | 24 | 24 | 0 |
|  | Millstream | Sep 04 | 19 | 18 | - | 37 | 71 | 54 |
|  |  | Sep 05 | 142 | 42 | - | 184 | 200 | 14.5 |
| Eel | Millstream | Sep 05 | 4 | 2 | 1 | 7 | 7 | 0 |
| Flounder | Goldsacs | Jun 04 | 16 | 2 | - | 18 | 18 | 0 |
|  |  | Mar 05 | 2 | 1 | - | 3 | 3 | 0 |
|  | Millstream | Sep 04 | 5 | 2 | - | 7 | 7 | 0 |
|  |  | Dec 04 | 3 | 1 | - | 4 | 4 | 0 |
| Minnow | Flood Relief | Jun 05 | 6 | 1 | - | 7 | 7 | 0 |
|  |  | Mar 05 | 530 | 440 | - | 970 | 2868 | 1332 |
|  | Goldsacs | Jun 05 | 2 | 1 | - | 3 | 3 | 0 |
|  |  | Sep 05 | 16 | 7 | - | 23 | 26 | 6.4 |
|  |  | Sep 04 | 278 | 177 | - | 455 | 745 | 180 |
|  | Millstream | Mar 05 | 8 | 5 | - | 13 | 15 | 4.8 |
|  |  | Sep 05 | 470 | 49 | 3 | 519 | 522 | 0 |
|  |  | Dec 05 | 8 | 2 | - | 10 | 10 | 0 |
| Pike | Millstream Railway | Sep 05 | 4 | 1 | 0 | 5 | 5 | 0 |
|  |  | Jun 05 | 4 | 2 | - | 6 | 6 | 0 |
| Roach | Millstream | Sep 05 | 54 | 17 | - | 71 | 77 | 8.5 |
| Stone loach | Goldsacs | Sep 05 | 11 | 6 | - | 17 | 20 | 6.7 |
|  | Millstream | Sep 04 | 16 | 7 | - | 23 | 26 | 6.4 |
|  |  | Sep 05 | 23 | 4 | 2 | 29 | 29 | 0 |
| Trout | Goldsacs | Jun 05 | 12 | 2 | - | 14 | 14 | 0 |
|  | Millstream | Dec 04 | 5 | 1 | - | 6 | 6 | 0 |

Chapter 4 - Side-Channel Characteristics and Species Assem6lages
Table 4.3 Common name, code, mean relative catch (in \%) and mean densities (in indiv $\mathrm{m}^{2}$ ) (based on single pass fishing) of fish sampled in the
side-channels over three years (4 quarterly samples per year) (2003-2005) by individual side-channel (RW, Railway; MS, Millstream; RU,
Rushton; LL, Luckford; GS, Goldsacs; HB, Holme Bridge; FR Flood Relief).

| Commonname | Code | Relative catch (\% of total capture) |  |  |  |  |  |  | Relative density (indiv/m ${ }^{2}$ ) |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | RW | MS | RU | LL | GS | HB | FR | RW | MS | RU | LL | GS | HB | FR |
| Bullhead | Cg | 0.02 | 8.37 | 0.50 | 0.03 | 8.66 | 0.07 | 0.01 | 0.00 | 0.81 | 0.03 | 0.02 | 0.73 | 0.01 | 0.02 |
| Dace | Ll | 0.12 | 4.87 | 0.03 | 1.05 | 3.22 | 0.08 | 0.04 | 0.01 | 0.47 | 0.00 | 0.52 | 0.27 | 0.02 | 0.08 |
| Eel | Aa | 1.53 | 4.75 | 2.99 | 0.18 | 3.93 | 1.20 | 0.07 | 0.17 | 0.46 | 0.17 | 0.09 | 0.33 | 0.25 | 0.13 |
| Flounder | Pf | - | 1.00 | - | - | 7.04 | - | - | - | 0.10 | - | - | 0.59 | - | - |
| Grayling | Tt | - | 0.29 | - | - | 0.05 | - | - | - | 0.03 | - | - | 0.00 | - | ${ }^{-}$ |
| Gudgeon | Gg | - | 0.39 | 0.09 | 0.03 | - | 0.05 | 0.00 | - | 0.04 | 0.00 | 0.01 | - | 0.01 | 0.00 |
| Lamprey | L | 0.06 | 0.21 | 0.79 | 0.03 | 6.70 | 0.01 | 0.00 | 0.01 | 0.02 | 0.04 | 0.02 | 0.56 | 0.00 | 0.00 |
| Minnow | Pp | 95.75 | 64.62 | 92.02 | 97.63 | 50.82 | 97.60 | 99.78 | 10.82 | 6.24 | 5.12 | 48.09 | 4.28 | 19.93 | 199.17 |
| Perch | Pef |  | 0.00 | 0.12 | 0.00 |  | 0.01 | 0.00 | - | - | 0.01 | 0.00 | - | 0.00 | 0.00 |
| Pike | El | 2.36 | 0.45 | 2.43 | 0.10 | 0.05 | 0.35 | 0.06 | 0.27 | 0.04 | 0.14 | 0.05 | 0.00 | 0.07 | 0.12 |
| Rainbow trout | Om | - | - | - | - | 0.03 | - | - | - | - | - | - | 0.00 | ${ }^{-}$ | ${ }^{-}$ |
| Roach | Rr | - | 0.85 | 0.18 | 0.74 | 0.42 | 0.47 | 0.01 | - | 0.08 | 0.01 | 0.36 | 0.04 | 0.10 | 0.02 |
| Salmon | Ss | - | 4.30 | - | 0.02 | 0.34 | 0.01 | - | - | 0.42 | - | 0.01 | 0.03 | 0.00 | - |
| 3 Spine stickleback | Ga | 0.06 | 0.09 | 0.09 | 0.00 | 0.34 | 0.01 | 0.01 | 0.01 | 0.01 | 0.00 | 0.00 | 0.03 | 0.00 | 0.03 |
| Stone loach | Bb | 0.09 | 8.31 | 0.73 | 0.16 | 5.78 | 0.14 | 0.01 | 0.01 | 0.80 | 0.04 | 0.08 | 0.49 | 0.03 | 0.03 |
| Sunbleak | Ld | - | - | 0.03 | - | - | - | - | - | - | 0.00 | - | - | - | - |
| Trout | St | - | 1.52 | - | 0.02 | 12.62 | 0.01 | - | 0.00 | 0.14 | 0.00 | 0.01 | 1.06 | 0.00 | 0.00 |

No difference was found across the 50 m stream sections using Bonferroni corrected two-way ANOVA ( F and P values provided in Appendix 10). Thus the four 50 m stream sections were used as replicates in subsequent analyses. A wide range of species diversity occurred between side-channels with the two faster flowing streams Goldsacs and Millstream exhibiting significantly higher diversity (ANOVA, Tukey test; $\mathrm{F}=34.08, \mathrm{P}<0.01, \mathrm{df}=6$ ) (Figure 4.5). Species richness was also significantly higher in the two flowing streams (ANOVA, Tukey test; $\mathrm{F}=102.88, \mathrm{P}<0.01, \mathrm{df}=6$ ). Diversity was significantly higher in side-streams in summer and autumn than spring and winter (ANOVA, Tukey test; $F=12.45, P<0.05$ ) and only remained high in spring and winter in Goldsacs and Millstream (Figure 4.6). Species richness varied less seasonally but was significantly higher in spring and winter (ANOVA, Tukey test; $F=21.38, P<0.05$ ), contrary to species diversity. Relative diversity was more variable between side-channels during the more diverse seasons, for example Rushton exhibited the lowest diversity in summer.


Figure 4.5 Species richness and Shannon's index of diversity for the full period of sampling in the side-channels (2003-2005). RW, Railway; MS, Millstream; RU, Rushton; LL, Luckford; GS, Goldsacs; HB, Holme Bridge; FR, Flood Relief.


Figure 4.6 Species richness ■, Shannon's index of diversity for side-channel fish assemblages during different seasons (2003-2005). RW, Railway; MS, Millstream; RU, Rushton; LL, Luckford; GS, Goldsacs; HB, Holme Bridge; FR, Flood Relief.

Minnow mean seasonal distribution over the three year period may explain the seasonal trend in diversity in the side-channels as the very high catch of minnows occurred in spring and winter (Figure 4.7). Although no significant size difference was found between seasons for all side-channels (two-way ANOVA, Tukey test; $F=$ $0.79, P>0.05, d f=3$ ), size distribution of minnows did vary between channels (twoway $A N O V A$, Tukey test; $F=3.70, P<0.05, d f=6$ ). Significantly younger (smaller size classes) minnow used Luckford and Railway than Goldsacs and Millstream. However, when considering these patterns it must be kept in mind that fishing efficiency was likely to be lower for smaller fish of all species.


Figure 4.7 Seasonal catch from electric fishing (bars) and size variation (points and standard deviation error bars) of minnow (Phoxinus phoxinus) in River Frome sidechannels (Mean of 2003-2005).

### 4.3.2.2 Modelling species assemblages with environment variables

DCA showed the gradient length to be 2.15 standard deviation units and so canonical correspondence analysis was appropriate. Canonical correspondence analysis yielded two canonical axes that modelled $80 \%$ of the total variation ( $58.9 \%$ and $21.1 \%$ for axes 1 and 2 respectively) indicating a reasonably strong relationship between the 15 fish species and eight significantly contributing environmental variables (Figure 4.8). The first axis differentiated species according to flow requirements, with limnophils such as pike and perch at the left extreme of the axis favouring slack flow habitats and rheophils such as grayling and salmon tending to cluster on the right of the ordination. High scores on the second axis were associated with wider streams and higher temperature and low levels of shade. Species favouring high flows were differentiated much more along this second axis than those favouring low flow environments. Some limnophilic species such as flounder, bullhead and stoneloach were also placed on the left of the ordination, indicating that they may be more associated with shallower streams (being directly opposite the depth vector) than slow flow (Figure 4.8). The three most important habitat variables (i.e. longest vectors; (Ter Braak and Verdonschot 1995)) were stream width, slack flow and sandy substrate.


Figure 4.8 Axes 1 and 2 of canonical correspondence analysis of the influence of environmental variables on fish assemblages sampled in March, June, September and December 2003 to 2005. Only those environmental variables that make a significant contribution are presented. (Cg, bullhead; Ll, dace; Aa, eel; Pf, flounder; Tt, grayling; Gg, gudgeon; L, brook lamprey; Pp, minnow; Pef, perch; El, pike; Rr, roach; Ss, salmon; $\mathrm{Ga}, 3$ spine stickleback; Bb , stone loach; St , trout).

General linear models gave mixed success at predicting species richness metrics and individual species catches (Tables 4.4 and 4.5). Species diversity could not be successfully described by habitat diversity indices (Table 4.6). It was possible to predict species richness and diversity with habitat variables (Table 4.4; Figure 4.9) but species diversity more weakly related to the environmental variables recorded (Table
4.4). Observed and predicted values of species richness presented the closest match of all predictions attempted.

Table 4.4 Regression coefficient and equation of general linear models of diversity indices and environmental variables.

|  | $R^{2}$ | Equation |
| :--- | :---: | :---: |
| Species | $74 \%$ | $\mathrm{~S}=-21.95+0.22 \mathrm{DO}+0.28$ Glide +0.25 Slack -0.03 Clay -0.02 |
| Richness $S$ |  | Sand -0.03 Silt +0.02 Branches/logs +2.0 Pike |
| Shannon | $54 \%$ | $\mathrm{H}^{\prime}=0.121+0.47$ Pike -0.005 Overhanging tree -0.005 Dense |
| diversity $H^{\prime}$ |  | Emergent vegetation +0.04 Temperature -0.005 Depth -0.004 |
|  |  | Dense submerged vegetation +0.04 Width +0.006 Glide |

Table 4.5 Regression coefficient and significance of general linear models of species catch and environmental variables.

|  | $R^{2}$ | Equation |
| :---: | :---: | :---: |
| Bullhead | 53\% | $\mathrm{B}=0.04205+0.009$ Glide - 0.001 Sand - 0.001 Dense Emergent Veg |
| Dace | 28\% | $\begin{aligned} & \mathrm{D}=-4.515+0.04 \mathrm{DO}+0.05 \mathrm{Temp}+0.006 \text { Branches } / \text { logs }+0.04 \text { Glide } \\ & +0.04 \text { Slack }+0.006 \text { Branches/logs }-0.002 \text { Overhanging tree } \end{aligned}$ |
| Eel | 44\% | $\mathrm{E}=0.3424+0.005$ Sand -0.007 Clay +0.02 Temp |
| Flounder | 52\% | F $=0.29-0.003$ Depth - 0.001 Dense Emergent Veg +0.004 Glide 0.002 Overhanging tree |
| Lamprey | 18\% | $\mathrm{L}=0.06+0.003$ Glide -0.01 Cobbles - 0.002 Dense Submerged Veg |
| Minnow | 14\% | $\mathrm{M}=0.22+0.01 \mathrm{DO}$ |
| Pike | 18\% | $\mathrm{P}=-1.22+0.01$ Glide +0.02 Slack +0.02 Width -0.004 Branches/logs |
| Roach | 9\% | $\mathrm{R}=-0.23+0.01 \mathrm{Temp}+0.22$ Pike +0.007 Branches/logs |
| Stone loach | 30\% | $\mathrm{S}=0.13+0.005$ Glide - 0.002 Dense Emergent Veg |
| Trout | 60\% | $\mathrm{T}=-0.21+0.01$ Sand + 0.04 DO |

Table 4.6 Regression coefficient and significance of the correlation between species and habitat diversity for richness $S$, Shannon diversity $H^{\prime}$.

|  | Species $S$ |  | Species $H^{\prime}$ |  |
| :--- | :---: | :---: | :---: | :---: |
|  | $R^{2}$ | $P$-value | $R^{2}$ | $P$-value |
| Habitat $S$ | $3.1 \%$ | 0.004 | $0.4 \%$ | 0.171 |
| Habitat $H^{\prime}$ | $0.7 \%$ | 0.098 | $0 \%$ | 0.406 |



Figure 4.9 Observed and predicted species richness for each assemblage sample (each point represents a single channel by season and year) taken, predicted from habitat variables. Results are shown for a "testing" sample after model development on a separate sample. The red line indicates the perfect fit line where $x=y$, the black line gives the actual regression fit. The two blue dots are outliers not included in the regression.

The relationship between species catch and environmental factors was also variable and $R^{2}$ values ranged from 9 to $60 \%$ (Table 4.5). In general it was the species most strongly associated with faster flow and sand substrate (bullhead, flounder and brown
trout see Figure 4.6) that were most successfully predicted with habitat variables (Figure 4.10). In the more successful models created, predicted catches were generally higher than the observed number of fish (Figure 4.10).


Figure 4.10 Observed and predicted species catch for each sample taken. Results are shown for a "testing" sample after model development on a separate sample. The red line gives the perfect fit where $x=y$, the black line gives the regression fit.

Species diversity appeared to be correlated with the relative temporal stability of sidechannel habitat (Figure 4.11). Diversity and stability relationship in the Railway ditch lay a long way outside the trend of the other streams (details of the habitat stability index and results are given in Appendix 11). While Pearson correlation coefficients
were not significant for all channels combined, they were significant following the removal of Railway (Table 4.7). Given the small sample size of only seven locations this suggests that with a larger sample of side-channels there could be a strong correlation between habitat stability and the diversity of fishes supported by the sidechannel.


Figure 4.11 Correlation between species richness and Shannon diversity indices and a habitat stability index (2003-2005). Note habitat stability is an inverse metric thus as the index increases stability decreases (see section 4.2.1 for calculation).

Table 4.7 Correlation coefficients and significance for relation between habitat stability and species diversity in all channels and also excluding the outlying Railway ditch.

|  | All side-channels |  | Side-channels without Railway |  |
| :--- | :---: | :---: | :---: | :---: |
|  | Correlation | $P$-value | Correlation | $P$-value |
| Species Richness $S$ | -0.083 | 0.86 | -0.931 | 0.007 |
| Shannon diversity $H^{\prime}$ | -0.606 | 0.149 | -0.852 | 0.031 |

### 4.3.3 MAIN RIVER ASSEMBLAGES

As fishing in the main river at East Stoke was conducted once annually in March (2003-2005), seasonal comparisons in assemblage and diversity were not possible. Likewise, the efficiency of electric fishing equipment used to sample the main river was inferior for catching small fish and thus direct comparison between main river and side-channel catch and diversity was not possible. A full list of the annual catch in the main river at East Stoke is given in Table 4.8.

Table 4.8 Summary of the annual catch by boom boat electric fishing in the main River Frome from East Stoke weir to Rushton Ditch.

|  | 2003 | 2004 | 2005 |
| :--- | :---: | :---: | :---: |
| Dace | 18 | 26 | 1 |
| Perch | 0 | 1 | 1 |
| Pike | 35 | 53 | 18 |
| Roach | 2 | 1 | 0 |

### 4.4 Discussion

Side-channels of the River Frome varied according to habitat variables. In particular flow was the main determining factor, with the two faster flowing streams Millstream and Goldsacs being grouped separately to other side-channels on the PCA ordination. Perhaps not surprisingly the fish composition structure varied between channels with unique assemblages present at each channel and species occurring at different relative densities from one location to another. In a relatively small geographic scale study such as this (side-channels distributed over 4 km of river), these differences are likely due to variation in the habitat available and suitability for each species rather than an upstream/downstream zonation.

As well as seasonal changes in side-channel habitat, daily changes in factors such as temperature, dissolved oxygen and pH altered the suitability of locations to fish, particularly in summer. Buffered summer water temperatures in the side-channels (as shown in Chapter 2) may have made these habitats more suitable to fish during the day, but low levels of oxygen after dark may have rendered them inaccessible at night (Wootton 1990). These daily oxygen variations may have altered the suitability of highly vegetated streams in particular, causing them to be used less by the less mobile species, resulting in an overall reduction in diversity. While diversity indices appeared low in winter and spring in many side-channels, this could be due to the very high catch of minnows in the side-channels. The dominance of minnows ( $>90 \%$ in five of the seven streams) led to a reduction in the apparent diversity of the streams during these seasons compared to summer and autumn. However species richness, unaffected by relative dominance, did not decrease proportionately and was actually higher in winter and spring.

Flow velocity was the most important factor in determining the catch of a fish species in a given location (CCA axis 1). Flow has often been documented as being a very important descriptor of species assemblages (Bartozova and Jurajda 2001, Cattaneo 2005, Junk et al. 1989). This is not only in terms of the flow regime being a major physical habitat determinant in a location as described in this study but also in terms
of maintaining connectivity of lateral habitats and supporting the requirements of species which have developed life history requirements to natural flow regimes (Bunn and Arthington 2002).

Predictive power of environmental variables on species catch was low however. This may have come from the fact that many other factors influence fish presence. This may include biotic factors such as competition (Wootton 1990) or more complex abiotic factors that were not measured during the study.

Species richness was successfully predicted from habitat variables. The determinants of species assemblages and diversity that appear most often in the literature are flow, stream order and habitat stability or complexity (Cattaneo 2005, Grenouillet et al. 2004, Jackson et al. 2001). Habitat complexity (diversity) was not correlated with species diversity in this study, but habitat stability was (positively). Species richness in particular was strongly correlated with habitat stability, with the exception of the outlying Railway ditch. According to the habitat stability index Railway ditch was comparatively stable, yet it only supported 8 species. Railway had the highest relative density of pike of the all side-channels and this high predation pressure may have had a limiting effect on species diversity (Wootton 1990). Pike were found to be one of the significant contributing variables for the relationship between environmental variables and species richness and diversity in the general linear models. The results suggest that habitats which undergo less seasonal and annual variability are able to support a wider range of species, which supports existing literature (Grossman et al. 1998, Oberdoff et al. 2001).

The presence of side-channels in the river system increased the range of habitats available to fish over a relatively small area. All side-channels supported a community of fishes illustrating utilisation of the good connectivity to these habitats. Diversity and species presence and relative density varied seasonally according, to some extent, to habitat variables. Understanding the functioning of these habitats within the system and for particular species may help to understand what drives the use of side-channels and thus which are particularly important to preserve and maintain. This is investigated further in Chapter 5.

## Chapter 5

Seasonality of Side-Channel Habitats
for Lowland River Fishes

### 5.1 Introduction

The ecological functioning of off-river habitats will change seasonally and ontogenetically (Baade and Fredrich 1998, Grift et al. 2001, Molls 1999). It depends on the functional requirements of the animal communities which are present (i.e. refuge, feeding, spawning and nursery areas) (Baade and Fredrich 1998, Borcherding et al. 2002, Brown et al. 2001, David and Closs 2002). Also, off-river habitats are very variable in annual connectivity. Existing channels may be reconnected during flooding or areas desiccated at other times of year may be inundated.

### 5.1.1 SPAWNING

Natural and man-made off-river habitats such as abandoned and partially abandoned floodplain channels, the floodplain itself or backwater lakes such as gravel pits are valuable reproductive locations for fish (Hohausová et al. 2003, Neumann et al. 1996, Ross and Baker 1983) and nursery habitats for juveniles (Copp 1997, Garner 1996, Lusk et al. 2001). They provide a potential patchwork of habitat, often harbouring a diverse fish fauna from which fishes can be recruited to populations in the main river (Copp 1997).

Reduced flows in backwaters and floodplains aid smaller fish with lower absolute swimming performance (Muller et al. 2000). Many side-channel and floodplain habitats contain large invertebrate populations (Armitage et al. 2003, Humphries et al. 1999) creating good feeding opportunities for fast growth (Neumann et al. 1996). Predation risk is greatest while at a small size so rapid growth reduces the period of vulnerability (Bayley 1995). Whilst a proportion of some fish populations use river backwaters as additional available spawning habitat, some depend almost exclusively on recruitment from these areas, such as common bream (Abramis brama) in the Lower Rhine, Germany (Borcherding et al. 2002, Molls 1999).

Many species in temperate regions respond to rising flows as a cue for spawning, although a combination of the onset of higher flows and temperature is often the
driver to stimulate spawning (Gozlan 1998, Lucas 1992, Welcomme 1985). Many freshwater fish species commonly anticipate high flow conditions and make their migration to spawn during or just prior to the rising water (Bayley 1995). Indeed, some species use or compensate for variations in flow by varying migration distances to compensate for the river discharge at the moment of spawning (Humphries et al. 1999). Rising water levels that inundate floodplain habitat or provide access to lentic, vegetated habitat are also important for phytophilic spawners such as pike and roach.

Ontogenetic shifts in floodplain habitat use have frequently been reported. While many fish begin their early life in these habitats and migrate to the main river channel only returning to spawn (Humphries et al. 1999, Neumann et al. 1996), more complicated use has also been revealed (Molls 1999). A proportion of the adult bream population in the Rhine river do not leave after moving to and spawning in oxbow lakes (Molls 1999). Thus, these fish demonstrated a dual use of the oxbows. First as a nursery habitat, leaving at $0+$ or $1+$ and then again as a resident habitat for adults over $4+$ having returned from the juvenile phase in the main river. A similar pattern was also found in the sofie (Chondrostoma toxostoma) in the Garonne River and its tributaries, France (Gozlan 1998).

### 5.1.2 FEEDING

There is increased recognition that floodplain habitats play a major role in the productivity of riverine communities (Bayley 1995). Use of the floodplain for feeding has frequently been reported (Borcherding et al. 2002, Humphries et al. 1999, Ross and Baker 1983, Sommer et al. 2001, Welcomme 1985) with many different feeding strategies catered for on the floodplain from plant and detritus feeders to top predators. A wide range of niches may be exploited within the rich, often seasonal environment of the floodplain.

Aquatic invertebrate prey density tends to be higher in river backwaters than in the main channel (Armitage et al. 2003, Hohausová et al. 2003, Neumann et al. 1996). Some individuals may take advantage of this high prey abundance and potentially reduced predation by moving into these areas to feed. Borcherding et al., (2002) reported segregation in a population of bream with a proportion moving into a side-
channel at night to feed. They concluded that there were competitive advantages to these individuals making daily migrations to these highly productive feeding areas.

If all requirements for an individual are catered for in an off-river habitat then some individuals may benefit by being resident. Grift et al., (2001) demonstrated that immigrant bream into a floodplain lake were both smaller and significantly lower in condition than residents. Significantly better growth rates were found in chinook salmon (Oncorhynchus tshawytscha) residing on the seasonally inundated floodplain. The floodplain resident salmon increased in size substantially faster than those in the river due to greater availability of drift invertebrates (Sommer et al. 2001). Masters et al., (2002) attributed use of the floodplain by riverine pike during over-bank floods to be due to fish utilising the additional feeding opportunity. However, as almost $80 \%$ of fish locations were made within 10 m of the nearest riverbank this behaviour is more likely related to sheltering from high flows.

### 5.1.3 SHELTER

The provision of shelter is also thought to be an important role of off-river habitats but this function has rarely been investigated. However, while few studies have focused on the use of off-river habitats for shelter, it has been considered as an aside to another study. This may be due to difficulty in demonstrating evidence of sheltering in comparison with the active tasks of feeding or spawning where physiological changes or activities can be quantified.

Off-river habitats are often important shelters from both high flow and predation experienced in the main river channel. Species responses to high flows vary greatly (David and Closs 2002), but many species, for example, threespot tilapia (Oreochromis andersonni), pink happy cichlids (Sargochromis giardi), coho salmon (Oncorhynchus kisutch) and roach undertake some form of sheltering behaviour in side-channels (Baade and Fredrich 1998, Erman et al. 1988, Thorstad et al. 2001).

Side-channels also offer sanctuary from predation. Borcherding et al., (2002) interpreted the fact that $90 \%$ of ditch users were $0+$ fish to be a result of them sheltering from higher predation risk in the main river. However, as previously
mentioned these habitats offer a good, lower flow environment for $0+$ fish (Humphries et al. 1999, Muller et al. 2000). This illustrates the difficulty in clarifying the exact function of off-river habitats to visiting fish.

As discussed, species are likely to use side-channels for different functions during different seasons and throughout their lifetime. Also different channel types may differ in the resources available and so may be used for functionally different reasons. The aim of this chapter is to investigate both the seasonal functions of the side streams studied and to assess the functions for which particular species require off-river habitats. It is hypothesised that fish species will vary in their use of side-channels and the functions they for which they use them. Different species will use a different network of side-channels and each side-channel will provide resources for a distinct community of fish. The use of side-channels by pike for spawning is also investigated. It is hypothesised that side-channels are an important spawning habitat for the population. It is also hypothesised that the pike population is structured as a result of the availability of off-river spawning grounds with some individuals exhibiting protandry (early arrival onto spawning grounds).

### 5.2 Materials and Methods

### 5.2.1 SAMPLING

Seven side-channels (Chapter 2) were sampled with electric fishing four times a year during 2003-2005. Five side-channels were monitored with PIT detectors during 2004-2005. Adult pike were radio-tracked at dawn, midday and dusk over a 13-day period four times per year to determine their seasonal home range (2002-2005). See Chapter 3 for further detail on the procedures used.

Several abiotic factors were monitored continuously. Daily temperature $\left({ }^{\circ} \mathrm{C}\right)$ was monitored every 15 minutes in each channel with a TinyTag data logger (Gemini Data Loggers, Chichester, UK). Side-channel depth (cm) was monitored every hour with water level loggers (Exeter University, Geography Department). Rainfall (mm) was measured daily at East Stoke. River discharge was monitored at 15 minute intervals at East Stoke gauging weir (Environment Agency data, using Venturi gauging weir). Light levels were monitored every hour with a light meter (Centre for Ecology and Hydrology) at East Stoke gauging weir (Appendix 12 provides a summary of this data).

### 5.2.2 ANALYSIS

Seasonal catch data (single pass electric fishing of the first 200 m of each sidechannel, see Chapter 3 for more detail) are presented as figures of the mean number of fish caught per season (averaged over 3 years) with mean and standard deviation of size on a second axis. The mean number of fish recorded by PIT detectors during each season (averaged over 2004 and 2005) is presented as a figure for each species, separated for each side-channel. Daily movement patterns are presented as the ratio of fish moving in and out of side-channels in hourly groups during each season for each species at each side-channel, with the total number of fish moving on a second axis. In this case the sum and not the mean number of fish was used for each season because once the data was split into 24 hour groups the number of fish was at times small.

The relationship of daily movements to abiotic factors (daily temperature $\left({ }^{\circ} \mathrm{C}\right.$ ), sidechannel depth ( cm ), change in side-channel depth from the previous day ( cm ), rainfall $(\mathrm{mm})$, river discharge $\left(\mathrm{m}^{3} \mathrm{~s}^{-1}\right)$, change in river discharge from the previous day $\left(\mathrm{m}^{3} \mathrm{~s}^{-1}\right)$ and maximum light level on a day (lux)) was investigated with linear regression for each species. This was also separated between seasons and side-channels. Change in temperature from previous day was also investigated but is not presented because it was not significant with daily movements of any fish species. Results were Bonferroni corrected for each species to account for the large number of regressions applied.

General linear models (StatSoft.Inc. 2006) were applied for each species to investigate the main drivers of fish movement into the side channels. Included in these were the factors mentioned above as well as season and side-channel.

Seasonal home range was determined for 23 adult pike for which 13-day dawn, midday and dusk locations had been collected ( $58-101 \mathrm{~cm}$ FL). Every individual fish is not represented for each season because some fish were not tracked in each season. The mean home range size (using $95 \%$ kernels, $95 \%$ clusters and range span; see Chapter 7) is presented with standard deviation error bars. Spawning period results are presented for all pike radio-tracking during spring 2004 and 2005.

Von Bertalanffy growth coefficients $\mathrm{L}_{\infty}$ and K were calculated for the full life of each fish (see Chapter 3). The following transformations were applied to normalise the data.

Transformations used were:

$$
\begin{array}{ll}
\mathrm{L}_{\infty} & \log _{10}\left(\mathrm{~L}_{\infty}\right) \\
\mathrm{K} & \log _{10}\left(\mathrm{~K}^{*} 10000+1\right)
\end{array}
$$

Values of $K$ were multiplied by 10000 because there were zeros and many low values. This allowed the data to be transformed while the proportions between the data points were maintained.

Nominal logistic regression (Sokal and Rohlf 1995) was used to investigate the relationship between individual growth or length and seasonal presence in spawning
channels. The correlation was tested for all fish combined and also males and females separated.

A general linear model was constructed with the aim of predicting the number of days an individual pike spent in a spawning channel per month. By using fish ID as the first factor in the model, it was created so that only the variation between individuals was being predicted (and not variation due to environmental factors for example).

Home range analysis was carried out in Ranges7. Statistical analyses were carried out in Minitab 14. Graphs were made in SigmaPlot 2000.

### 5.3 Results

### 5.3.1 BULLHEAD

Bullhead were caught in all seven side-channels, although they were only caught in very small numbers in all streams except Goldsacs and Millstream. Highest catches of bullhead were in autumn in both Goldsacs and Millstream (Figure 5.1). There was no significant variation in bullhead size between Goldsacs and Millstream (GLM Tukey adhoc; $P=0.07, T=-1.804, d f=1$ ). Bullhead size in spring was significantly larger than autumn (GLM Tukey adhoc; $P=0.003, T=3.469, d f=3$ ), no other seasonal size differences were found.


Figure 5.1 Seasonal catch (bars) and size variation (points and standard deviation error bars) of bullhead (Cottus gobio) in River Frome side-channels (Mean of 20032005).

### 5.3.2 BROOK LAMPREY

Brook lamprey were caught mainly in Goldsacs although a very small number were caught in all other side-channels. There was no significant variation in size between seasons in Goldsacs (Figure 5.2). Summer was the time of highest catch and winter the lowest, although water clarity in Goldsacs in winter may have reduced fishing efficiency.


Figure 5.2 Seasonal catch (bars) and size variation (points and standard deviation error bars) of brook lamprey (Lampetra planeri) in River Frome side-channels (Mean of 2003-2005).

### 5.3.3 FLOUNDER

Goldsacs and Millstream also supported large numbers of flounder. Flounder were not found in any of the other side-channels sampled. The highest catch of flounder in Millstream occurred in autumn, while in Goldsacs flounder catch peaked in summer (Figure 5.3). After peaking in summer in Goldsacs flounder decreased in number in each of the other three seasons. Flounder size gradually increased from summer onwards and was significantly different between all seasons except autumn and winter (GLM Tukey adhoc; $P<0.05, T>2.01$, $d f=3$ in all cases). Flounder caught in Millstream were significantly larger than those in Goldsacs (GLM; $P<0.001, T=$ $8.046, d f=1)$.


Figure 5.3 Seasonal catch (bars) and size variation (points and standard deviation error bars) of flounder (Platichthys flesus) in River Frome side-channels (Mean of 2003-2005).

### 5.3.4 EEL

Eel used side-channels throughout the year and were present to some extent in every side-channel sampled (Figure 5.4). Recorded catch was generally lowest in winter suggesting this was the period of least use. Highest catch was in summer in Goldsacs, Holme Bridge, Rushton and Millstream, whereas eel catch was highest in spring in Flood Relief and Luckford, suggesting different functional use of these two groups of side-channel (Figure 5.4). There was little variation in the mean size of eel between seasons with only eel in side-channels in spring being larger than summer (GLM Tukey adhoc; $P=0.03, T=-2.777, d f=3$ ). Although fishing efficiency of small eel was probably lower than for other species. Eel in Goldsacs were significantly smaller than those in all other side-channels except Railway (GLM Tukey adhoc; $P<0.03, T$ $>3.116, d f=6$ in all cases). Eel in Luckford were significantly larger than those in Millstream and Railway (GLM Tukey adhoc; $P<0.003, T<-4.097, d f=6$ in both cases) and those in Holme Bridge were larger than those in Railway (GLM Tukey adhoc; $P=0.01, T=-3.353, d f=6$ ).

Eel demonstrated the strongest nocturnal in/out movements in spring and summer in the Millstream (Figure 5.5). Eel moved out of the Millstream in early morning and back in during the evening and overnight in spring and summer. Small amounts of movement were apparent in autumn, with no real directional gradient and there was almost no movement during the winter (Figure 5.5). Between 20 and 30 eel were
recorded at the PIT detectors in spring, summer and autumn with only 8 individuals in winter (Figure 5.6). Despite a high number of individuals recorded in autumn a lower number of daily movements were made throughout the season, indicative of lower activity during this time. Eel movements were most strongly related to abiotic factors in autumn. Channel depth, river discharge and temperature were significantly positively associated with eel movements at Millstream (Table 5.1). A general linear model ( $R^{2}=27 \%$ ) found eel use of the Millstream to vary according to season ( $p=$ 0.027 ). Temperature was a positively related to eel side-channel use ( $p=0.033$, coefficient $=1.137$ ).


Figure 5.4 Seasonal catch (bars) and size variation (points and standard deviation error bars) of eel (Anguilla anguilla) in River Frome side-channels (Mean of 20032005).


Figure 5.5 Ratio of eel (Anguilla anguilla) daily side-channel in and out movements during different seasons (bars) and total number of eel detected (line) (2004-2005).


Figure 5.6 Seasonal activity of eel (Anguilla anguilla) at each PIT monitored sidechannel used (2004-2005). Data reflect the numbers of different individual eel recorded in different seasons.

Table 5.1 Significance values of simple linear regression analysis of various abiotic factors on the level of eel (Anguilla anguilla) movement based on automated PIT records at side-channels during spring, summer, autumn and winter (2004-2005). Values represent day-to-day variations within a season. Cross-hatched areas $\mathbb{Q}$ represent periods lacking in sufficient data to perform analysis (sample size of fish detected too low or no abiotic data available) and black areas $\square$ indicate significance at an 0.002 Bonferroni corrected $\alpha$ level. Actual $R^{2}, P$-values, $n$ and coefficients are given in Appendix 13.

| EEL | Millstream | Millhead | Railway | Rushton | Luckford |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | Sp S A W | Sp S A W | Sp S A W | Sp S A W | Sp S A W |
| Mean Daily Temperature ( ${ }^{\circ} \mathrm{C}$ ) |  |  |  |  |  |
| Side-Channel Depth (cm) |  |  |  |  |  |
| Side-Channel Depth Change |  |  |  |  |  |
| Rainfall (mm) |  |  |  |  |  |
| River Discharge ( $\mathrm{m}^{3} \mathrm{~s}^{-1}$ ) |  |  |  |  |  |
| River Discharge Change |  |  |  |  |  |
| Max Daily Light (lux) |  |  |  |  |  |

### 5.3.5 DACE

Dace showed differential use of side-channels with small dace using the Flood Relief throughout the year and larger dace using Goldsacs and Luckford but mainly during spring and winter (Figure 5.7). Millstream was used throughout the year but by different size classes of fish in different seasons. Different size classes of fish used the side-channels in every season (GLM Tukey adhoc; $P<0.02, T>2.978, d f=3$ in all cases). Fish size was different between most side-channels except as mentioned previously between Flood Relief and Holme Bridge, Luckford and both Goldsacs and Holme Bridge (GLM Tukey adhoc; $P<0.02, T>2.914, d f=5$ in all cases). No dace were recorded in the Railway ditch.


Figure 5.7 Seasonal catch (bars) and size variation (points and standard deviation error bars) of dace (Leuciscus leuciscus) in River Frome side-channels (Mean of 2003-2005).

Very little or no movement was recorded at Luckford in the summer and winter with small numbers present during spring and autumn (Figure 5.8). Dace showed a diurnal pattern of movement at the Millhead in spring and summer and Millstream in spring. They moved in during the morning and out in the afternoon/evening (Figure 5.8). Less dace visited Millhead in autumn and winter with no real diel pattern. Dace did not use Millstream appreciably in the winter. While many fish visited Millstream in autumn, there was not a strong directional gradient according to the time of day (Figure 5.8).

No abiotic factors were found to affect the use of side-channels by dace using linear regression. (Table 5.2). A general linear model of all variables and also season and side-channel found a corresponding result. Only dace use of different side-channels and during different seasons were found to significantly differ ( $p<0.001$ and $p=$ 0.01 respectively; $R^{2}=26 \%$ ).


Figure 5.8 Ratio of dace (Leuciscus leuciscus) daily side-channel in and out movements during different seasons (bars) and total number of dace detected (line) (2004-2005). The peak of dace detected in the Millhead in summer (peak was 192 dace) is not shown in full to avoid loss of detail in all other seasons/locations.

Table 5.2 Significance values of simple linear regression analysis of various abiotic factors on the level of dace (Leuciscus leuciscus) movement at side-channels based on automated PIT records during spring, summer, autumn and winter (2004-2005). Values represent day-to-day variations within a season. Cross-hatched areas $\mathbb{\mathbb { D }}$ represent periods lacking in sufficient data to perform analysis (sample size of fish detected too low or no abiotic data available) and black areas indicate significance at an 0.001 Bonferroni corrected $\alpha$ level. Actual $R^{2}, P$-values, $n$ and coefficients are given in Appendix 13.

| DACE |  | ill | tre |  |  | Mil | hea |  |  | ilw |  |  | h |  |  |  | for |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  | S | A |  |  | S | A |  |  |  |  |  |  |  |  |  | A |  |
| Mean Daily Temperature ( ${ }^{\circ} \mathrm{C}$ ) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Side-Channel Depth (cm) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Side-Channel Depth Change |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Rainfall (mm) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| River Discharge ( $\mathrm{m}^{3} \mathrm{~s}^{-1}$ ) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| River Discharge Change |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Max Daily Light (lux) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |

Dace showed a much higher level of movement than was suggested by fish catches in Luckford in autumn (Figures 5.7 and 5.9). Luckford is a long stream and it is possible for fish to move further upstream than the 200 m section fished during the sampling. Radio tagged dace were found up to 800 m up Luckford during this study. Movement at Millstream was also much higher than capture for dace in spring. Millstream also extended a long distance past the 200 m sampled section and potential spawning habitat was located upstream.


Figure 5.9 Seasonal activity of dace (Leuciscus leuciscus) at each PIT monitored sidechannel used (2004-2005). Data reflect the numbers of different individual dace recorded in different seasons.

### 5.3.6 ROACH

Roach showed similar use of side-channels to dace in some cases, for example in their seasonal catch in Goldsacs and Luckford (Figure 5.10). Likewise the peak of roach catch in Millstream occurred in autumn, although at a much lower level then dace and use was only by large roach. Juvenile roach used the low flowing channel Holme Bridge, although the mean size and standard deviation showed high size variation. Roach using side-channels in summer were significantly smaller than during all other seasons (GLM Tukey adhoc; $P<0.01, T<-3.089, d f=3$ in all cases). Dace using the Millstream were significantly larger than those using all other side-channels, those using Luckford were significantly larger than those using Flood Relief and Holme Bridge (GLM Tukey adhoc; $P<0.003, T>4.270, d f=5$ in all cases).

Roach also showed a much higher level of movement than was suggested by fish catches in Luckford in autumn (Figures 5.10, 5.11). This may be for the same reason of a large amount of available habitat upstream of the 200 m sampling section as dace.


Figure 5.10 Seasonal catch (bars) and size variation (points and standard deviation error bars) of roach (Rutilus rutilus) in River Frome side-channels (Mean of 20032005).


Figure 5.11 Seasonal activity of roach (Rutilus rutilus) at each PIT monitored sidechannel used (2004-2005). Data reflect the numbers of different individual roach recorded in different seasons.

The majority of daily roach activity at Luckford PIT detector occurred during autumn (Figure 5.12). During this time there was little diel pattern in the movement, but greatest activity (movement in both directions) occurred during late morning. Visits to Millhead occurred throughout the year, although the number of fish was relatively low. During spring there is a diurnal pattern of movement out in the afternoon/evening and in during the morning (Figure 5.12). The sample was too low during other seasons to describe diel activity. Roach were active at Millstream during spring, summer and autumn but not winter. The strongest diurnal pattern shown by roach was in Millstream in spring when they moved in at dawn and out at dusk with very few roach not following this trend (Figure 5.12). In summer and autumn there was no such pattern at the Millstream.

Roach movements were only related to rainfall in the summer, with increasing rainfall associated with increased movement of roach into the Millstream (Table 5.3). Although in a number of seasons the sample was not large enough to investigate the relationship statistically. A general linear model $\left(R^{2}=25 \%\right)$ found temperature to increase movements of roach into side-channels ( $p=0.043$, coefficient $=0.304$ ) and movements to vary between seasons ( $p=0.013$ ).


Figure 5.12 Ratio of roach (Rutilus rutilus) daily side-channel in and out movements during different seasons (bars) and total number of roach detected (line) (2004-2005).

Table 5.3 Significance values of simple linear regression analysis of various abiotic factors on the level of roach (Rutilus rutilus) movement at side-channels based on automated PIT records during spring, summer, autumn and winter. Values represent day-to-day variations within a season. Cross-hatched areas $\mathbb{N}$ represent periods lacking in sufficient data to perform analysis (sample size of fish detected too low or no abiotic data available) and black areas $\square$ indicate significance at a 0.002 Bonferroni corrected $\alpha$ level. Actual $R^{2}, P$-values, $n$ and coefficients are given in Appendix 13.


### 5.3.7 PIKE

Pike were found in all side-channels throughout the year, except Goldsacs (Figure 5.13). Presence in the side-channels appeared to be higher in spring and winter, although it was only found to be significantly larger in spring than summer (GLM Tukey adhoc; $P=0.027, T<-3.205, d f=3$ ). Catch in Railway and Rushton ditches, where spawning was observed, was particularly high during spring. Pike using sidechannels in autumn were significantly smaller than those in spring or winter (GLM Tukey adhoc; $P<0.001, T>3.724, d f=5$ in both cases). Pike in the Millstream were significantly larger than all other side-channels except Luckford, pike in Luckford were also larger than those in Railway (GLM Tukey adhoc; $P<0.002, T<-3.753, d f=$ 5 in all cases).

In some cases pike PIT-recorded movement and catch (measured by electric fishing) were the same in a given season, for example in Millstream (Figures 5.13 and 5.14). However, often the point sampling carried out by electric fishing (i.e. one day in a
three month period) did not compare well to the pattern of movement shown by PITmonitoring (Figures 5.13 and 5.14). Peaks in movement into Rushton ditch in spring in particular was missed by catch sampling (Figure 5.13) but highlighted well with PIT telemetry (Figure 5.14).


Figure 5.13 Seasonal catch (bars) and size variation (points and standard deviation error bars) of pike (Esox lucius) in River Frome side-channels (Mean of 2003-2005).


Figure 5.14 Seasonal activity of pike (Esox lucius) at each PIT monitored sidechannel used (2004-2005). Data reflect the numbers of different individual pike recorded in different seasons.

Pike used Luckford throughout the year but less in the winter time. Once again the daily pattern of movement varied seasonally (Figure 5.15). The little movement that occurred in winter was during the day. Summer and spring movements showed no real pattern and movements were made throughout the 24 hours. In autumn pike moved out of Luckford in late morning and into Luckford during afternoon and early morning. Movement at Railway and Rushton ditches was more seasonally structured (Figure 5.15). Much higher levels of movement were seen in spring (GLM Tukey adhoc; $P<0.003, T>4.534, d f=3$ ) (Figures 5.14 and 5.15). In both Rushton and Railway, spring movement was during daylight hours (Figure 5.15). Fewer fish visited Rushton and Railway ditches during other seasons with no real directional gradient apparent. Pike used Millstream and Millhead less and use was spread throughout the whole year with no daily direction pattern (Figure 5.15).

Only movement of pike into and out of the Millstream was found to be associated with abiotic factors. Channel depth and discharge increased activity at the Millstream in spring and rainfall increased movement into the Millstream in autumn (Table 5.4). Both rainfall and discharge were found to influence pike side-channel use with a general linear model ( $p=0.022$ and $p=0.008$ respectively; $R^{2}=27 \%$ ).


Figure 5.15 Ratio of pike (Esox lucius) daily side-channel in and out movements during different seasons (bars) and total number of pike detected (line) (2004-2005). Spring movements are shown on a different scale due to much higher levels of activity

Table 5.4 Significance values of simple linear regression analysis of various abiotic factors on the level of pike (Esox lucius) movement at side-channels based on automated PIT records during spring, summer, autumn and winter (2004-2005). Values represent day-to-day variations within a season. Cross-hatched areas $\mathbb{\mathbb { C }}$ represent periods lacking in sufficient data to perform analysis (sample size of fish detected too low or no abiotic data available) and black areas $\square$ indicate significance at a 0.0005 Bonferroni corrected $\alpha$ level. Actual $R^{2}, P$-values, $n$ and coefficients are given in Appendix 13.

| PIKE | Millstream | Millhead |  |  |  | Railway |  |  |  | Rushton |  |  |  | Luckford |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Sp S A W | Sp | S | A | W |  |  | A | W |  | S | A | W | Sp | S | A | W |
| Mean Daily Temperature ( ${ }^{\circ} \mathrm{C}$ ) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Side-Channel Depth (cm) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Side-Channel Depth Change |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Rainfall (mm) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| River Discharge ( $\mathrm{m}^{3} \mathrm{~s}^{-1}$ ) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| River Discharge Change |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| Max Daily Light (lux) |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |

### 5.3.8 PIKE POPULATION USE OF FLOODPLAIN HABITATS FOR SPAWNING

Adult pike activity (home range area/range span size per season), monitored by radio telemetry ( $n=23 ; 58-101 \mathrm{~cm}$ FL), tended towards increasing and became more variable between individuals in spring (Figure 5.16). Both core and outer measurements of home range increased, showing that adult pike both increased their exploratory movements as indicated by an increase in outer home range area and range span and also the area in which they spent most of their time (the core).

Of all confirmed spawning events of radio-tagged pike (active spawning observed; $n$ = 7) $71 \%$ occurred in a side-channel, suggesting a high level of side-channel use for pike spawning. This value may be biased as spawning events were harder to confirm in the river because spawning could not be observed in deeper water and only increases in activity noted. However, $15 \%$ of the whole PIT tagged pike population
sample visited Railway or Rushton (two historical spawning ditches; CEH and Jerome Masters unpublished data) during the 2005 spawning season.


Figure 5.16 Seasonal activity of radio-tagged adult pike (Esox lucius) measured in terms of home range for core ( $95 \%$ cluster) and outer ( $95 \%$ kernel) and also range span (2002-2005). Spring $n=16$, summer $n=12$, autumn $n=14$ and winter $n=12$. Error bars represent standard deviation.

As well as an increase in radio-tagged pike home range size, visits to side-channels increased during the spawning season (Figure 5.15). Activity at the side-channels began in the first half of February and ended in June (Figures 5.17 and 5.18). Male and female pike entered in early February but in both years females left and did not return until early March, while activity by males continued from early February through the rest of the spawning season. The ratio of total PIT tagged female pike to total PIT tagged male pike was $1: 1.4$. Yet two to four times more males visited sidechannels than females (Female to male ratio 2004; 1:3.7 and 2005 1:2.9, illustrated in Figures 5.17 and 5.18), indicative of the spawning behaviour of pike whereby one female is often joined by between one and four males to spawn (Pictures 5.1, 5.2 and 5.3). Visits started during the same fortnight (early February) in both years and peaked during the same fortnight (late April). However the period of side-channel visits lasted longer in 2005 than 2004 with a small number of pike still making visits in July 2005, while visits ended in early June 2004.


Figure 5.17 Side-channel visits by male and female pike (Esox lucius) during the 2004 spawning season. Total number of pike detected by PIT readers in each fortnight is presented. Shading emphasises each month.


Figure 5.18 Side-channel visits by male and female pike (Esox lucius) during the 2005 spawning season. Total number of pike detected by PIT readers in each fortnight is presented. Shading emphasises each month.


Picture 5.1 Spawning pike (Esox lucius) in Flood Relief showing a larger female (bottom) being attended by 2 smaller males both on her right (top).


Picture 5.2 Spawning pike (Esox lucius) in Railway ditch showing a larger female (bottom) being attended by a smaller male (top).


Picture 5.3 Paired spawning pike (Esox lucius) in Railway ditch showing a larger female (bottom) being attended by a smaller male (top).

The number of male and female pike present in the spawning channels varied by season. From electric fishing samples in the lower 200 m of the side-channels, the sex ratio differed seasonally for all three years combined (Figure 5.19). In spring and autumn it was approximately even ( $59 \%$ and $45 \%$ male, respectively) while in winter the sex ratio was much more skewed towards males, with $79 \%$ of all pike being male.


Figure 5.19 Percentage ratio of male $\square$ to female $\square$ pike (Esox lucius) in Railway and Rushton ditches (when sex was determined). As only 3 pike of known sex were caught in these ditches in June of all three years this was not deemed an adequate sample to determine a sex ratio. The sample size is given for all seasons.

The results of nominal regression highlighted fork length and growth coefficients ( $\mathrm{L}_{x}$ and K ) as strong predictors of the difference in PIT-recorded side-channel presence of males between winter and spring (Table 5.5). Growth and size of males in Railway and Rushton was lower in winter than spring (Table 5.5). There was no such relationship for females or when sex was not included. All models of fork length and growth constants were found to fit the data significantly (Pearson Goodness of Fit, Table 5.5). Thus males, and not females, in Railway and Rushton ditches were significantly smaller and slower growing in winter than they were in spring.

Table 5.5 Results of nominal logistic regression investigating seasonal differences in growth coefficients by sex. Only pike (Esox lucius) caught in the historical spawning channels (Railway and Rushton; CEH and Jerome Masters unpublished data) were included in this seasonal analysis. Results are reported only for the comparison of spring and winter as it is the differences between these that are important to the hypothesis of early arrival.

| Predictor | Sex | Coefficient | Odds <br> Ratio | $P$-Value | Goodness of <br> fit (Pearson) |
| :---: | :---: | :---: | :---: | :---: | :---: |
| $\mathrm{L}_{\infty}$ | Both | 0.18 | 1.2 | 0.52 |  |
|  | F | -0.77 | 0.46 | 0.239 | $P>0.1$ |
|  | M | -1.26 | 0.28 | $\mathbf{0 . 0 2 6}$ |  |
|  | Both | 0.49 | 1.63 | 0.530 |  |
| K | F | -0.76 | 0.47 | 0.243 | $P>0.1$ |
|  | M | -1.27 | 0.28 | $\mathbf{0 . 0 2 5}$ |  |
|  | Soth | 0.006 | 1.01 | 0.785 |  |
|  | Size (FL) | -0.76 | 0.47 | 0.244 | $P>0.1$ |
|  | M | -1.24 | 0.29 | $\mathbf{0 . 0 2 9}$ |  |

A general linear model using the variables K, age and length described $65 \%$ of the variation between individuals in time spent in ditches (Coefficients are given in Table 5.6). Railway and Rushton were included in the model as they are historical spawning locations where spawning has been observed, while Millstream, Millhead and Luckford were not.

Table 5.6 General linear model describing individual variation in the number of days per month spent in a side-channel of the River Frome study reach by individual pike (Esox lucius) measured by PIT telemetry (2004-2005).

|  | $P$-Value | Coefficient |
| :---: | :---: | :---: |
| Constant | -0.238 |  |
| K | $\mathbf{0 . 0 2}$ | 5.13 |
| Age | 0.06 | 1.0004 |
| Length | $\mathbf{0 . 0 0 0 6}$ | 0.24679 |

Adult pike monitored by radio telemetry (that usually resided in the main river channel) often did not spawn in the nearest side-channel to their main river home range, but travelled further to another side-channel (Table 5.7). In all but one cases they travelled upstream to spawn. Nine radio tagged fish were observed visiting sidechannels during both spring 2004 and spring 2005. All of these nine fish visited the same side-channel in both years.

Table 5.7 Travel distances of pike (Esox lucius) to their nearest side-channel and to the side-channel they actually visited. RW, Railway; FR, Flood Relief; LL, Luckford; HD, Hedgemans ditch ( 2 km upstream of East Stoke).

| Fish | Distance home range to <br> nearest side-channel (m) | Nearest <br> channel | Distance home range to <br> spawning location (m) | Channel <br> visited |
| :---: | :---: | :---: | :---: | :---: |
| $\mathbf{E}$ | 67 | RW | 67 | RW |
| $\mathbf{Z a}$ | 102 | RW | 102 | RW |
| $\mathbf{A b}$ | 111 | FR | 991 | None |
| $\mathbf{F}$ | 111 | RW | 1380 | HD |
| $\mathbf{H}$ | 116 | RW | 116 | RW |
| $\mathbf{O}$ | 157 | RW | 3080 | River upstream |
| Be | 161 | RW | 161 | RW |
| $\mathbf{X}$ | 201 | LL | 1409 | RU |
| Fd | 280 | RW | 280 | RW |
| $\mathbf{Y}$ | 286 | RW | 286 | RW |
| $\mathbf{B}$ | 327 | RW | 327 | RW |
| $\mathbf{C}$ | 363 | RW | 938 | HD |
| Ze | 401 | RU | 401 | RU |
| K | 494 | FR | 591 | RU |
| Ka | 159 | RW | 159 | RW |
| J | 58 | RW | 58 | RW |
| $\mathbf{L}$ | 41 | RW | 38 | River |

[^0]
### 5.4 Discussion

Eel demonstrated strongly nocturnal behaviour, supporting existing literature (Baras et al. 1998, Schulze et al. 2004, Tesch 1977). However contrary to Baras (1998) movements extended throughout the night and did not peak soon after darkness and then reduce. Eel generally moved into the Millstream during the evening and to a smaller extent overnight and out in early morning although there was general movement in both directions. This suggests that Millstream may have been used for feeding during nocturnal activity and eel moved out to the main river during daytime inactivity. Daily activity reduced in autumn, while the total number of eel moving during the season did not. Adult eel make their seaward migrations in autumn (Tesch 1977). In autumn on the River Frome many eel only passed through the PIT detectors once in a downstream direction, seemingly on their way to sea. Eel seaward migrations tend to occur on high flows (Behrmann-Godel and Eckmann 2003) and in this study only eel movements in autumn were positively correlated with river discharge and channel depth. As $34 \%$ of eel moving through the Millstream PIT detector in autumn had not been detected at any other time these may have been migrating individuals from further upstream in the Millstream that had not made daily nocturnal forays between the millstream and river. Thus it is proposed that the Millstream provided residence habitat for many eel, some of which migrated to sea in autumn.

Some roach activity occurred at dawn and dusk which is in agreement with daily activity recorded in the literature (Baade and Fredrich 1998, Jacobsen et al. 2004), but often sample sizes were too small to show a clear pattern. Autumn was the peak of side-channel use but while this was reflected in daily activity at Luckford, there was not such an increase in daily activity at Millstream. This suggests that roach made frequent trips between Luckford and the river in autumn, while once in the Millstream they remained there. Large numbers of roach collect in the Eel pool (CEH unpublished data), which is a large pool with some flow but many sheltered, slack areas, about 1 km up the Millstream. Baade and Fredrich (1998) also found roach to be located more often in stagnant waters during colder months. It could be suggested
from this that Millstream provides shelter and feeding habitat, while Luckford is only suitable for sheltering.

Roach activity increased at Millstream, Millhead and Luckford in spring suggesting that areas of all were used for spawning activity. Roach are known to spawn in sidechannels and reed beds (Mann 1973). While Luckford and Millhead provided the vegetated habitat used by roach to spawn (Baade and Fredrich 1998, Wootton 1990) this was not available in the lower sections of the Millstream. However it is possible that roach moved up through the Millstream to spawn in the suitable habitat in Millhead further upstream. Nursery habitat was also provided by the non-flowing side-channels. Holme Bridge appeared to offer a sanctuary away from the higher flows and predation risk of the main river, possibly with a larger supply of invertebrates for feeding (Armitage et al. 2003). Holme Bridge may therefore have also been used for spawning. Although mature roach were never caught there during electric fishing sampling, this side-channel was not monitored with PIT telemetry, which was shown to better reflect transient activities such as spawning, in dace and pike.

Dace use of side-channels was similar in many respects to roach. Dace too used Luckford heavily in autumn, although unlike roach they also were present in spring. This spring use has been described as "resting-up" post-spawning (Clough et al. 1998). Use of tributaries upstream of East Stoke by post-spawning dace in the River Frome had previously been documented by Mann and Mills (1986) by fin clipping of dace spawning in Millstream and recapture in Tadnoll Brook. It was thought that sidechannels were used for sheltering and not for feeding during resting-up as $66 \%$ of the dace caught in these areas had empty stomachs and the mean fullness was $<1$ item per fish (Clough et al. 1998). Dace showed diurnal activity patterns with increased movement in both directions at dawn and dusk as has often been described in cyprinids (Clough 1997, Lucas and Baras 2001). Dace exhibited the diel movement patterns out of the side-channel at night, returning in the morning as documented between the Millhead and main river channel by Clough (1997).

Catch data suggested that the Flood Relief was used by young dace ( $<10 \mathrm{~cm}$ ) at a low level throughout the year. This is similar to the ditch use by the juvenile roach in this
study and the habitat described by Baras and Nindaba (1999). They concluded that these shallow, low flowing and heavily vegetated habitats were selected because they were warmer, provided good cover from predation and better feeding due to a higher abundance of micro-invertebrates (Baras and Nindaba 1999).

Pike were present throughout the year in all side-channels except Goldsacs. In some cases, for example Rushton and Railway in spring, movements were clearly made during the day (in both directions). At other channels pike moved at dawn and dusk, for example Luckford in autumn and Millhead in spring. However, in many cases activity occurred throughout the day and night. Pike have been shown to be flexible in the timing of daily feeding activity in different locations (Cassleman and Lewis 1996, Diana 1980, Jepsen et al. 2001, Lucas et al. 1991). On the River Frome feeding activity has previously been linked to movement at dawn and dusk (Beaumont et al. 2005). It is possible that pike were feeding in Luckford during autumn and Millhead in spring. Here, a strong diurnal pattern of activity was apparent. Combined with the knowledge that large numbers of roach used Luckford in autumn and large numbers of dace used Millhead in spring it may be hypothesised that pike were feeding in these locations, particularly as cyprinids were previously found to form the major component of pike diet on the River Frome (Mann 1976b).

Pike movements at Millstream were made throughout the 24 hours, which may suggest that pike were primarily moving in to Millstream to shelter from unsuitable conditions in the main river, when those conditions arose. The stronger positive correlation between movements into the Millstream and abiotic factors such as rainfall, river discharge and channel depth supports this premise. In many other cases the reason for moving into or out of a side-channel is less clear, perhaps as pike use them for a number of different functions simultaneously. As there was a continued presence of pike in most of the side-channels throughout the year it is probable that some individuals were resident in these areas.

Fifteen percent of the PIT tagged pike population visited Railway or Rushton during the 2005 spawning season. These were the only two side-channels monitored by PIT detectors that radio-tagged pike were observed spawning in. Deaths and emigration since tagging and immaturity of some individuals must be taken into account. Also,
the fact that a number of other suitable unmonitored side-channels (known from radio telemetry) were available in the area must be considered. With this in mind the conservative estimate of $15 \%$ of the population spawning in these two locations can be considered as a large part of the spawning population.

Male pike were more active and spent longer in spawning channels than females. This behaviour has previously been described in pike and was thought to demonstrate a polygynous mating strategy (Frost and Kipling 1967, Lucas 1992). Slow growing males were present in the spawning ditches even earlier in the season than their faster growing counterparts. No such relationship was found for female pike. A strong sex bias towards males was found in pike in side-channels prior to the spawning season (December). The mature males in the side-channels at this time were small and also slower growing. Thus they had matured at small size and not young age. It is possible that slow growing pike were resident in side-channels, however in most cases different individuals were caught in each season (determined through PIT tagging). It is also possible that slow growing pike moved into the side-channels for shelter in winter when much of the vegetation cover in the main river had died back. However, the strong sex bias toward males and the proportion of these producing milt and ready to spawn already in December suggests an alternative or additional rationale for the increasing number of pike using side-channels in winter. It is suggested that some slow growing males may have been exhibiting protandry; the early arrival of males to spawning grounds to gain spawning opportunities that may not have been available to them after larger males had arrived (Morbey and Ydenberg 2001).

Duration of time spent on the spawning grounds could be successfully predicted by a combination of size, age and growth revealing that slow growing pike remained longer in the side-channels. This may again be due to pike sheltering in the sidechannels or it may infer that the males may be using "sneaky behaviour" as a second strategy to maximise their spawning potential. By lingering in the side-channels during the spawning season the slow growing pike may maximise their chances of being present when a female entered. With the present data conclusions cannot be drawn as to which of these possibilities, if any, are the drivers of side-channel use by slow growing male pike. However, hypotheses drawn here can be used as a basis for further research. Clearer understanding could come from radio tracking all individuals
utilising a spawning ditch in order to monitor all interactions, which may indicate whether slow growing males were participating in spawning behaviour or merely sheltering away from the main river channel. Further, genetic analysis of offspring would determine which pike and, indeed how many, fathered them.

All male and female radio-tagged pike that spawned in both 2004 and 2005 returned to their spawning location of previous years. Spawning site fidelity has long been acknowledged in pike (Carbine and Applegate 1946, Frost and Kipling 1967, Karas and Lehtonen 1993, Rosell and MacOscar 2002). Recently however it has been suggested that pike return to their natal site to spawn (Miller et al. 2001). In this case there may be large consequences of the destruction or alteration of a spawning area, depending on the obligatory nature of the fidelity. If reproductive success is highly dependent on returning to a particular spawning site, the loss of that site would result in the loss of reproductive output from an entire spawning population. Constructions of levees and draining of the floodplain of the Illinois River caused the pike population to diminish (Starrett 1972). The author attributed this to loss of the spawning grounds.

In many systems, including the present study site, some spawning channels are only available under high flows. Others, following several years without significant flooding, may become blocked and impassable. In this case fish may either spawn elsewhere or skip spawning for a year. Skipped spawning has been reported in a number of fishes including three-spine stickleback (Lam et al. 1978), Atlantic salmon (Schaffer and Elson 1975), and yellow perch (Perca flavescens) (Holmgren 2003), although data on skipped spawning is limited due to the difficulties in clearly identifying non-spawning females. Females may retain their eggs if a suitable spawning location cannot be found and energy saved by not spawning in a poor year may lead to increased survival and a higher probability of spawning in subsequent years (Rideout et al. 2005). A number of female pike in this study visited spawning channels (Railway and Rushton) in only one of the two years they were monitored. While it is possible that some spawning events were missed, the large number observed visiting in only one year and the high level of spawning site fidelity exhibited by all pike between years suggests that at least some females skipped spawning. It is likely that in unsuitable years female pike skip spawning until the
following year. However if spawning channels are lost through lack of maintenance or destruction the whole pike population will be affected, particularly if pike will not spawn at other locations than their natal site.

All side-channels were utilised by some or all of the fish species investigated during this study. The niche provided by each varied according to the species, season and type of channel and for roach, dace and pike different side-channels were used for different functions. For example, dace used Millstream for spawning, Luckford for resting post-spawning and Flood Relief as a nursery habitat. Thus the full mosaic of habitats available was exploited by the fish community. This corroborates other studies that have shown the benefits of accessible off-river habitats to fishes and the losses as a result of their removal or inaccessibility (Hohausová et al. 2003, Neumann et al. 1996, Penczak et al. 2003, Schiemer 1999, Schmutz and Jungwirth 1999, Scott and Nielsen 1989).

Goldsacs and Millstream supported a number of species that were not present elsewhere in side-channels. Goldsacs in particular supported the only large numbers of brook lamprey, a Biodiversity Action Plan species, in the area sampled, elevating the importance of this habitat within the system. Goldsacs may also be a freshwater nursery ground for flounder which peaked in abundance in summer when they move in from sea (Greenhalgh 1999) and reduced in number thereafter. Flounder were smallest in summer and gradually increased in size during the following three seasons.

The level of contribution of lateral habitats to the survival of the population was considered for pike, but further investigation of population structure and use of sidechannels may provide extra information as to the necessity of these habitats within the river system and the functionality for particular species. Two case studies on the functional use of side-channels by pike and dace are described in the following two chapters.

## Chapter 6

## Population Structure and Side-

Channel Use: A Case Study of a
Rheophilic Species

### 6.1 Introduction

Dace are a rheophilic cyprinid found in fast flowing rivers and streams throughout England and E Wales (Wheeler 1969). They congregate in large shoals at all life stages, except the largest individuals which may be more solitary (Bagenal 1973). They attain a maximum size of $20-25 \mathrm{~cm}$ or up to 600 g (Davies et al. 2004). Dace are opportunistic feeders, feeding on insects and crustaceans and allocanthous material and grazing on algae (Davies et al. 2004). Unlike most other British cyprinids dace are thought to continue to feed opportunistically, at a reduced rate, throughout the winter (Davies et al. 2004). Dace are known to undertake migrations of up to 30 km and are able to osmoregulate in the salinities found in tidal reaches (Cowx 2001).

Dace are the first cyprinids to spawn each year, with spawning taking place between February and early April. Onset of spawning migration is thought to depend on water temperature (Mann 1974, Alabaster and Lloyd 1982) and photoperiod (Brook and Bromage 1989). Dace are lithophilous spawners, spawning communally over gravelly, well oxgenated shallows of rivers and streams (Davies et al. 2004). Females release a single batch of eggs during the two to three week spawning period (Mann and Mills 1986, Mills 1991). Female fecundity is a function of length and varies between 6550 and 9500 eggs for 20 cm females (Cowx, 2001).

Lakes, rivers and streams offer a mosaic of habitats that influence the persistence and structure of fish populations. Over the year, fish move between different habitats, depending on their requirements and activities, such as feeding, spawning or sheltering (Borcherding et al. 2002, Harden Jones 1968, Northcote 1978). These seasonal patterns can be regarded as an attempt to optimise habitat use, for example for spawning, foraging, or as a trade-off between predator avoidance and resource use (Krebs and Davies 1997). Separate age classes of fish or those at different of developmental stages have different resource requirements and thus their ecological needs differ.

A key to understanding spatially structured populations is in the movements of individuals between patches or habitats (Szacki 1999). The extent and scale of movements are of particular significance in understanding mobile populations, yet monitoring these movements is dependent on the method used to obtain measurements and movements are often underestimated (Szacki 1999). Defining population patches and habitats used by riverine fish and monitoring movement between them at the population level is a difficult task due to the difficulty of identifying individuals. So few studies have considered population structure and population scale movement in fishes (but see (Dunham and Rieman 1999, Gotelli and Taylor 1999, Koizumi and Maekawa 2004)). However, now the use of PIT telemetry opens a new possibility in characterising spatial population structure of fish.

The fate of mammal and bird populations fragmented by habitat destruction and isolation has been commonly investigated using theoretical models such as metapopulation theory, attracting interest from conservation managers as well as population ecologists (Hanski 1994, Lindenmayer et al. 2001, Lindenmayer et al. 1995). Yet, although little studied, the effects of habitat fragmentation on the population dynamics of stream-dwelling organisms like fish are arguably even more serious than terrestrial organisms because the colonisation route is restricted linearly. Thus, construction of movement barriers (e.g. dams, thermal plumes or impassable road culverts) may compromise their persistence, particularly in upstream areas, through the destruction of dispersal and migration routes (Jager et al. 2001, Morita and Yamamoto 2002, Reyes-Gavilan et al. 1996).

This study addressed the issues discussed above by using a number of different methods (electric fishing captures, mark-recapture, PIT telemetry and radio telemetry) to monitor dace spatial population structure within the river system. Identification of individual dace with telemetry methods allowed repetitive use of habitats to be monitored at a finer temporal and spatial scale than previous studies. The aims of this chapter were to investigate functional main river and side-channel use by dace and understand the structure of the dace population in the Frome Catchment.

### 6.2 Materials \& Methods

### 6.2.1 SAMPLING

Seven side-channels (Chapter 2) were sampled with electric fishing four times a year during 2003-2005. Five side-channels were monitored with PIT detectors during 2004-2005. Dace were radio-tracked daily during spring 2004 and 2005 and autumn 2004. See Chapter 3 for further detail on the procedures used.

### 6.2.2 ANALYSIS

Data presented for dace catch from electric fishing captures in side-channels or the main river use combined data from 2003 to 2005. Data for main river PIT telemetry is presented from July 2004 to June 2005 (after which time sensitivity of this detector decreased and became unreliable). This time period was associated with the availability of a large sample of dace tagged in the main river and side-channels. Results from side-channel PIT telemetry use data from January 2004 to December 2005 except for Luckford as the PIT detector was built in February 2005. Main river (FDX, 12 mm tag) and side-channel (HDX, 23 mm tags) PIT detectors read different tags so in most cases records were mutually exclusive and recorded different fish (except in the instances of double tagged fish - see Chapter 3).

Due to missing weeks in the data one week per month was selected for PIT data from each side-channel (as close to the middle of the month as available data would allow) and all movements during that week noted. Use of one week was necessary because there were occasions when a PIT detector operated successfully for only one week out of the month. This provided a standardised dataset that could be compared across months, seasons and locations. This was done for 2004 and 2005 and averaged for each season, except for Luckford where the reader was only installed for 2005.

The main river PIT detector was not directional i.e. it recorded the presence of a fish but not the direction in which it was travelling. However, all fish were initially tagged
and released downstream of the detector. For side-channel PIT telemetry, movements in and out were combined and only one in and one out movement were counted per fish per day to prevent influence of fish loitering in the vicinity of detectors. While numbers of fish moving through PIT detectors are presented it must be kept in mind that only $5 \%$ of the population $>12 \mathrm{~cm}$ was tagged (see Chapter 5).

Seasons for dace were divided according to the dace spawning season in the Frome (Mann 1974). Thus:

| Spring | February - April |
| :--- | :--- |
| Summer | May - July |
| Autumn | August - October |
| Winter | November - January |

Mature and immature dace were defined by size according to Mann (1974). Mann gave 16.3 cm as the mean size of maturation of males and 17.9 for females. As it was only possible to tell the sex of dace around spawning time an average size at maturity for males and females was taken:

|  | Size |
| :--- | :--- |
| Immature | $0-17.09 \mathrm{~cm}$ |
| Mature | $17.1+\mathrm{cm}$ |

Statistical analyses were carried out in Minitab 14. Graphs were made in SigmaPlot 2000.

### 6.3 Results

### 6.3.1 POPULATION STRUCTURE OF DACE USING SIDE-CHANNELS

Within the side-channels dace fork length was significantly different between all seasons (ANOVA, Tukey test; $F=56.66, P<0.001, d f=3$ ). A wider size range of dace were present in the side-channels during spring and summer than during the latter half of the year (Figure 6.1).


Figure 6.1 Seasonal size distributions of dace (Leuciscus leuciscus) in side-channels from 2004 to 2005. Lengths are given as fork length. The box plots show the median, $10^{\text {th }}, 25^{\text {th }}, 75^{\text {th }}$ and $90^{\text {th }}$ percentiles. Remaining data points are values outside these boundaries.

Of the dace using side-channels only the smallest individuals were captured in the non-flowing channels (Figure 6.2). While the size range of dace in the three flowing channels was less variable, ANOVA post-hoc tests show that all were significantly different from each other (ANOVA, Tukey test; $F=205.72, P<0.001, d f=3$ ). Dace catch in side-channels varied greatly between seasons, with greatest numbers of dace caught in a single channel peaking in September and reaching a minimum in June (Figure 6.3). Very few mature dace were caught in side-channels in June, while in contrast almost half of the catch in March was mature fish.


Figure 6.2 Size distribution of dace (Leuciscus leuciscus) caught in side-channels (2004-2005). The box plots show the median, $10^{\text {th }}, 25^{\text {th }}, 75^{\text {th }}$ and $90^{\text {th }}$ percentiles. Remaining data points are values outside these boundaries.

Seasonal side-channel use also varied between channels (Figure 6.3). In March mostly mature fish were captured in the Millstream, with some other mature fish visiting other channels also. Immature dace were present in all side-channels. Lowest use of the side-channels occurred in June and then almost exclusively by immature individuals. Millstream was heavily used by mature and immature dace alike in September. Indeed, in September very few fish were caught in other side-channels. By December the importance of Millstream had diminished and Luckford supported the
greatest number of dace. Once again, as in summer, relatively fewer mature fish were caught in the side-channels.


Figure 6.3 Seasonal distribution of immature and mature dace (Leuciscus leuciscus) in the side-channels as a percentage of total fish captured (2004-2005). Dace capture for all non-flowing channels has been pooled due to low sample size.

### 6.3.2 SPATIAL DISTRIBUTION OF THE DACE POPULATION

Movements in the main river measured by the East Stoke PIT detector peaked in May with elevated movement detected between March and July (Figure 6.4). There was little repeat detection of individuals during this time. When separated by maturity, immature fish constituted a larger number of those movements than mature dace (Figure 6.4). While both showed increased activity between March and July the largest peak was of immature and not mature dace. No such peak in May was
observed through the Millhead PIT detector which is immediately adjacent to the main river detector.


Figure 6.4 Seasonal movement of mature and immature dace (Leuciscus leuciscus) through the main river PIT detector (2004-2005).

Many long distance movements were recorded by PIT telemetry with $24 \%(n=13)$ of dace recorded as moving through the East Stoke main river PIT detector having been PIT tagged in Wareham, the tidal reach some 10 km downstream. When recaptures were considered for all locations, $11.5 \%(n=3)$ of all East Stoke (main river reach and side-channel) recaptures were fish that had moved upstream from Wareham (Table 6.1).

Approximately $3 \%$ of dace were recaptured in the large scale area (i.e. East Stoke or Wareham) in which they were tagged, and $1 \%$ of dace had made the long distance movement between tidal and upstream locations (Table 6.1). There was no difference according to location or direction of movement. Although based on small numbers of recaptures these results suggest that one quarter of all tagged dace move between the tidal and upstream areas of river (Table 6.1).

Table 6.1 Movements of recaptured dace (Leuciscus leuciscus), demonstrating those that remained in the general tagging area and those that moved between the upstream and tidal reaches. Three values are given; the total number of fish tagged, the total number of fish re-caught and recaptures as a percentage of all fish tagged in that area.

| Tagging <br> Location | Recapture <br> Location | Total number <br> tagged in the area | Total number <br> recaptured | \% of total tagged <br> in that area |
| :--- | :---: | :---: | :---: | :---: |
| East Stoke | East Stoke | 852 | 26 | 3.1 |
| Wareham | Wareham | 321 | 9 | 2.8 |
| East Stoke | Wareham | 852 | 9 | 1.1 |
| Wareham | East Stoke | 321 | 3 | 0.9 |

Radio telemetry of a smaller ( $n=32$ ) sample of dace gave differing results on the mobility of individuals between upstream and tidal sections. During this part of the study $54 \%$ of all individuals visited Wareham. In both spring and autumn the same proportion of the radio-tagged population (33\%) used side-channels, despite the different number of fish sampled (Table 6.2). However, dace were twice as likely to visit Wareham in spring as in autumn (Table 6.2). Dace that visited Wareham travelled significantly further than those that did not ( $t$-test, $P<0.001$ ) (Figure 6.5b). There was no significant difference between distance moved by dace in spring and autumn ( $t$-test, $P>0.10$ ) (Figure 6.5a). Dace that used side-channels did not travel significantly further than those that did not $(t$-test, $P>0.10)$ (Figure 6.5c).

Table 6.2 Number and percentage of radio-tagged adult dace (Leuciscus leuciscus) using a side-channel or visiting the tidal reach in spring or autumn (all dace were tagged in the East Stoke area).

|  | Spring |  |  |  | Autumn |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | n | $\%$ | n | $\%$ |  |  |
| All fish | 15 | 100 |  | 9 | 100 |  |
| Use side-channel | 5 | 33 |  | 3 | 33 |  |
| Visited Wareham | 10 | 67 |  | 3 | 33 |  |



Figure 6.5 Mean distance moved (mean of maximum distance moved by each fish during tracking period) by radio tracked dace (Leuciscus leuciscus) (a) during different seasons, (b) by those that visited and did not visit the tidal reach and (c) by those that used side-channels and those that did not (2004-2005). Error bars represent standard deviation.

### 6.3.3 POPULATION STRUCTURE INFLUENCES ON SPATIAL DISTRIBUTION

Most dace detected at side-channels were originally tagged in Millstream, even for those detected at Luckford, 3 km downstream of the Millstream entrance (Figure 6.6). More movements at Luckford were recorded from dace tagged at Luckford than were detected elsewhere, as may be expected. Ten percent of fish PIT detected at the Millhead and Millstream were first tagged at Luckford. No fish tagged in Goldsacs visited Luckford, despite it being 2 km upstream of Goldsacs and nearer to Goldsacs than any other side-channels in the East Stoke reach. Millstream and Millhead entrances which were visited by fish tagged in Goldsacs are 5 and 8 km upstream respectively. In all three locations $5 \%$ of fish PIT detected had been tagged in Wareham.


Figure 6.6 Proportions of fish detected at each side-channel PIT detector tagged in different locations. Total numbers of dace are given under each title.

Dace fork length in Luckford was found to be significantly smaller than both Millhead and Millstream when ANOVA was carried out on $\log _{10}$ transformed fork lengths ( $A N O V A$, Tukey test; $\mathrm{F}=5.11, P=0.006, d f=2$ ). There was no difference between Millhead and Millstream. A higher proportion of immature dace visited Luckford than Millhead or Millstream (Figure 6.7).


Figure 6.7 Percentage of mature and immature dace visiting each site (2004-2005).

Individual use of side-channels varied considerably (Figure 6.8). More than half ( $55 \%$ ) of TIRIS PIT tagged individuals were never recorded on side-channel PIT detectors (those fish that were tagged outside PIT monitored areas). Of those that did use the side-channels, most ( $33 \%$ ) visited only one of the three that were monitored, with a small number visiting two or even all three. Similarly $58 \%$ of all fish using side-channels visited only during one season (Figure 6.9) with a decreasing number visiting in two, three or four seasons ( $29 \%, 9 \%$ and $4 \%$ respectively).


Number of side-channels visited
Figure 6.8 The number of different side-channels fitted with a PIT detector visited by individual dace (Leuciscus leuciscus) (2004-2005).


Figure 6.9 The number of different seasons individual dace (Leuciscus leuciscus) were recorded visiting side-channels by PIT telemetry (2004-2005).

A number of fish were found to leave a side-channel site, enter the main river and then to return to the same side-channel during the same season the following year
(Figure 6.10). Returns to Millstream, a known spawning location, were much lower in all seasons including spring than returns to Millhead which reached $30 \%$ (of all fish tagged or recorded there previously) in autumn and winter 2005.


Figure 6.10 Returns of individual dace (Leuciscus leuciscus) in 2005 to the same channel as during the same season of 2004.

As mentioned previously the majority of fish detected used only one monitored sidechannel during one season. Of those that visited more, individuals were equally likely to visit one side-channel in two seasons, two side-channels in one season or two sidechannels in two seasons Table 6.3. A small number of individuals visited a higher number of side-channels or during a larger number of seasons or both.

Table 6.3 The number of PIT-monitored side-channels used by individual dace (Leuciscus leuciscus) and the number of seasons in which they visit these habitats (2004-2005).

|  | No. seasons |  |  |  |
| :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 |
| 1 channel | 284 | 35 | 3 | 1 |
| 2 channels | 38 | 43 | 10 | 3 |
| 3 channels | 1 | 3 | 4 | 2 |

In order to summarise the complex dace movements and side-channel use, a diagram illustrating these movements was prepared (Figure 6.11). Physical recapture data was used rather than detections from PIT telemetry so that more locations could be included and compared. As recapture data was used, data are only presented for fish $>12 \mathrm{~cm}$ which could be PIT tagged. Estimates are conservative as they do not take mortality into account.


Figure 6.11 Schematic diagram of dace movements between locations. Data are presented for physical recapture of fish to allow inclusion of sites without monitoring by PIT detector. No percentages were available for non-flowing ditches because these fish using these were too small to mark for recapture analysis.

### 6.4 Discussion

Dace used the entire mosaic of habitats available to them within the river system. Over the year, fish moved between habitats, depending on their requirements and activities. More immature than mature dace were found in the side-channels though the proportion varied seasonally and between side-channels. Lateral side-channels often contain large numbers of juvenile fish (Copp 1997, Garner 1996, Lusk et al. 2001) sheltering from the larger and more abundant predators of the main river channel (Borcherding et al. 2002). Though non-flowing ditches supported low numbers of dace in this study, those that were present were mostly small, $0+$ dace. This suggests their significance only as secondary nursery habitats, supporting few fish. Large, primary nursery areas were found in the main river in an oxbow lake out of the main flow (CEH, unpublished data).

Mixing of dace occurred to some extent between all locations, both between river and off-river habitats and also long distance movements along the main river. The peak in main river upstream movements happened in May, despite first upstream movements occurring in February. A study on the Yorkshire Derwent also found that over a full year of continuous PIT telemetry in a fishway entrance, over $80 \%$ of recorded upstream movement by dace ( $12-19 \mathrm{~cm}$ ) was between April and July (Lucas et al., 2000; M.C.Lucas, personal communication). It is likely that these movements in the River Frome were feeding migrations as they coincided with known increases in the abundance of invertebrates at that time (Clough 1998) and were made by substantial numbers of both mature and immature dace.

The majority of movement occurred between the tidal reach Wareham and upstream section at East Stoke. There were discrepancies in the recorded proportion of individuals between the different monitoring methods used, something that has been acknowledged as an issue with this type of study (Szacki 1999). Estimates of mixing ranged from between $5 \%$ to $50 \%$ of the population. This is in line with a previous study using passive marking techniques (visible implant alpha-numeric tags, visible implant elastomer and fin clipping) which found that $28 \%$ of individuals moved
between the two areas (Clough 1998). Substantial variation in estimates of large scale movements was evident. Differences between East Stoke main river ( $24 \%$ ) and sidechannel (5\%) PIT detection may reflect different proportions of fish from Wareham travelling further up the main river or into one of the 5 monitored side-channels. Physical recapture proportions were intermediate to the two PIT detection estimates (11.5\%). Radio tracking estimates of the proportion of dace visiting Wareham were by far the highest ( $67 \%$ in spring, but only $33 \%$ in autumn). These fish could be constantly monitored so more movements, and thus more long scale travel, were recorded as opposed to the more passive forms of monitoring. However, during the study few radio-tagged dace returned upstream following a downstream migration to Wareham. It is possible that fish moving downstream were unwell and unable to maintain position further upstream. However, many fish made localised upstream movements in Wareham and some were observed shoaling with other dace. Those that did not move following downstream travel to Wareham (implying death) were not included in analysis. Despite the high level of variability between estimates of population mixing, it is most likely that estimates made from radio-tracking were most accurate as this technique enabled a higher proportion of all movements, and particularly long-distance movements to be recorded (Szacki 1999).

While this study has addressed movement between habitat patches, it was over a short time period when considering population scale issues. However, it has gone some way towards establishing the complex spatial and temporal structure of a lowland river dace population. The study provides evidence of a quite structured population with some age structuring. There is also indication of the use of alternative strategies by adults with some mobile individuals travelling between habitats and other less mobile components of the population remaining in one area. A portion of the population utilised the side-channels, whilst others moved to the tidal reach and another group remained solely in the main river channel. Both linear and lateral connectivity were found to be contributory components to the life history diversity of the dace population.

This study concurs with other literature in that small changes in the distribution or accessibility of good and poor habitat may precipitate relatively large changes in species' viability (Doak 1995, Pulliam et al. 1992). If, as suggested, this is the case for
dace then destruction of lateral habitats or blockage of movement throughout the system with obstacles such as dams or weirs could have substantial impacts on population characteristics and persistence.

## Chapter 7

## Home Range Estimation within

Complex Restricted Environments:
Importance of Method Selection in
Detecting Seasonal Change

### 7.1 Preface

The work presented in this chapter has previously been submitted as a multi-authored paper to Ecological Applications. Those involved in the earlier work have been acknowledged in the acknowledgements section of this thesis. While in the original submitted thesis it was included as an appendix it was felt that it added much to the thesis, particularly as an introduction to the analysis techniques used in Chapter 8. For this reason it has been included in the body of the thesis as an additional chapter. Some of the pike telemetry data used in the manuscript was collected prior to the onset of this project.

### 7.2 Introduction

For many animal populations, systematic radio-tracking can provide data on spatial behavior and demography more rapidly and with less bias than more traditional methods such as visual observations or mark recapture (White and Garrott 1990). The advantages of such tracking are at their greatest for elusive species, including fish.

Quantitative analysis of home range size, shape and core structure has become fundamental to understanding the movements and behavior of animals (White and Garrott 1990, Fisher 2000, Crook et al. 2001, Broomhall et al. 2003, Fuller et al. 2005). The use of telemetry data to estimate home ranges is now common-place for studies of resource use (Aebischer et al. 1993, Terry et al. 2000, Johnson et al. 2004, Markus and Hall 2004), social interactions (Sliwa 2004, Sunde and Bolstad 2004), activity (Taylor and Skinner 2003), predation (Madsen and Shine 1996, Kraus and Rodel 2004, Yoder et al. 2004). Home ranges can be considered as the spatial expressions of the behaviors carried out by animals to survive and reproduce (Burt, 1943) that may change between seasons or year on year. Thus as spatial and temporal representations of an animal's requirements they are well suited to use investigating applied issues of animal ecology. Home range assessment has helped to answer a wide range of applied ecological questions relating to spatial behavior including reserve design (Wielgus 2002), conservation planning (Locke 1996, Johnson et al. 2004), habitat management analyses (Peach et al. 2004), assessing habitat suitability for reintroductions (Schad et al. 2002), controlling spread of disease (Woodroffe et al. 2006) and interactions of native with non-native species (Kenward and Hodder 1998).

There is a wide choice of possible methods for estimating home range. One group consists of ellipses or contours based on density distributions (Dalke and Sime 1938, Jennrich and Turner 1969, Worton 1989), which are derived from all the locations and hence tend to be influenced by outliers that represent the excursions of an individual. The second group creates polygons (Dalke and Sime 1938, Jennrich and Turner 1969, Worton 1989) that minimize linkage distances between pairs of locations. Polygon
methods include an excursive-sensitive minimum convex polygon around the peripheral locations, but can also exclude the influence of outliers by peeling or with cluster analysis to define range cores. The various density and linkage methods differ in their ability to (i) estimate home range outlines that conform to the observed locations; (ii) derive statistics describing the range structure; and (iii) achieve stable estimates with few locations (Harris et al. 1990, Kenward 1992, Robertson et al. 1998).

Methods of estimating home range use were primarily designed for species that move freely throughout the landscape (Dalke and Sime 1938, Calhoun \& Casby 1958, Dixon and Chapman 1980). However, many species are tightly associated with restricted and fragmented habitats such as woodland (Redpath 1995, Major and Gowing 2001) or confine their movements to largely linear pathways. Mammals, such as weasels (Mustela nivalis), inhabiting field edges in agricultural areas tend to remain confined to these linear corridors rather than venturing far into cultivated fields (MacDonald et al. 2004). Species such as river otter (Lutra canadensis) (Blundell et al. 2001), watervole (Arvicola terrestris) (Barreto and MacDonald 2000, Fedriani et al. 2002) and bald eagle (Haliaeetus leucocephalus) (Harmata and Montopoli 2001) often associate themselves with rivers and shorelines and many freshwater fish are limited to rivers. The use of standard home range methods in these cases may estimate outlines that include large areas of unusable habitat. This is particularly true where narrow corridors of usable habitat are highly convoluted, as occurs for braided river channels and meanders. The result is bias in size and home range structure statistics that indicate foraging movements.

Recently some investigators have attempted to develop new techniques to address this issue using simulated datasets (Burgman and Fox 2003, Matthiopoulos 2003, Getz and Wilmers 2004), however these techniques have not become widely available or used in ecological management. Few applied studies of species with linear patterns of movement maximize the information gained from the data by employing the most appropriate methods to quantify home ranges. Currently, while most work using linear home range estimation has focused on fish in rivers (Bridcut and Giller 1993, Baras 1997, Masters et al. 2002, Bahr and Shrimpton 2004), novel analysis has been applied mainly to river otters (Blundell et al. 2001, Sauer et al. 1999). Despite an increasing
number of telemetry studies on river fish species and related management issues, such as interactions between natives and non-natives, most either make no estimates of home range or calculate very basic measures, likely missing much of the ecological or behavioral information (Vokoun 2003). With potentially rapid alteration in habitats and movements of animals through climate change, coupled with strict targets for management through legislation such as the Water Framework Directive (European Commisssion 2000), the need to make best use of hard-won data from radio tracking is now greater than ever.

In this study, we test the effectiveness in a restricted environment of commonly used and widely available home range analysis techniques, including the clipping of range outlines to include only the water usable by an aquatic species. Pike (Esox lucius Linnaeus), like many fish species, are known to alter their behavior seasonally (Cook and Bergersen 1988). We postulate that size and internal structure of home ranges of river-dwelling pike, would differ between seasons to reflect differences in behavior, and that these differences would be shown more effectively by some analysis methods than by others.

### 7.3 Materials \& Methods

### 7.3.1 STUDY AREA

The study was conducted on the lower River Frome, Dorset, UK. The river is largely unmodified, with a meandering main channel in its lower reaches, but also has a manmade millstream within the study site. Most fish locations were collected within a 2 km stretch of river (UK national grid references SY867863 to SY882870), although fish were also tracked outside this stretch

### 7.3.2 RADIO TAGGING AND TRACKING

Data from 23 adult pike were used to test the range estimators. Radio tags (TW-5 tags, Biotrack Ltd., Wareham BH20 5AX, UK) were implanted into the body cavity of the pike ( $58-101 \mathrm{~cm} \mathrm{FL}$ ) as described in Beaumont et al. (2002). Fish locations were determined to within 1 m by triangulation from within 10 m on the river bank, using a Sika radio-receiver (Biotrack Ltd.) and a hand-held three-element Yagi antenna

Data were collected between May 2000 and September 2003, during March, June, September and December to provide home range estimates for each season. Fish were located at dawn, midday and dusk every day over a 13 day period, resulting in standard home range data sets of 39 locations per fish in each season. The timing of dawn, midday and dusk locations was selected as a result of a pilot study that demonstrated the diel activity of pike to be at dawn and dusk (Masters et al. 2002; Hodder et al. in press). In addition, fish were routinely located two or three times per week during the entire study period in order to prevent loss of mobile individuals.

One range was used from each fish per season. If data were collected from an individual in more than one year, the average range area for that season was used. In order to enable direct seasonal comparison of river channel habitat use ranges estimated during flooding were not included as these may have included floodplain habitat not available to fish in all seasons. In total, 16 records were available for spring, 12 for summer, 14 for autumn and 12 for winter (Table 7.1).

Table 7.1 Summary of home range tracks for each individual fish used for estimation of home range area.

|  | Fish ID Code |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12 | 13 | 14 | 15 | 16 | 17 | 18 | 19 | 20 | 21 | 22 | 23 |
| Spring |  | 1 |  | I | I |  |  | I | 1 | I | 1 | I | I | I | I | I | I |  |  | 1 | 1 |  | I |
| Summer | I |  | I | I | 1 | I |  |  | I |  | 1 | I |  |  | I | I | I |  |  |  |  | I |  |
| Autumn |  |  |  | 1 | I |  | 1 | 1 | I |  | 1 | I | I |  | 1 | I | I | I | I | I |  |  |  |
| Winter |  |  |  |  | I |  |  | I | I |  |  | I | I |  | 1 | I | 1 | I |  | I | I |  |  |

### 7.3.3 HOME RANGE AND MOVEMENT ANAL YSES

Four methods of area-based range analysis were used. All were methods readily available in commercial software (as reviewed by Larson (2001)) that are widely applied (Kernohan et al. 2001, Walls and Kenward 2001, Fuller et al. 2005) to give trade-offs between precision in conforming to peripheral and core locations (Table 7.2) and sample-size requirements (Robertson et al. 1998). All were estimated with RANGES 6F (Anatrack Ltd., Wareham. BH20 5AX, UK), with abbreviations and classification following Kenward et al (2001).

Table 7.2 Summary of home range estimation techniques and their advantages and disadvantages for use in restricted environments.

| Home range Technique | Sample size considerations | Periphery/core representation | \% locations included | Short name | Original reference |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Density methods |  |  |  |  |  |
| Ellipses | Size can be stable with as few as 1215 locations (well separated in time). | Outline conforms poorly to locations and is sensitive to outlying locations. | 95\% | E95 | $\begin{aligned} & \text { (Worton } \\ & \text { 1989) } \end{aligned}$ |
| Kernels | Size can be stable with as few as 1520 locations. | Contours conform to multinuclear data but not to abrupt change in location density and are affected by outlying locations. | $\begin{aligned} & 95 \% \\ & 50 \% \end{aligned}$ | $\begin{aligned} & \text { K95 } \\ & \text { K50 } \end{aligned}$ | (Dalke and Sime 1938) |
| Link distance methods |  |  |  |  |  |
| Minimum convex polygon | Sample-sizedependent but often stable with $30+$ locations. | Single polygon conforms only to peripheral locations. | 100\% | X100 | $\begin{aligned} & \text { (Kenward } \\ & 1987 \text { ) } \end{aligned}$ |
| Clusters (core influenced) | Requires $30+$ locations for stability. | Polygons fit core location patches and provide internal structure statistics. | 85\% | CX85 | (Kenward et al. 2001) |

### 7.3.4 DENSITY METHODS

Jennrich-Turner ellipses, estimated to include $95 \%$ of the density distribution ( $\mathrm{E}_{95}$ ), are least precise in conforming to locations, but can give stable estimates of total range area with 12-15 locations that have low spatio-temporal correlation (Kenward, 2001). Kernel contours containing $95 \%$ of the estimated density distribution ( $\mathrm{K}_{95}$ ) were estimated using least squares cross-validation with a fixed smoothing multiplier of 1 and a $40 \times 40$ matrix as a total area measure with more flexible conformation. A widely-used estimator of core size (Fuller et al. 2005) is the kernel contour $\mathrm{K}_{50}$, containing $50 \%$ of density distribution (Table 7.2).

### 7.3.5 LINK DISTANCE METHODS

As an estimate of total range size, a single convex polygon around $100 \%$ of the locations ( $\mathrm{X}_{100}$ ) provides greatest comparability with other publications (Harris et al. 1990), but its conformation to the range periphery requires large sample sizes to
minimize sample-size-dependence. Similarly, core polygons derived by cluster analysis require large samples of locations for stability (Robertson et al. 1998). Cluster polygons were estimated around $85 \%$ of the locations in each range $\left(\mathrm{Cx}_{85}\right)$ as this excluded peripheral locations for $95 \%$ of the ranges (Kenward et al. 2001). In addition to area and number of cores, cluster analysis provides an index of range patchiness: the sum of areas of the separate cores as a fraction of the area of a single convex polygon around all the clusters decreases from 1, if locations are all in one nucleus, to a small fraction if nuclei are far apart (Harris et al. 1990).

### 7.3.6 AREAS AND DISTANCES IN RESTRICTED ENVIRONMENTS

A problem with analyzing movements in restricted environments is that measurement of distances by straight lines may underestimate travel distances for example, in rivers, by cutting across curves, islands and junctions with tributaries. RANGES 6F avoids this problem by estimating inter-location distances along a river midline for travel estimates and the nearest-neighbor distances used in cluster analysis. Similar functions may be performed in widely used software such as ArcView. A more pervasive problem with analyses in rivers is that home range outlines often extend beyond the river banks (Figure 7.1), so that inaccuracy (over-estimation of area) may mask biologically meaningful relationships. Cluster analysis minimizes this error (Hodder et al. in press) but cannot entirely eliminate it. Outlines for all six range area estimators were therefore clipped to a map of the river banks that was imported to RANGES 6F from ArcView GIS 3.2 (ESRI Inc, Redlands, California). An excursionexcluded range span was calculated as the summed midline lengths of $85 \%$ cluster cores. The clipped $\mathrm{X}_{100}$ measures maximum river area covered by excursions and is equivalent to maximum linear distance at constant river width.


Figure 7.1 Example of out-of-river error from locations of the same fish fitted with (a) $85 \%$ clusters and (b) $95 \%$ kernels (solid line) and $50 \%$ kernels (dotted line).

### 7.3.7 COMPARING ESTIMATORS

Two approaches were used to compare the performance of different spatial estimators. 'Discrimination testing' compared the ability of different estimators (of area, span, patchiness and distance between consecutive locations) to detect variation between seasons. When testing for seasonal variation with six estimators of range area in the same data set, there was an enhanced probability of Type I error. Therefore a

Bonferroni correction was applied. 'Error testing' compared, for each method, the extent of unusable environment (out-of-river) error, its coefficient of variance and its correlation with range span (as an independent estimate of range size). The index of out-of-river error was total range area divided by clipped area.

Seasonal variation in range areas, patchiness and distances between consecutive locations was investigated with a global test across all seasons, followed by pair-wise comparison of seasons. A Friedman test of overall seasonal variation in range size was used for seven fish that contributed a range in all four seasons, followed by Mann-Whitney U-tests of range size differences between pairs of seasons for all fish. Wilcoxon tests of estimators for each season paired across the same fish gave similar results, albeit at lower significance levels due to reduced sample sizes.

### 7.4 Results

Friedman tests demonstrated overall differences in area between seasons only when calculated with $\mathrm{Cx}_{85}$ and $\mathrm{Cx}_{85}$ clipped ( $P<0.01$ ). However, range area was found also to differ significantly between pairs of seasons for $\mathrm{K}_{95}$ and $\mathrm{K}_{50}$ using Mann-Whitney U-tests (Figure 7.2). All methods estimated the smallest ranges in autumn. For the kernel estimators, summer range areas were significantly larger ( $P<0.001$ ) than autumn range areas for both kernel and cluster estimators. These seasonal patterns remained the same after clipping. There were no significant seasonal differences in range span (Figure 7.2g).


Figure 7.2 Mean home range area (with standard deviation bars) estimated with nonclipped methods (black) and clipping (grey) for three excursion-sensitive range estimation techniques; (a) $100 \%$ polygons, (b) $95 \%$ ellipses and (c) $95 \%$ kernels, followed by two coring methods; (d) $85 \%$ clusters and (e) $50 \%$ kernels with (f) excursion-excluded range span. Letters, indicate Bonferroni corrected significant differences between the seasons labeled; Spring - P, Summer - S, Autumn - A and Winter - W. Note different scales for the top, middle and bottom plots.

Excursion sensitive range estimators $\mathrm{X}_{100}, \mathrm{E}_{95}$ and $\mathrm{K}_{95}$ gave larger range areas than core estimators $\mathrm{Cx}_{85}$ and $\mathrm{K}_{95}$, by a factor of ten (Figure 7.2). However, clipping increased comparability between range cores and excursion-sensitive methods.

Ellipses produced the highest proportion of out-of-river error within each range (Fig. 3a). However the two kernel estimators, particularly $\mathrm{K}_{50}$, had the highest variability in out-of-river error created during range estimation (Figure 7.3b). Clusters ( $\mathrm{C}_{85}$ ) gave lowest and least variable errors.


Figure 7.3 (a) Index of out-of-river error, as the proportion of the clipped range area in the total unclipped range area, and (b) its coefficient of variation.

Range span provides an index for investigating the extent to which out-of-river error depends on range size, because the greater the span along a curvaceous river, the more meanders the outline is likely to encompass. The influence of range size on the amount of out-of-river error included into estimation of range size varied greatly between different estimators from $8 \%$ to $73 \%$ (Figure 7.4). Error of core range estimators was less correlated with range span (maximum $R^{2}=44 \%$ ) than for excursion sensitive kernels and polygons (maximum $R^{2}=73 \%$ ) but not ellipses ( $R^{2}=$ $8 \%$ ).


Figure 7.4 Regression of error index on range span for (a) $100 \%$ polygons, (b) $95 \%$ ellipses, (c) $95 \%$ kernels, (d) $50 \%$ kernels and (e) $85 \%$ clusters.

Overall distances traveled differed between seasons in a Friedman test ( $P<0.05$ ). Distances moved between locations in spring were significantly longer than in autumn and winter $(P<0.01)$. Distances were highly variable in summer, because some pike appeared to exhibit less excursive activity than others, and therefore only showed tendencies ( $0.05<P<0.01$ ) to differ from spring and winter (Figure 7.5).


Figure 7.5 Boxes show median, $25^{\text {th }}$ and $75^{\text {th }}$ percentiles, with whiskers at $10^{\text {th }}$ and $90^{\text {th }}$ percentiles (and outliers as dots) for mean midline distances between consecutive locations ( $\log _{10}$ transformed). . Upper case letters indicate differences significant at the $1 \%$ level, with lower case letters for the $5 \%$ level.

Variation in home range patchiness indicated differences in core range structure between seasons (Figure 7.6). Home ranges were significantly less patchy in autumn than summer ( $P<0.01$ ), when fish activity was less tightly focused in the core areas.


Figure 7.6 Home range patchiness of $85 \%$ clusters. Low scores indicate the patchiest ranges. Upper case letters indicate differences significant at the $1 \%$ level, with lower case for the $5 \%$ level.

### 7.5 Discussion

The area that is repeatedly traversed by an animal tends not to be used either in a uniform manner, or as defined by a normal distribution (White and Garrott 1990). Typically, there are one or more patches of intensive use surrounded by an area used for excursions (Kenward 1987, 2001) or for patrolling territorial boundaries (Kruuk 1978). An ability to identify the number of high-use cores, and their placement with regard to habitats and other animals, can reveal much more about foraging and sociality than can mere range size (Fuller et al. 2005) and is crucial to many management and conservation applications. Relative size and usage of peripheral areas can indicate the effort directed to exploratory behavior, mating and territoriality, as well as quantifying the habitats and neighbors available to an individual (Allouche et al. 1999).

In this study, cores defined by cluster analysis showed the strongest variation between seasons, the lowest and least variable inclusion of unusable habitat, the second lowest correlation with range span, and also showed seasonal variation in core fragmentation. The least variation due to estimation processes might be expected in cluster cores, because the exclusion of outliers eliminates a small number of locations with very variable placement due to behavior that may differ from core activities (Kenward et al. 2001). Areas of clusters are totally uninfluenced by excluded outliers. In contrast density estimators include effects of all locations in every calculation, although distance functions of harmonic mean estimators (Dixon and Chapman 1980) minimize this effect (Kenward et al. 2001) and peripheral polygons are defined by outliers. With least methodological variance, cluster areas should be best able to show biological variance, in this case due to seasonality. Moreover, their tight definition by polygons minimizes the expansion into unusable habitat that results from smoothing implicit in density estimation. A similar restriction to local nearest-neighbor linkages may give equally good performance from the convex-hull approach of (Getz and Wilmers 2004) if an algorithm becomes more generally available.

For estimating a home-range periphery, a $\mathrm{K}_{95}$ contour gave better detection of seasonal variation than either an $E_{95}$ ellipse or a peripheral convex polygon ( $\mathrm{X}_{100}$ ).

This resulted despite the peripheral polygon having least (and relatively invariable) out-of-river error, probably because only the kernel contours could define multinuclear range outlines. The moderate sensitivity to outliers of $\mathrm{K}_{95}$, as opposed to the high sensitivity of $\mathrm{E}_{95}$ and total dependence of $\mathrm{X}_{100}$ may also have been advantageous.

The extent of unusable-environment error would be expected to depend both on the home range estimator and the behavior of each animal. Expansive estimators and far ranging individuals will tend to cause more unused area to be included in the home range calculation than those that occupy smaller areas. Ellipse estimates tended to be very large (Figure 7.2), such that error was large and relatively invariable (Figure 7.3), which apparently minimized scope for correlation with range span (Figure 7.4). Yet in no cases was there a tendency for seasonal variation to be better defined by clipping of unusable habitat. This poor result from an intuitively attractive analysis approach was surprising. If confirmed for other species, this finding implies that unclipped estimates of range area might serve for fish in both linear and non-linear (e.g. lakes) habitats. Indeed, unclipped kernel and cluster analyses showed significant range expansion of home ranges when the river Frome flooded adjacent fields (Hodder et al. in press). However, clipping might be more necessary for habitat analyses and when range overlaps are used to investigate sociality.

Seasonal differences in range size reflected variation partly in linear distances traveled and partly in the patchiness of range structure. Range span alone however, was not informative in terms of pike behavior. It did not show the significant differences in space use between seasons demonstrated by area based range analysis methods in this study or the spring peak and autumn dip in movement described in Cook and Bergersen's (1988) radio telemetry study in a Colorado reservoir. A high level of range span variation between individuals hindered detection of statistically significant seasonal trends, as it did for peripheral convex polygon areas, with which range span correlated strongly (Figure 7.4). Differences in river width between seasons might also have contributed to area-based methods having greater statistical significance than range span in seasonal differences.

Therefore, although range span has traditionally been the most common analysis technique of animals in restricted habitats (Powell 2000), linear distance measures of movement should be complemented by estimates of home range. For pike in the River Frome, this showed that all range area estimators were smallest in autumn (Figure 7.2), at which time cores were least fragmented (Figure 7.6) and inter-location movements smallest (Figure 7.5). Cluster analysis showed that core ranges were largest and most fragmented in summer, when inter-location distances were at their most variable. However, the largest peripheral range areas, with statistically significant differences from autumn for the most moderately excursion-sensitive methods ( $\mathrm{K}_{95}$ ) were in spring, when inter-location distances were at their largest.

### 7.5.1 APPLICATION

In a review of how twelve estimators of home range area scored subjectively in seven performance criteria, Kernohan et al. (2001) rated kernels first and clusters second. Our objective comparison of four methods in a complex restricted environment concurs with this result for estimating peripheral outlines, but we rate cluster polygons as the more appropriate for estimating range cores because of their greater ability to eliminate influence of outlying locations and their generation of structure statistics. A multifaceted approach with more than one technique enables the most thorough interrogation of the data and also helps to overcome some of the drawbacks inherent in the different methods of home range analysis. Continued estimation of range span will provide an index that remains comparable with previous studies in restricted environments, such as rivers and hedgerows.

Maximizing data interrogation and comparability of results between studies should aid the application of findings to management situations For example, improved study and analysis of space-use interactions between native and non-native fish species in river systems could contribute usefully to understanding mechanisms of observed declines in native species (e.g. Marchetti et al. 2004). Furthermore, improved data interrogation and comparability of space-use statistics in restricted habitats would enhance their utility for meta-analyses that can give broad ecological insights. Current progress in the analysis of telemetry data (Börger et al. 2006), should permit even greater exploitation of available information and provision of resources for managers.

Management may also require predictive modeling, for which it is important to minimize sources of error and bias.

## Chapter 8

## Population Structure and Side-

Channel Use: A Case Study of a
Limnophilic Species

### 8.1 Introduction

Pike are a limnophilic predator typically found in lakes, slow-flowing rivers and canals with shallow to moderately deep waters and a high density of vegetation (Raat 1988). Pike are found throughout Britain, except Northern Scotland and occur throughout Europe, Asia and North America (Davies et al. 2004). Pike exhibit sexual size dimorphism with females being larger than males (Raat 1988). Females attain sizes of up to 130 cm or 19 kg in Britain (Davies et al. 2004). They are thought to be solitary but not territorial, with some overlap in their range areas (Cowx 2001). Pike are opportunistic predators, with their food type varying according to abundance and seasonal availability (Chapman et al. 1989). They feed on invertebrates, fish and other vertebrates and are known to be strongly cannibalistic, with cannibalism forming a large part of the predation pressure on young pike in some populations (Mann 1982).

Pike spawn in shallow, sheltered, vegetated water between March and May (Cowx 2001). The onset of spawning is controlled by photoperiod and increasing temperature (Lucas 1992). There is some evidence that pike migrate to the same spawning ground each year (Frost and Kipling 1967, Bregazzi and Kennedy 1980). Spawning takes place in daylight with between one and three males accompanying a single female (Raat 1988). Eggs are shed indiscriminately over vegetation over a distance of 50 100m (Raat 1988). Female fecundity is approximately $15-30$ eggs per gram of body weight fecundity varies widely (Frost and Kipling 1967, Mann 1976b).

Fish populations are often structured by the environmental pressures that they face. Thus habitat structure and diversity influences the spatial distribution and ecology of fish populations. Different life history stages often use different habitats. For example, sofie (Chondrastoma toxostoma) spawn in river tributaries. Juveniles move out into the main river and do not return until they spawn, young mature sofie migrate from the main channel to spawn in the tributaries and then return to the main channel and older sofie remain in the tributaries as residents (Gozlan 1998). Population spatial structure is not just driven by different age groups, but individuals within a population may adopt different space use strategies. This may mean use of segregated habitats
such as the main channel and side-channels or it may mean use of different habitats within the main river channel. Habitats differ in their quality for the individuals using them. Individuals in poorer quality habitats exhibit slower growth and lower condition. For instance, some bream are resident floodplain lakes of the Lower Rhine, while other individuals, which are smaller and in lower condition, remain in the river or make trips to the floodplain lakes (Grift et al. 2001).

The floodplain of the River Frome provides a varied network of off-river habitats available to fish. Each of these lateral habitats supports a community of fishes (Chapter 4) and provides different functional uses for many fishes throughout the year (Chapter 5). Dace used a wide range of habitats available on the Frome, but as a rheophilic species largely utilised only flowing side-channels (Chapter 6). As a limnophilic species, pike could be expected use a different range of the side-channels available on the River Frome.

Pike are also a top predator and unlike dace which batch spawn, pike undergo mate selection with one female spawning with between one to three males (Raat 1988). In pike, unlike many fish species, there are no secondary sexual characteristics for females to differentiate between males (Wootton 1990). Instead, as in many other species, a likely selective trait for spawning pike is the size of the fish (Downhower and Brown 1980, Maekawa et al. 1996, Rowland 1989). Thus small male pike are likely to compete less well for spawning than larger males.

This chapter addresses the hypothesis that a limnophilic, top predator population will be spatially structured as a result of habitat variability. This may be applied to use of different areas within the main river channel and also differential use of main river and side-channel habitats. Individual's home ranges will be compared with respect to their growth characteristics to investigate population structure using the common definition of a home range as "the area traversed by an individual in its normal activities of food gathering, mating and caring for young" (Burt 1943). It is likely that the size and quality (availability of feeding and sheltering resources) of an animal's home range will affect the excursions made outside its everyday area (Powell 2000). So the relationship between home range size and an individual's vagility, or inherent tendency to move, was investigated in pike as it has been documented previously in
both birds and mammals (Bowman 2003, Bowman et al. 2002). Indeed, Bowman (2002; 2003) was able to predict vagility from home range size. Bowman et al (2002) argued that differences in behaviour and physiology between species affect vagility in mammals, such that home range size and dispersal distance covary proportionately. These previous studies investigated this relationship between species, while this investigation focussed on differences between individuals of a single species. The relationship between an individual pike's growth rate and it's presence in a sidechannel was also investigated.

### 8.2 Materials \& Methods

### 8.2.1 SAMPLING

Seven side-channels (Chapter 2) were sampled with electric fishing four times a year during 2003-2005. Five side-channels were monitored with PIT detectors during 2004-2005. Adult pike were radio-tracked at dawn, midday and dusk over a 13-day period four times per year to determine their seasonal home range (2002-2005) and also 2-3 times per week to monitor excursive activity and maintain contact (20022005). See Chapter 3 for further detail on the procedures used.

### 8.2.2 ANALYSIS

The relationship between seasonal home range size and annual excursions was investigated. The sample size was limited to 14 pike for which both seasonal home range and annual excursions were available. Adult pike (mean FL $\pm$ S.D. $69.5 \pm 12 \mathrm{~cm}$, range $52-93 \mathrm{~cm}$ ) were located in December, March, June and September at dawn, midday and dusk every day over a 13 day period, resulting in standard home range data sets of 39 locations per fish. In addition, fish were routinely located two or three times per week during the entire 3.5 year study period.

Annual home range area was calculated using cluster analysis, objectively excluding all excursive activity (see example in Figure 8.1). An iterative process excluded the location with the most extreme linkage distance if it was beyond $1 \%$ of the distribution estimated by the remainder, and repeated this process until all distances were within the $1 \%$ alpha-level on a normal distribution. Ninety nine percent cluster analysis (Kenward 1987) was used to ensure that only the clearest excursions were excluded. The number of excursions and mean excursion distance (m) outside this home range were calculated along a river midline using the three times weekly tracking data over a full year (Figure 8.1).


Figure 8.1 Example of the calculation of annual and seasonal home range and of the calculation of excursion distance.

Ninety five percent cluster analysis (Kenward 1987) was used to estimate the seasonal home range areas as it expresses the core structure of a range once excursive movements are excluded (Hodder et al. 1998). Core ranges estimated from the seasonal 13-day tracks for each pike were then compared to their annual number of excursions and mean excursion distance (Figure 8.1).

Home range size for all four seasons was regressed against both mean excursion distance and annual number of excursions. All variables were on a $\log _{10}$ scale to correct for non-normality of the data. The potential for predicting mean excursion distance from seasonal home range area was then investigated. An isometric relationship between the linear dimension of home range (square root of home range size) and median dispersal distance exists in mammals (Bowman et al. 2002) of the form:

$$
\text { median dispersal distance }=7 \sqrt{H R}
$$

Where:

$$
\mathrm{HR}=\text { Home range }
$$

Vagility is defined as the tendency of animals to move. While there are clear differences between excursions (long distance movements where the individual returns to its original location) and dispersal (long distance movements away from a
location with no return) they can both be used as measures of vagility. The equation above was used to test the predictability of individual pike vagility from home range size. For this study the prediction was tested against mean excursion distance rather than median dispersal distance as used by Bowman (2002) because measures of pike dispersal were not available. Residuals of the $\log _{10}$ (home range area) vs. $\log _{10}$ (body size) regression and the $\log _{10}$ (mean excursion distance) vs. $\log _{10}$ (body size) regression were used to assess, by regression, the relationship between home range area and mean excursion distance independently of body size.

In order to investigate this further seasonal home ranges were obtained between May and August for 51 pike, (this included the 14 pike used for the vagility investigation.) (mean FL $\pm$ S.D. $51.4 \pm 25.5 \mathrm{~cm}$, range $9.2-105 \mathrm{~cm}$ ). Between 30 and 39 fixes were included in all home range estimates. In most cases 39 fixes were used (Hodder et al. in press) but for some of the smallest pike fewer were available. Home range size stabilised with fewer locations for small pike. So estimates made from 30-35 fixes for small pike were comparable to 39 -fix home ranges estimated for large pike. Sex was known for 36 of the 51 pike radio-tracked (female $n=17$; male $n=19$ ) and growth data was available for 45 .

Von Bertalanffy growth coefficients $\mathrm{L}_{\infty}$ and K were calculated for each pike (see Chapter 3 and Chapter 5 for further details). Except where specified, $85 \%$ clusters were used appropriately as home range estimates (see Chapter 7). The following transformations were applied to normalize the data. It was not possible to totally normalise $\mathrm{L}_{\infty}$ but after transformation the variances were very much reduced and it tended towards normality ( $P>0.05$ ).

Transformations used were:

| Cluster area | $\log _{10}($ cluster area $* 10000+1)$ |
| :--- | :--- |
| $\mathrm{L}_{\infty}$ | $\log _{10}\left(\mathrm{~L}_{\infty}\right)$ |
| K | $\log _{10}\left(\mathrm{~K}^{*} 10000+1\right)$ |

Values were multiplied by 10000 (before adding 1) when there were zeros and many low values in order to transform the data while maintaining the proportions between the data points.

Seasons were defined according to the pike spawning season, so that all spawning months were described together as Spring. Thus:

| Spring | March - May |
| :--- | :--- |
| Summer | June - August |
| Autumn | September - November |
| Winter | December - February |

The condition coefficient of individual fish was calculated with the formula:

$$
\mathrm{C}=10^{5}\left(\mathrm{~W} / \mathrm{L}^{3}\right)
$$

Where:
C $=$ condition coefficient
W = weight
$L=$ fork length

No pike were caught in Goldsacs stream and Millstream functionality differed from that of the other streams utilised by pike (see Chapter 5 for further detail). Thus for this chapter only (Figures 8.7 and 8.8 and Tables 8.2 and 8.3), side-channels Railway, Rushton, Luckford, Flood Relief and Holme Bridge were grouped as 'ditches' for analyses into condition and growth factors, with Millstream and also the main river channel used for comparison.

Statistical analyses were carried out in Minitab 14. Graphs were made in SigmaPlot 2000.

### 8.3 Results

### 8.3.1 PIKE SPATIAL BEHAVIOUR

Winter and spring home range area of the 14 adult pike studied were strongly positively correlated with mean annual excursion distance ( $R^{2}=0.91, P<0.0001$ and $R^{2}=0.74, P=0.001$ respectively , while summer and autumn were not ( $R^{2}=0.37, P$ $=0.15$ and $R^{2}=0.08, P=0.42$ respectively) (Figure 8.2). Seasonal home range areas
were not correlated to the number of excursions made throughout the year $\left(R^{2}<0.22\right.$, $P>0.18$ in all cases).

No significant differences were found between observed and predicted mean excursion distances (Paired $t$-test: $P>0.10$ in all cases). So, mean excursion distance of pike was successfully predicted from winter and spring home range area with Bowman's (2002) equation developed for mammals, despite the small sample size of pike.


Figure 8.2 Relationship between residuals of seasonal cluster home range areas and residuals of mean annual excursion distance (i.e. after effects of body size were removed) demonstrating the seasonal relationships between home range area and excursion distance.

Pike length was not correlated with either variation in seasonal home range area ( $R^{2}<$ $0.01, P>0.34$ in all cases) or mean excursion distance ( $R^{2}=0.00, P=0.925$ ). The relationship between seasonal home range area and mean excursion distance showed no change when the effect of body size was removed (Fig. 8.2), showing that body
size did not influence the relationship between home range size and mean excursion distance. No significant difference was found between males $(n=6)$ and females ( $n=$ 9) for either home range size or mean excursion distance ( $t$-test; $P>0.29$ in all cases), although sample sizes were low. The relationship between individual growth variables and home range size were investigated for a larger sample of pike (including those that did not have information on annual excursion distance).

On a large scale, ranging nearly the full size range of pike, home range size increased with the size and age of pike (Figures 8.3 and 8.4). Home range increased at a faster rate with size increases in smaller/younger pike, slowing to reach a plateau in fish over 80 cm or 9 years of age.


Figure 8.3 Cluster home range size according to individual fish size. Home range is expressed as $\log _{10}$ (Home range size * 10000).


Figure 8.4 Cluster home range size according to individual fish age. Home range is expressed as $\log _{10}$ (Home range size * 10000).

Home range size was correlated with growth throughout a pike's life time (Figure 8.5). This was illustrated by both the growth constant K and $\mathrm{L}_{\infty}$ being positively correlated with cluster home range area ( K is inversely proportional to growth rate, hence the apparent negative correlation with home range size in Figure 8.5). The theoretical maximum size of an individual $\left(\mathrm{L}_{\infty}\right)$ gave a stronger correlation with home range size than growth coefficient $K$.


Figure 8.5 Relationship between growth coefficients $L_{\infty}$ and $K$ (expressed as $\log _{10}$ of the variables) with cluster home range size. Home range is expressed as $\log _{10}$ (Cluster * 10000).

Male and female pike growth rate did not differ significantly ( $t$-test, $\mathrm{K} ; P=0.708, \mathrm{~L}_{\infty}$; $P=0.456$ ). Male cluster home ranges were significantly smaller than female home ranges ( $t$-test; $P=0.002$ ). Male home ranges were also more variable in size than females' as illustrated by a higher coefficient of variation (Table 8.1). However, males were generally smaller than females (male mean fork length $=51.5 \mathrm{~cm}$, S.D. $=19.7$;
female mean fork length $=70.23 \mathrm{~cm}$, S.D. $=17.84$ ). To account for this, size was added as a covariate in the ANCOVA when testing the correlation differences between fish size and home range size. Male home range size was found to be more strongly related to fish size than female ( $A N C O V A$; Clusters $R^{2}=0.57, P=0.023$ ). This was not the case for age ( $A N C O V A$; Clusters $R^{2}=0.40, P=0.19$ ).

Table 8.1 Home range characteristics for male and female pike

|  | Male | Female |
| :--- | :--- | :--- |
|  |  |  |
| Mean home range size (Ha) | 0.0234 | 0.0724 |
| Coefficient of variation | $24.03 \%$ | $13.84 \%$ |

Incorporation of pike life history parameters (i.e. growth, sex, size and age) by stepwise multiple regression against cluster home range size found that the best (explaining most variance) combination was size, $\mathrm{L}_{\infty}$ and sex ( $R^{2}=0.66, P<0.001$; size $P=0.001 ; \mathrm{L}_{\infty} P=0.005$; sex $P=0.069$ ).

### 8.3.2 DIFFERENTIAL SIDE-CHANNEL USE

Both mean fork length and mean weight differed according to the location in which the pike was caught between all locations (Figure 8.6a and b). The condition coefficient was significantly lower for fish caught in ditches (excluding Millstream, see section 6.1.2) than those in the river (Figure 8.6c) but no significant differences were found between Millstream and either the ditches or main river. Likewise individual growth indicators varied according to catch location. $L_{\infty}$ was lower in ditch pike than Millstream and river pike (Figure 8.6d). K indicated lower growth in fish caught in ditches than the main river (Figure 8.6e). Pike caught in the main river and Millstream exhibited similar growth and condition characteristics. Pike caught in the ditches attained a smaller size at age than those in the main river or Millstream (Figure 8.7). Indeed it took pike in the ditches two years longer to reach a given length than those in the main river or Millstream.


Figure 8.6 Mean (a) length, (b) weight, (c) condition coefficient, (d) $\mathrm{L}_{\infty}$ and (e) growth coefficient K of fish in the ditches, Millstream and main river with standard deviation error bars. Asterisks indicate significant differences at $\alpha=0.001$ level (ANOVA; Tukey test, $F>13.23$ in all cases).


Figure 8.7 Back-calculated fork length at age for pike caught in the side-channels, Millstream and main river.

Age, fork length and growth were all found to be good predictors of pike presence in side-channels using binary logistic regression (Table 8.2). Fork length strongly predicted presence in side-channels (based on electric fishing capture data), with decreasing size increasing the likelihood (odds) of being caught in a side-channel (as the odds ratio decreases or increases from 1 the likelihood of the influence of the predictor increases). Likewise, younger pike were more likely to be caught in sidechannels. Odds ratios suggested that the probability of faster growing pike being caught in a side-channel was only 0.39 or less than that of capture elsewhere. A small $\mathrm{L}_{\infty}$ was strongly representative of side-channel caught pike. In the case of the growth constant K , as K is inverse to growth rate the odds ratios were also inverse. In most cases the coefficients were found to differ significantly from 0 and the prediction of most models was high. Growth constant K was a weaker predictor than age, fork length or $\mathrm{L}_{\infty}$.

Table 8.2 Results of binary logistic regression modelling of growth on presence or absence from ditch habitats. The estimated coefficient is the likelihood of success (presence in ditch) according to the predictor. The odds ratio can be interpreted as the odds of slow growers being present in the ditch versus the odds of fast growers being present. Significant $P$-values (in bold italics) indicate that there is sufficient evidence that the coefficients are not zero. Concordance is a comparative measure of model prediction allowing comparison between models.

| Predictor | Coefficient | Odds ratio | P-value | Concordance |
| :--- | :--- | :--- | :--- | :--- |
| Age | -0.93 | 0.39 | $<\mathbf{0 . 0 0 1}$ | 78.8 |
| Fork length | -6.16 | 0.00 | $<\mathbf{0 . 0 0 1}$ | 84.7 |
| L $\infty$ | -5.35 | 0.00 | $<\mathbf{0 . 0 0 1}$ | 82.3 |
| K | 0.86 | 2.37 | 0.072 | 55.5 |

### 8.4 Discussion

Like dace (Chapter 6), pike used a wide range of habitats available to them within the river system. Over the seasonal cycle, pike move between different habitats, for feeding, spawning or sheltering (Chapter 5). Individuals have distinct spatial distributions and habitat use along the Frome, either in terms of the time spent in lateral channels or of the size and structure of their home range.

Vagility was related to pike winter and spring home range size. This is similar to bird and mammal dispersal (included in Bowman et al's (2002) study were gray wolf, Canis lupus, Eastern chipmunk, Tamias striatus, cougar, Felis concolor and meadow vole, Microtus pennsylvanicus) for which home range size is a reliable predictor of tendency to move (Bowman et al. 2002, Walls et al. 1999). However no relationship was found with the number of excursions. This suggests that while all pike travel outside their home range as frequently, some travel further than others. The strong relation between seasonal home range size and vagility indicates that there are common factors dictating the tendency of an individual to move.

The relationship between home range size and vagility for birds is influenced, at least in part, by social pressures (Walls et al. 1999). Growth rate was faster in pike with larger home ranges, suggesting that these individuals were able to acquire more energy than was expended while using the greater area. Social hierarchy has been noted in pike, with dominant and subdominant individuals (Hawkins et al. 2005). It is possible that fast growing individuals gain advantage over conspecifics early in life and become dominant members of the population. As pike are not territorial, but intimidation between individuals does occur (Eklov and Diehl 1994, Nilsson et al. 2000, Raat 1988), it is possible that faster growing pike gain access to more areas (and thus have a larger home range) through intimidation and as a result encounter more feeding opportunities.

The lack of correlation between mean excursion distance and summer or autumn home range size indicates a difference in activities between winter/spring and summer/autumn periods. In temperate climates highest pike competition for resources occurs during winter and spring. In winter prey productivity in the river is low (Chapman 1968) and availability of cover for these ambush predators is reduced due to die back of vegetation (Dawson et al. 1991). In spring competition increases during the spawning season for access to spawning grounds and mate selection (Billard 1996) with larger pike likely to be more successful spawners. Increased competition between pike during winter/spring is likely to increase the mobility of pike, with more dominant individuals able to travel between more areas (Sutherland 1996). This is supported by the highly significant correlation between home range size and an individuals' tendency to move during these highly competitive seasons.

Growth rate was more strongly linked to male than to female pike home range size suggesting that behaviour differing between the sexes may be the root of the fish size to home range size relationship. It is possible that size may a selective trait for mate selection in pike with bigger males gaining more spawning opportunities. This would suggest that pike which grow faster are better reproductive competitors. Male home range size was also more variable than that of females, suggesting that inter-male competition was more significant in pike than inter-female competition. Inter-male competition is often higher in species where female selection determines male reproductive success (Krebs and Davies 1997). These findings may suggest a
reproductive basis to the differences in home range characteristics seen in male and female pike. In the case of inter-male reproductive competition, male pike would likely employ reproductive strategies to maximise their individual spawning success. Male pike arrived earlier onto spawning grounds than females (Chapter 5) which could be suggestive of protandrous behaviour; the early arrival of males onto the spawning grounds to wait for females.

Home range size correlated well with both the age and size of an individual. At the start of life, home range size increased quickly with respect to age/size. This increase in home range size slowed and eventually reached a plateau at about 80 cm or 9 years old. This change in home range size with time closely follows, and probably results from, the pike growth curve. In the exponential phase of growth early in life pike need to increase their home range size quickly as both their body size and energy requirements increase. This allometric relationship has previously been shown in birds (Bowman 2003), mammals (Bowman et al. 2002, Mysterud et al. 2001, Swihart 1986), lizards (Perry and Garland 2002) and other fish (Minns 1995) but has always considered cross-species relationships and not within-species relationships as was the case in this study.

Pike resident in lateral habitats were smaller, slower growing and in poorer condition. Pike caught in side-channels took a full two years longer to reach the same size as pike from the main channel. Indeed the relationship was strong enough that growth characteristics could be used to predict presence or absence in side-channels (Table 8.3) and also described $65 \%$ of the individual variation in the time spent in sidechannels (Chapter 5). There was less difference in these growth variables between the main river and Millstream probably because in terms of pike habitat, the millstream was closer to being a 'small river' than a slow flowing side-channel.

Like the River Frome dace population, the pike population was structured by the surrounding environment. Side-channel habitats were used by pike for spawning, but were particularly important for slow-growing males which arrived early in the spawning season. The main channel population was also behaviourally structured with faster-growing pike accessing larger areas of the river.

## Chapter 9

General Discussion and Conclusions

During the history of the River Frome human alterations have been made which changed the form of this semi-natural system. Changing land-use such as cultivation on the floodplain and bank stabilising has affected the river system and the fish populations it supports. Fish, particularly dace, abundance has declined over the last 15 years (observations from River Frome angler CPUE) with no known cause, though it is likely that human activities have impacted on the populations. Increased water abstraction or farming practises leading to siltation of gravel beds (Heywood and Walling 2003) may have damaged important dace spawning areas for example. Where natural off-river habitats have been lost through bank stabilisation to prevent oxbow lake formation or infilling to create pastureland, man-made creations in the form of drainage ditches have taken their place as low-flow lateral habitats. Now field drainage occurs through underground pipes, drainage ditches no longer function for their original purpose and are not regularly cleared or have been filled becoming inaccessible for fish. This trend of loss of natural and man-made connectivity is occurring in most temperate floodplain river systems and threatens many fish populations (Gore and Shields 1995, Olson and Dinerstein 1998).

Lateral connectivity in the River Frome provides a mosaic of habitats used to different degrees and during different parts of the life-cycle by a wide range of fishes (Chapters 4 and 5). Current literature on riverine habitat ecology and rehabilitation following anthropogenic change ascribes increasing importance of connectivity to lateral habitats for lowland river fish populations (Pretty et al. 2003). Rehabilitated off-river channels lead to increased fish abundance while in-stream structures have much lower effect (Langler and Smith 2001, Pretty et al. 2003). This study characterised the functioning of lateral side-channel habitats of the River Frome by fish communities. This will enable a more rational management of the Frome floodplain and beyond of floodplain rivers generally.

Ideally, due to the diversity of fish species requirements for off-river habitats (Chapters 4 and 5) the total preservation of floodplain systems and the maintenance or restoration of all lateral connection would be possible. However, funds, human resource requirements and the level of degradation that has already occurred in main riverine habitats mean that this is often not possible (Schiemer 1999) and a more
realistic and practical approach of rehabilitation must be adopted with the most important habitats being highest priority (Gore and Shields 1995).

Defining which lateral habitat is the most 'important' for the system depends on the criteria selected for the definition. Some habitats support a larger abundance or range of fish than others and could merit protection for this reason. Some authors have found that physical criteria may not be enough to determine ecological need and contribution to the system (Pretty et al. 2003). Yet this study found that species diversity in side-channels was positively correlated with their habitat stability (Chapter 4) and so an index of stability could indicate ecological contribution. This is a potentially useful tool for habitat managers and policy makers who may not have the resources to sample and monitor fish populations. They could determine the significance of a series of lateral habitats by taking simple habitat measurements such as those collected in this study over a period of time to create a stability index to rate each location.

Some lateral habitats support significant species, such as Goldsacs which contained a large population of brook lamprey a Biodiversity Action Plan (BAP) species. As no other side-channel in the study supported such a density of brook lamprey Goldsacs is a notable habitat for that reason. However, conservation solely for individual species may not provide the range of habitats necessary for a healthy riverine ecosystem. A combination of conservation of the most significant habitats to the overall system and those for particular species is therefore recommended. Understanding the functioning of these habitats within the system and for particular species may help to understand what drives the use of side-channels and thus which are particularly important to preserve and maintain.

As mentioned, Goldsacs is an important habitat for brook lamprey and it is also an important nursery habitat for flounder on the Frome. Eel, another BAP species, occur in many of the side streams sampled, particularly Holme Bridge and the Millstream. Eel were resident in the Millstream but moved between here and the main river during nocturnal activity. Dace used Millstream for spawning, Luckford for resting postspawning and Flood Relief as well as a natural oxbow near Luckford as a nursery habitat. Some dace also moved into both Luckford and Millstream in autumn.

Railway and Rushton provided a great deal of the local spawning habitat for pike. Pike were present at a lower level in most channels (except Goldsacs where they were not recorded) throughout the year.

This study on the River Frome can be used to create an example of a management plan for preserving the most important aspects of the floodplain. The two flowing streams Goldsacs and Millstream were the most stable habitats with the highest species diversity, Luckford was next most stable. Goldsacs also supported the only BAP species in the study, the brook lamprey. Luckford and Millstream were important habitats for dace supporting both spawning and sheltering individuals. Thus these three side-channels must be considered the three most important side-channels monitored during the study. However, this does not take into account the needs of the pike population for which drainage ditches are important spawning areas. So Rushton (more stable and species diverse than Railway) must also be included as an important lateral habitat for conserving. Finally, if preservation of all lateral habitats within the system were not possible Goldsacs, Luckford, Rushton and Millstream would be the most significant to preserve, in that order. A large number of reserve selection algorithms have been developed for application in prioritising relative importance of habitats for preservation or conservation (Nicholls and Margules 1993, Rothley 2006, Van Teeffelen et al. 2006). Further investigation into the relative importance of the side-channels on the River Frome could be continued using these algorithms.

The complex and mobile nature of the dace population structure showed that connectivity (both lateral and longitudinal) is an important element in the preservation of dace populations in the river system (Chapter 6). Habitat diversity and in particular access to certain habitats such as side-channels or oxbows may have disproportionate importance that is not always evident from species' abundance (Hoopes and Harrison 1998). Small changes in the distribution and accessibility of different habitats may precipitate relatively large changes in the species' viability (Doak 1995, Pulliam et al. 1992), an essential consideration for management plans. Should blockages to fish passage or loss of lateral connectivity prevent population mixing or reduce access to habitats necessary for life history requirements, the dace population as a whole is likely to suffer.

The life history strategies shown in pike highlight a variety of different connectivityrelated movements and migrations (Chapter 7). Behavioural strategies relating to habitat use and spatial distribution rely on both lateral and longitudinal connectivity. Different spawning strategies developed within the population are based on the nature and availability of lateral spawning sites. Behavioural diversity may enable the pike population to cope better with situations of change (Krebs and Davies 1997). For example, residents in poorer, side-channel habitats provide a source of colonisers should a population crash occur in the main river population. Side-channels are particularly important spawning habitat during unusually high flow years when eggs in the main river would be washed away. As pike appear to exhibit spawning-, and perhaps, natal-site fidelity conservation of a range of spawning habitats for pike is important so that the gene pool is not artificially reduced by lack of spawning areas. If this were to happen those individuals arriving early into the spawning-channels may be more likely to spawn successfully because in the limited habitat available those larvae that hatch early will have a size advantage (and thus lower threat from cannibalisation) to those hatching later in the season. In that case the trade-off between the cost of moving into the poorer feeding habitat (poorer for adult pike) earlier and spending longer there would be repaid by greater successful reproductive output.

Studying the dynamics of the pike and dace populations at the level of an individual within the population provides a much higher level of detail on the use and functionality of the side-channels, as well as the spatial distribution of individuals in the river system. The populations of the two species were structured very differently and had very different requirements from the lateral habitats. The need for a diverse range of habitats to be maintained throughout the river system and for integrated, catchment-wide management has been illustrated. In view of that, the final contribution of this thesis will be to consider practical implications and guidelines for conservation management of lateral connectivity of river systems.

### 9.1 Practical Implications

Lateral connectivity has been shown to be of importance to the ecology of a floodplain river system both in this study and others (Bayley 1995, Borcherding et al. 2002, Copp 1997, Gore and Shields 1995, Ross and Baker 1983, Slipke et al. 2005). According to several authors, restoration of lateral connectivity holds more benefits to the ecological integrity of lowland river systems than rehabilitation of in-stream structures (Buijse et al. 2002, Pretty et al. 2003, Schmutz and Jungwirth 1999). Human alterations to the ecosystem, such as poor water quality, damming, river regulation or abstraction all contribute to a cumulative negative effect on the communities and species resident in the river system. In particular, flood induced disturbances are a key element for maintaining biodiversity, creating characteristic patch dynamics and spatial heterogeneity (Schiemer 1999), as well as being a necessary part of the life history of some species (Bayley 1995, Gozlan 1998, Lucas 1992, Welcomme 1985). According to the flood pulse concept, seasonal flooding is not a disturbance but a typical and crucial event of floodplain rivers and connectivity (Junk et al. 1989). With so many human requirements on the river system and the associated changes, provision or maintenance of diverse and complex lateral habitats is especially important to enable the persistence of communities or populations which may be impacted by other changes.

The Water Framework Directive (WFD) requires "prevention of further deterioration and protection and enhancement of the status of water resources" (2000). The guidance document on the implementation of the WFD to rivers and lakes (WFD CIS Guidance Document No. 10 2003) recommends physical rehabilitation of river habitats, including lateral connectivity, to enable "recovery to a level of biodiversity and ecological functioning equivalent to unmodified natural water bodies". Such a recommendation should mean that as the WFD comes into operation the restoration of lateral connectivity will be a frequently used management practice applied in many ecosystems throughout Europe. Merging a system-oriented view with detailed analysis of functional requirements by populations, as approached in this study, will allow the development of the most appropriate management procedures. Creating
interdisciplinary teams of ecologists, hydrologists, environmental engineers and local stakeholders and combining the restoration of connected lateral habitats with continued assessment of the success of the management should ensure the greatest benefit to the riverine biota.

Studies such as this can increase knowledge and understanding of the complex interactions between fish communities and the lateral habitats available to them. Investigation of different dimensions of side-channel use, from assemblages through to individual fish, begins to provide an insight into the functionality of these habitats and the mosaic of these habitats provided in a floodplain system upon which management practices can be based. This study provided strong evidence for the need for a variety of different off-river habitats to enable the continuation of a balanced ecosystem and provision for all species habitat requirements to ensure high biodiversity of the sort in natural systems.

## Appendices

## Appendix 1 Table of PIT tagged fish

| ID | US PIT tag | UK PIT tag | Date tagged | Species | Fork length on first capture | First Catch Location |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | 94698332 | DC001C-B287 | 16/04/2003 | Pike | 56 | Main River Frome us road bridge |
| 19 |  | DC001B-E4CC | 17/03/2003 | Pike | 12.4 | Railway 1 |
| 20 | 124086156 | DC001B-E619 | 17/03/2003 | Pike | 11.2 | Railway 1 |
| 65 | 94698696 |  | 19/03/2003 | Dace | 22.7 | Luckford Lake 4 |
| 77 | 94698688 |  | 19/03/2003 | Dace | 19.3 | Luckford Lake 4 |
| 78 | 94698689 |  | 19/03/2003 | Dace | 19.4 | Luckford Lake 4 |
| 79 | 94698690 |  | 19/03/2003 | Dace | 18.9 | Luckford Lake 4 |
| 80 | 94698691 |  | 19/03/2003 | Dace | 22 | Luckford Lake 4 |
| 81 | 94698693 |  | 19/03/2003 | Dace | 22.2 | Luckford Lake 4 |
| 82 | 94698695 |  | 19/03/2003 | Dace | 25.6 | Luckford Lake 4 |
| 83 | 94698699 |  | 19/03/2003 | Roach | 18.8 | Luckford Lake 3 |
| 84 | 94698700 |  | 19/03/2003 | Roach | 17.9 | Luckford Lake 3 |
| 85 | 94698701 |  | 19/03/2003 | Roach | 19 | Luckford Lake 3 |
| 86 | 94698702 |  | 19/03/2003 | Dace | 22.6 | Luckford Lake 3 |
| 87 | 94698703 |  | 19/03/2003 | Dace | 21.5 | Luckford Lake 3 |
| 88 | 94698704 |  | 19/03/2003 | Dace | 23.1 | Luckford Lake 3 |
| 107 | 94698705 |  | 19/03/2003 | Dace | 21 | Luckford Lake 2 |
| 108 | 94698706 |  | 19/03/2003 | Dace | 18 | Luckford Lake 2 |
| 109 | 94698707 |  | 19/03/2003 | Dace | 21.2 | Luckford Lake 2 |
| 110 | 94698708 |  | 19/03/2003 | Dace | 22.8 | Luckford Lake |
| 111 | 94698709 |  | 19/03/2003 | Dace | 16.2 | Luckford Lake 2 |
| 112 | 94698710 |  | 19/03/2003 | Dace | 20.2 | Luckford Lake 2 |
| 113 | 94698711 |  | 19/03/2003 | Pike | 33.1 | Luckford Lake 2 |
| 114 | 94698698 |  | 19/03/2003 | Pike | 34.2 | Luckford Lake 3 |
| 115 | 94698692 |  | 19/03/2003 | Pike | 45.5 | Luckford Lake 4 |
| 117 | 94698713 |  | 19/03/2003 | Pike | 60.5 | Luckford Lake 1 |
| 118 | 94698712 |  | 19/03/2003 | Pike | 65 | Luckford Lake 1 |
| 135 |  | DC001B-F278 | 20/03/2003 | Dace | 9.3 | Holme Bridge 3 |
| 139 |  | DC001B-EECB | 20/03/2003 | Roach | 11.4 | Holme Bridge 3 |
| 140 |  | DC001B-E8CD | 20/03/2003 | Roach | 12 | Holme Bridge 3 |
| 142 |  | DC001B-EEF2 | 20/03/2003 | Roach | 12.2 | Holme Bridge 3 |
| 144 |  | DC001C-0C80 | 20/03/2003 | Roach | 11.5 | Holme Bridge 3 |
| 149 |  | DC001C-0AEE | 20/03/2003 | Roach | 7.5 | Holme Bridge 1 |
| 150 |  | DC001C-0BB5 | 20/03/2003 | Dace |  | Holme Bridge 1 |
| 151 |  | DC001C-085B | 20/03/2003 | Roach | 8.5 | Holme Bridge 1 |
| 152 |  | DC001C-0198 | 20/03/2003 | Dace |  | Holme Bridge 1 |
| 153 |  | DC001B-FAC4 | 20/03/2003 | Dace | 9.3 | Holme Bridge 1 |
| 155 |  | DC001B-ED9C | 20/03/2003 | Roach | 12 | Holme Bridge 1 |
| 156 | 94698683 |  | 20/03/2003 | Pike | 18.8 | Holme Bridge 4 |
| 157 | 94698682 |  | 20/03/2003 | Pike | 13.5 | Holme Bridge 4 |
| 158 | 94698679 |  | 20/03/2003 | Pike | 52.5 | Holme Bridge Extra |
| 159 | 94698681 |  | 20/03/2003 | Pike | 13.5 | Holme Bridge Extra |
| 160 | 94698685 |  | 20/03/2003 | Pike | 14 | Holme Bridge 4 |
| 161 | 94698684 |  | 20/03/2003 | Pike | 28.5 | Holme Bridge 4 |
| 163 |  | DC001B-F450 | 20/03/2003 | Roach | 8.5 | Holme Bridge 2 |
| 165 | 94698680 |  | 20/03/2003 | Pike | 12.5 | Holme Bridge Extra |
| 166 | 94698653 |  | 24/03/2003 | Dace | 14.6 | ESMS |


| 167 | 94698655 | 24/03/2003 | Dace | 22.8 | ESMS |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 168 | 94698656 | 24/03/2003 | Dace | 22 | ESMS |
| 169 | 94698657 | 24/03/2003 | Dace | 23.6 | ESMS |
| 170 | 94698654 | 24/03/2003 | Dace | 19.8 | ESMS |
| 171 | 94698652 | 24/03/2003 | Dace | 18 | ESMS |
| 172 | 94698651 | 24/03/2003 | Dace | 20.2 | ESMS |
| 173 | 94698650 | 24/03/2003 | Dace | 20.5 | ESMS |
| 174 | 94698658 | 24/03/2003 | Dace | 24 | ESMS |
| 175 | 94698662 | 24/03/2003 | Dace | 22.7 | ESMS |
| 176 | 94698659 | 24/03/2003 | Dace | 23.4 | ESMS |
| 177 | 94698661 | 24/03/2003 | Dace | 18.3 | ESMS |
| 178 | 94698660 | 24/03/2003 | Dace | 17.4 | ESMS |
| 179 | 94698663 | 24/03/2003 | Dace | 18.5 | ESMS |
| 180 | 94698664 | 24/03/2003 | Dace | 17.5 | ESMS |
| 181 | 94698666 | 24/03/2003 | Dace | 19.5 | ESMS |
| 182 | 94698670 | 24/03/2003 | Dace | 21.6 | ESMS |
| 183 | 94698667 | 24/03/2003 | Dace | 24 | ESMS |
| 184 | 94698671 | 24/03/2003 | Dace | 24.9 | ESMS |
| 186 | 94698672 | 24/03/2003 | Dace | 21 | ESMS |
| 187 | 94698647 | 24/03/2003 | Dace | 17.4 | ESMS |
| 188 | 94698668 | 24/03/2003 | Dace | 19.5 | ESMS |
| 189 | 94698669 | 24/03/2003 | Dace | 25.5 | ESMS |
| 190 | 94698665 | 24/03/2003 | Dace | 23 | ESMS |
| 191 | 94698620 | 24/03/2003 | Roach | 14.1 | ESMS |
| 192 | 94698621 | 24/03/2003 | Dace | 19.2 | ESMS |
| 193 | 94698622 | 24/03/2003 | Dace | 19.5 | ESMS |
| 194 | 94698630 | 24/03/2003 | Dace | 17.1 | ESMS |
| 195 | 94698629 | 24/03/2003 | Dace | 19.1 | ESMS |
| 196 | 94698628 | 24/03/2003 | Dace | 20.4 | ESMS |
| 197 | 94698627 | 24/03/2003 | Dace | 19 | ESMS |
| 198 | 94698626 | 24/03/2003 | Dace | 15.8 | ESMS |
| 199 | 94698625 | 24/03/2003 | Dace | 19.2 | ESMS |
| 200 | 94698624 | 24/03/2003 | Dace | 21.2 | ESMS |
| 201 | 94698623 | 24/03/2003 | Dace | 18.4 | ESMS |
| 202 | 94698632 | 24/03/2003 | Dace | 20.7 | ESMS |
| 203 | 94698631 | 24/03/2003 | Dace | 20.2 | ESMS |
| 204 | 94698633 | 24/03/2003 | Dace | 20.9 | ESMS |
| 205 | 94698634 | 24/03/2003 | Dace | 18 | ESMS |
| 206 | 94698635 | 24/03/2003 | Dace | 18.9 | ESMS |
| 207 | 94698637 | 24/03/2003 | Dace | 18.1 | ESMS |
| 209 | 94698639 | 24/03/2003 | Dace | 18.9 | ESMS |
| 210 | 94698638 | 24/03/2003 | Dace | 17.4 | ESMS |
| 211 | 94698640 | 24/03/2003 | Dace | 20.7 | ESMS |
| 212 | 94698641 | 24/03/2003 | Dace | 21.2 | ESMS |
| 213 | 94698642 | 24/03/2003 | Dace | 24.3 | ESMS |
| 214 | 94698643 | 24/03/2003 | Dace | 20.6 | ESMS |
| 215 | 94698644 | 24/03/2003 | Dace | 18 | ESMS |
| 216 | 94698646 | 24/03/2003 | Roach | 17.8 | ESMS |
| 217 | 94698645 | 24/03/2003 | Roach | 18.5 | ESMS |
| 218 | 94698619 | 24/03/2003 | Dace | 15.3 | ESMS |
| 219 | 94698618 | 24/03/2003 | Roach | 13.6 | ESMS |
| 220 | 94698403 | 24/03/2003 | Dace | 20.4 | ESMS |
| 221 | 94698402 | 24/03/2003 | Dace | 19.2 | ESMS |
| 222 | 94698648 | 24/03/2003 | Roach | 14.5 | ESMS |


| 223 | 94698649 |  | 24/03/2003 | Roach | 14.5 | ESMS |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 224 | 94698677 |  | 24/03/2003 | Pike | 61.6 | ESMS |
| 225 | 94698674 |  | 24/03/2003 | Pike | 21.3 | ESMS |
| 226 | 94698673 |  | 24/03/2003 | Pike | 59 | ESMS |
| 227 | 94698398 |  | 24/03/2003 | Pike | 34.3 | ESMS eel trap |
| 228 | 94698399 |  | 24/03/2003 | Pike | 44.5 | ESMS eel trap |
| 229 | 94698678 |  | 24/03/2003 | Pike | 77 | ESMS Lily Pool |
| 230 | 94698380 |  | 24/03/2003 | Roach | 13 | ESMS eel trap |
| 231 | 94698381 |  | 24/03/2003 | Roach | 13.7 | ESMS eel trap |
| 232 | 94698382 |  | 24/03/2003 | Roach | 12.8 | ESMS eel trap |
| 233 | 94698383 |  | 24/03/2003 | Roach | 12.3 | ESMS eel trap |
| 234 | 94698384 |  | 24/03/2003 | Dace | 14.6 | ESMS eel trap |
| 235 | 94698385 |  | 24/03/2003 | Roach | 15.3 | ESMS eel trap |
| 236 | 94698386 |  | 24/03/2003 | Dace | 14 | ESMS eel trap |
| 237 | 94698387 |  | 24/03/2003 | Roach | 13.2 | ESMS eel trap |
| 238 | 94698388 |  | 24/03/2003 | Roach | 13.5 | ESMS eel trap |
| 239 | 94698389 |  | 24/03/2003 | Roach | 14.1 | ESMS eel trap |
| 240 | 94698390 |  | 24/03/2003 | Perch | 14.2 | ESMS eel trap |
| 241 | 94698391 |  | 24/03/2003 | Perch | 15.7 | ESMS eel trap |
| 242 | 94698392 |  | 24/03/2003 | Roach | 18.2 | ESMS eel trap |
| 243 | 94698393 |  | 24/03/2003 | Roach | 12.7 | ESMS eel trap |
| 244 | 94698394 |  | 24/03/2003 | Roach | 14.2 | ESMS eel trap |
| 245 | 94698395 |  | 24/03/2003 | Perch | 21.7 | ESMS eel trap |
| 246 | 94698396 |  | 24/03/2003 | Roach | 51 | ESMS eel trap |
| 247 | 94698397 |  | 24/03/2003 | Roach | 17.7 | ESMS eel trap |
| 248 | 94698400 |  | 24/03/2003 | Roach | 16.1 | ESMS |
| 249 | 94698401 |  | 24/03/2003 | Roach | 13 | ESMS eel trap |
| 250 | 94698377 |  | 25/03/2003 | Pike | 60.4 | Main River Frome |
| 251 | 94698379 |  | 25/03/2003 | Pike | 61 | Main River Frome |
| 252 | 94698376 |  | 25/03/2003 | Pike | 72.9 | Main River Frome |
| 253 | 94698375 |  | 25/03/2003 | Pike | 47.1 | Main River Frome |
| 254 | 94698360 |  | 25/03/2003 | Pike | 86.2 | Main River Frome |
| 255 | 94698373 |  | 25/03/2003 | Pike | 65 | Main River Frome |
| 256 | 94698361 |  | 25/03/2003 | Pike | 79.8 | Main River Frome |
| 257 | 94698374 |  | 25/03/2003 | Pike | 18.1 | Main River Frome |
| 258 | 94698359 |  | 25/03/2003 | Dace | 21.1 | Main River Frome |
| 260 | 94698356 |  | 25/03/2003 | Dace | 21.5 | Main River Frome |
| 261 | 94698358 |  | 25/03/2003 | Dace | 18.8 | Main River Frome |
| 262 | 94698366 |  | 25/03/2003 | Dace | 19 | Main River Frome |
| 263 | 94698365 |  | 25/03/2003 | Dace | 18.7 | Main River Frome |
| 264 | 94698364 |  | 25/03/2003 | Dace | 17.3 | Main River Frome |
| 265 | 94698363 |  | 25/03/2003 |  | 18 | Main River Frome |
| 266 | 94698362 |  | 25/03/2003 | Dace | 21.3 | Main River Frome |
| 267 | 94698367 |  | 25/03/2003 | Dace | 21.5 | Main River Frome |
| 268 | 94698368 |  | 25/03/2003 | Dace | 17.5 | Main River Frome |
| 269 | 94698369 |  | 25/03/2003 | Dace | 19.1 | Main River Frome |
| 270 | 94698370 |  | 25/03/2003 | Dace | 22.6 | Main River Frome |
| 271 | 94698371 |  | 25/03/2003 | Dace | 23 | Main River Frome |
| 272 | 94698372 |  | 25/03/2003 | Roach | 21.2 | Main River Frome |
| 273 | 94698378 |  | 25/03/2003 | Dace | 18.8 | Main River Frome |
| 274 | 94698350 | DC001C-9126 | 26/03/2003 | Pike | 17.5 | ESMS Millhead |
| 275 | 94698349 |  | 26/03/2003 | Pike | 17.5 | ESMS Millhead |
| 276 | 94698353 | DC001C-6FCC | 26/03/2003 | Pike | 59.3 | ESMS Millhead |
| 277 | 94698352 | DC001C-7FEB | 26/03/2003 | Pike | 45.4 | ESMS Millhead |


| 278 | 94698351 | DC001C-B4FC | 26/03/2003 | Pike |
| :---: | :---: | :---: | :---: | :---: |
| 279 | 94698355 | DC001C-B331 | 26/03/2003 | Pike |
| 280 | 94698354 | DC001C-A832 | 26/03/2003 | Pike |
| 281 | 94698347 | DC001C-B1C0 | 01/04/2003 | Pike |
| 282 | 94698348 | DC001C-B7EA | 01/04/2003 | Pike |
| 283 | 94698346 | DC001C-77DF | 15/04/2003 | Pike |
| 284 | 94698314 |  | 14/04/2003 | Pike |
| 285 | 94698318 |  | 16/04/2003 | Dace |
| 286 | 94698319 |  | 16/04/2003 | Roach |
| 287 | 94698323 | DC001C-A06E | 16/04/2003 | Dace |
| 288 | 94698334 | DC001C-AE21 | 16/04/2003 | Pike |
| 289 | 94698322 | DC001C-9DAA | 16/04/2003 | Pike |
| 290 | 94698321 | DC001C-B8FD | 16/04/2003 | Pike |
| 291 | 94698320 | DC001C-A6C8 | 16/04/2003 | Pike |
| 292 | 94698331 | DC001C-AB5D | 16/04/2003 | Pike |
| 293 | 94698315 |  | 16/04/2003 | Pike |
| 294 | 94698317 | DC001C-9E28 | 16/04/2003 | Pike |
| 295 | 94698330 | DC001C-7BA4 | 16/04/2003 | Pike |
| 296 | 94698328 | DC001C-B604 | 16/04/2003 | Pike |
| 297 | 94698329 | DC001C-B66C | 16/04/2003 | Pike |
| 298 | 94698327 | DC001C-842E | 16/04/2003 | Pike |
| 299 | 94698326 | DC001C-77E3 | 16/04/2003 | Pike |
| 300 | 94698325 | DC001C-B990 | 16/04/2003 | Pike |
| 301 | 94698316 |  | 16/04/2003 | Pike |
| 302 | 94698313 | DC001C-BA53 | 01/05/2003 | Pike |
| 303 | 94698307 | DC001C-7A21 | 06/05/2003 | Pike |
| 304 | 94698312 | DC001C-BAO7 | 06/05/2003 | Pike |
| 305 | 94698309 | DC001B-F3EB | 06/05/2003 | Pike |
| 306 | 94698308 | DC001C-B64A | 06/05/2003 | Pike |
| 307 | 94698324 | DC001C-BBB9 | 16/07/2003 | Dace |
| 308 | 94698305 | DC001C-6DEB | 20/05/2003 | Pike |
| 309 | 94698303 | DC001C-A332 | 20/05/2003 | Pike |
| 310 | 94698306 | DC001C-A355 | 20/05/2003 | Pike |
| 311 | 94698310 | DC001C-A053 | 20/05/2003 | Pike |
| 312 | 94698300 | DC001C-78D7 | 23/06/2003 | Pike |
| 313 | 94698302 | DC001C-A24D | 23/06/2003 | Pike |
| 314 | 94698301 |  | 23/06/2003 | Perch |
| 315 |  | DC001C-B470 | 23/06/2003 | Dace |
| 316 |  | DC001C-7418 | 23/06/2003 | Dace |
| 317 |  | DC001C-7297 | 23/06/2003 | Dace |
| 318 |  | DC001C-B865 | 23/06/2003 | Dace |
| 319 | 94698299 | DC001C-FDC0 | 23/06/2003 | Pike |
| 320 | 94698298 | DC001B-E5B3 | 23/06/2003 | Pike |
| 321 | 113943627 | DC001C-7F6C | 24/06/2003 | Pike |
| 322 | 94698297 | DC001C-7BA3 | 24/06/2003 | Pike |


| 16.1 | ESMS Millhead |
| :---: | :---: |
| 62.8 | ESMS Millhead |
| 62.3 | ESMS Millhead |
| 81 | ESMS Millhead |
| 56 | ESMS |
| 55.4 | ESMS |
| 73.2 | Main River Frome |
| 17.2 | Main River Frome |
| 19.2 | Main River Frome |
| 17.9 | Main River Frome Main River Frome us |
| 63.9 | road bridge |
| 62.1 | Main River Frome |
| 58.8 | Main River Frome Main River Frome Road |
| 78.3 | bridge |
| 51.8 | Main River Frome us road bridge |
| 64.8 | Main River Frome |
| 101 | Main River Frome |
| 48 | road bridge |
| 45.5 | Main River Frome us road bridge |
| 25.7 | Main River Frome us road bridge |
| 21.3 | Main River Frome us road bridge |
|  | Main River Frome us |
| 23 | road bridge <br> Main River Frome us |
| 21.7 | road bridge |
| 22.6 | Main River Frome |
| 95.5 | ESMS Millhead ESMH above smolt |
| 64.5 | counter |
| 71 | Main River Frome T4 Main River Frome T9 - |
| 58 | T10 |
| 78 | Main River Frome T2 Main River Frome |
| 23.2 | Roadbridge/Ducks Egg |
| 47.5 | ESMH |
| 22.2 | ESMH |
| 45.5 | ESMH |
| 68.5 | ESMH |
| 24.8 | Rushton 1 |
| 29.8 | Rushton 1 |
| 25.8 | Rushton 1 |
| -9 | ESMS 1 |
| -9 |  |
| 11 |  |
| 11.5 |  |
| 19.7 | Flood Relief 3 |
| 37.1 | Flood Relief 2 |
| 18.5 | Luckford Lake 2 |
| 23.9 | Railway 2 |


| 94698296 |  | 24/06/2003 | Pike |
| :---: | :---: | :---: | :---: |
| 94698289 | DC001C-898D | 24/06/2003 | Pike |
|  | DC001C-B5B3 | 25/06/2003 | Roach |
|  | DC001C-8BCF | 25/06/2003 | Roach |
|  | DC001C-785A | 25/06/2003 | Roach |
|  | DC001C-8A73 | 25/06/2003 | Pike |
|  | DC001C-7130 | 25/06/2003 | Roach |
|  | DC001C-A85D | 25/06/2003 | Roach |
|  | DC001C-7B92 | 25/06/2003 | Roach |
| 94698252 | DC0024-EFB3 | 08/09/2003 | Roach |
| 94698251 | DC0024-DD87 | 08/09/2003 | Dace |
| 94698284 | DC0024-EEF7 | 08/09/2003 | Dace |
| 94698283 | DC0024-E725 | 08/09/2003 | Roach |
| 94698282 | DC0024-E17F | 08/09/2003 | Roach |
| 94698281 | DC0024-CCFE | 08/09/2003 | Dace |
| 94698280 | DC0024-DED2 | 08/09/2003 | Roach |
| 94698278 | DC0024-DDF9 | 08/09/2003 | Dace |
| 94698279 | DC0024-DACD | 08/09/2003 | Roach |
| 94698277 | DC0024-E87F | 08/09/2003 | Dace |
| 94698276 | DC0024-EEBB | 08/09/2003 | Roach |
| 94698271 | DC0024-CB6D | 08/09/2003 | Dace |
| 94698272 | DC0024-CFC9 | 08/09/2003 | Dace |
| 94698273 | DC0024-CE47 | 08/09/2003 | Dace |
| 94698274 | DC0024-D2CC | 08/09/2003 | Roach |
| 94698288 | DC001B-EE42 | 17/03/2003 | Pike |
| 94698286 |  | 08/09/2003 | Pike |
| 94698287 |  | 08/09/2003 | Pike |
| 94698246 | DC0024-E965 | 08/09/2003 | Roach |
| 94698248 | DC0024-F332 | 08/09/2003 | Dace |
| 94698250 | DC0024-CC8A | 08/09/2003 | Dace |
| 94698254 | DC0024-DB26 | 08/09/2003 | Roach |
| 94698253 | DC0024-E9FC | 08/09/2003 | Roach |
| 94698249 | DC0024-E54C | 08/09/2003 | Dace |
| 94698285 | DC0024-F32C | 08/09/2003 | Dace |
| 94698247 | DC0024-CEFB | 08/09/2003 | Dace |
| 94698245 | DC0024-F030 | 08/09/2003 | Roach |
| 94698258 | DC0024-E430 | 08/09/2003 | Dace |
| 94698257 | DC0024-E91F | 08/09/2003 | Roach |
| 94698255 | DC0024-ECC1 | 08/09/2003 | Dace |
| 94698256 | DC0024-EFB0 | 08/09/2003 | Roach |
| 94698259 | DC0024-E8B4 | 08/09/2003 | Dace |
| 94698260 | DC0024-D552 | 08/09/2003 | Roach |
| 113943601 | DC0024-ES34 | 08/09/2003 | Dace |
| 94698265 | DC0024-E9BC | 08/09/2003 | Dace |
| 94698264 | DC0024-E2F7 | 08/09/2003 | Roach |
| 11394360 | DC0024-F1E7 | 08/09/2003 | Dace |
| 113943602 | DC0024-CF25 | 08/09/2003 | Dace |
| 94698266 | DC0024-E602 | 08/09/2003 | Perch |
| 113943610 | DC0024-EC68 | 08/09/2003 | Roach |
| 113943604 | DC0024-DCA5 | 08/09/2003 | Dace |
| 94698262 | DC0024-CBE1 | 08/09/2003 | Roach |
| 94698261 | DC0024-DFF5 | 08/09/2003 | Roach |
| 94698268 | DC0024-D649 | 08/09/2003 | Dace |
| 113943605 | DC0024-CA56 | 08/09/2003 |  |

17 Luckford Lake 1
13.2 Railway 2
9.5 Holme Bridge 4
9.4 Holme Bridge

14 Holme Bridge 4
13.6 Holme Bridge 4
10.2 Holme Bridge 4
9.1 Holme Bridge Extra
8.5 Holme Bridge Extra
23.7 ESMH
20.5 ESMH

16 ESMH
18.6 ESMH
21.6 ESMH
19.6 ESMH
18.2 ESMH
17.1 ESMH
16.9 ESMH
16.3 ESMH
21.9 ESMH
25.6 ESMH
25.7 ESMH
14.4 ESMH
17.8 ESMH
11.4 Railway 4
19.3 Railway 4
17.5 Railway 3
17.3 ESMH
21.5 ESMH
21.4 ESMH
16.4 ESMH
18.3 ESMH
19.5 ESMH
22.5 ESMH

21 ESMH
14.3 ESMH
16.9 ESMH
16.5 ESMH
25.8 ESMH
15.5 ESMH
15.4 ESMH
15.1 ESMS
15.2 ESMH
20.5 ESMH
13.9 ESMH
13.6 ESMH
19.5 ESMH
23.5 ESMH
13.2 ESMH
14.6 ESMH
15.3 ESMH
13.9 ESMH
17.2 ESMH

16 ESMH

| 377 | 94698269 | DC0024-EFID | 08/09/2003 | Dace | 15.5 | ESMH |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 378 | 94698267 | DC0024-EBB3 | 08/09/2003 | Roach | 15.6 | ESMH |
| 379 | 113943606 | DC0024-EC90 | 08/09/2003 | Dace | 23.2 | ESMH |
| 380 | 113943598 | DC0024-EF58 | 08/09/2003 | Roach | 18.5 | ESMH |
| 381 | 94698270 | DC0024-DGAF | 08/09/2003 | Dace | 24.2 | ESMH |
| 382 | 113943608 | DC0024-CBAC | 08/09/2003 | Roach | 18 | ESMH |
| 383 | 113943600 | DC0024-D29D | 08/09/2003 | Roach | 16.3 | ESMH |
| 384 | 113943599 | DC0024-F19F | 08/09/2003 | Dace | 14.9 | ESMH |
| 385 | 94698342 | DC0024-DGAC | 08/09/2003 | Dace | 15.4 | ESMH |
| 386 | 113943609 | DC0024-EBF3 | 08/09/2003 | Dace | 22.2 | ESMH |
| 387 | 113943607 | DC0024-E13F | 08/09/2003 | Dace | 16.5 | ESMH |
| 388 | 113943611 | DC0024-DEA6 | 08/09/2003 | Dace | 14.9 | ESMH |
| 389 | 94698244 | DC0024-ED9B | 08/09/2003 | Dace | 14.6 | ESMH |
| 390 | 94698242 | DC0024-E780 | 08/09/2003 | Dace | 21.1 | ESMH |
| 391 | 94698243 | DC0024-F1C5 | 08/09/2003 | Roach | 18.6 | ESMH |
| 392 | 94698238 | DC0024-E83B | 08/09/2003 | Dace | 13.9 | ESMH |
| 393 | 94698240 | DC0024-EA57 | 08/09/2003 | Dace | 14.3 | ESMH |
| 394 | 94698239 | DC0024-EFCF | 08/09/2003 | Dace | 16.7 | ESMH |
| 395 | 94698236 |  | 09/09/2003 | Trout | 22.8 | ESMS 1 |
| 396 | 94698233 |  | 09/09/2003 | Trout | 21.4 | ESMS 1 |
| 397 | 94698234 |  | 09/09/2003 | Eel | 49.7 | ESMS 1 |
| 398 | 94698229 |  | 09/09/2003 | Eel | 32 | ESMS 1 |
| 399 | 94698232 |  | 09/09/2003 | Eel | 32.7 | ESMS 1 |
| 400 | 94698230 |  | 09/09/2003 | Eel | 37.2 | ESMS 1 |
| 401 | 94698235 |  | 09/09/2003 | Trout | 28 | ESMS 1 |
| 402 | 94698231 |  | 09/09/2003 | Eel | 33.4 | ESMS 1 |
| 403 | 94698237 |  | 09/09/2003 | Eel | 41.3 | ESMS 1 |
| 404 | 94698177 | DC001C-0A92 | 09/09/2003 | Dace | 13.8 | ESMS 2 |
| 405 | 94698178 | DC001B-E867 | 09/09/2003 | Dace | 14.3 | ESMS 2 |
| 406 | 94698157 | DC001B-F6E3 | 09/09/2003 | Dace | 12.1 | ESMS 2 |
| 407 | 94698179 | DC001B-EF00 | 09/09/2003 | Dace | 16.5 | ESMS 2 |
| 408 | 94698187 | DC001C-07BF | 09/09/2003 | Dace | 15.9 | ESMS 2 |
| 409 | 94698191 | DC0024-D1A3 | 09/09/2003 | Dace | 14.3 | ESMS 2 |
| 410 | 94698167 | DC001B-E3F3 | 09/09/2003 | Dace | 13.6 | ESMS 2 |
| 411 | 94698192 | DC001B-F781 | 09/09/2003 | Dace | 14.5 | ESMS 2 |
| 412 | 94698188 | DC001C-0118 | 09/09/2003 | Dace | 14.9 | ESMS 2 |
| 413 | 94698149 | DC001B-E2DE | 09/09/2003 | Dace | 15.9 | ESMS 2 |
| 414 | 94698171 | DC001B-F18E | 09/09/2003 | Dace | 13.7 | ESMS 1 |
| 415 | 94698174 | DC001B-EFEB | 09/09/2003 | Dace | 15.5 | ESMS 2 |
| 416 | 94698166 | DC001B-F3A6 | 09/09/2003 | Dace | 15.1 | ESMS 2 |
| 417 | 94698183 | DC001B-EF45 | 09/09/2003 | Pike | 16.5 | ESMS 2 |
| 418 | 94698181 | DC001C-038A | 09/09/2003 | Dace | 18.7 | ESMS 2 |
| 419 | 94698175 | DC0024-E57C | 09/09/2003 | Dace | 14.8 | ESMS 2 |
| 420 | 94698184 | DC001B-E97C | 09/09/2003 | Dace | 15.6 | ESMS 2 |
| 421 | 94698176 | DC001C-08B3 | 09/09/2003 | Dace |  | ESMS 2 |
| 422 | 94698170 | DC001B-EE85 | 09/09/2003 | Dace | 14 | ESMS 2 |
| 423 | 94697175 | DC001B-E499 | 09/09/2003 | Dace | 14.8 | ESMS 2 |
| 424 | 94698165 | DC001B-FDE5 | 09/09/2003 | Dace | 15.9 | ESMS 2 |
| 425 | 94698168 | DC001C-055D | 09/09/2003 | Dace | 14.9 | ESMS 2 |
| 426 | 94698169 | DC001B-E722 | 09/09/2003 | Dace |  | ESMS 2 |
| 427 | 94698198 | DC0024-C90E | 09/09/2003 | Trout | 27 | ESMS 2 |
| 428 | 94698208 | DC0024-D22C | 09/09/2003 | Dace | 16.1 | ESMS 2 |
| 429 | 94698180 | DC0024-EAEA | 09/09/2003 | Dace | 14.5 | ESMS 2 |
| 430 | 94698193 | DC0024-EC34 | 09/09/2003 | Dace | 14 | ESMS 2 |


| 431 | 94698182 | DC0024-D324 | 09/09/2003 | Dace | 12.9 | ESMS 2 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 432 | 94698189 | DC0024-D46E | 09/09/2003 | Dace | 19.3 | ESMS 2 |
| 433 | 94698186 | DC0024-E8FB | 09/09/2003 | Dace | 21.5 | ESMS 2 |
| 434 | 94698190 | DC0024-D4B7 | 09/09/2003 | Dace | 21.5 | ESMS 2 |
| 435 | 94698185 | DC0024-D993 | 09/09/2003 | Dace | 15 | ESMS 2 |
| 436 | 94698173 | DC00TB-E559 | 09/09/2003 | Dace | 16.8 | ESMS 2 |
| 437 | 94698199 | DC0024-E646 | 09/09/2003 | Dace | 16.1 | ESMS 2 |
| 438 | 94698194 | DC0024-CAE0 | 09/09/2003 | Dace | 16.3 | ESMS 2 |
| 439 | 94698200 | DC0024-D3BB | 09/09/2003 | Dace | 16.7 | ESMS 2 |
| 440 | 94698211 | DC0024-F3BF | 09/09/2003 | Dace | 15 | ESMS 2 |
| 441 | 94698202 | DC0024-E910 | 09/09/2003 | Dace | 14.8 | ESMS 2 |
| 442 | 94698203 | DC0024-E132 | 09/09/2003 | Dace | 13.4 | ESMS 2 |
| 443 | 94698205 | DC0024-CFE2 | 09/09/2003 | Dace | 15 | ESMS 2 |
| 444 | 94698206 | DC0024-E5FC | 09/09/2003 | Dace | 14 | ESMS 2 |
| 445 | 94698207 | DC0024-DC73 | 09/09/2003 | Dace | 18.4 | ESMS 2 |
| 446 | 94698210 | DC0024-CAFF | 09/09/2003 | Dace | 13.9 | ESMS 2 |
| 447 | 94698209 | DC0024-CEE2 | 09/09/2003 | Dace | 14.5 | ESMS 2 |
| 448 | 35471119 | DC0024-D013 | 09/09/2003 | Dace | 15.2 | ESMS 2 |
| 449 | 94698197 | DC0024-E0A5 | 09/09/2003 | Dace | 15.1 | ESMS 2 |
| 450 | 94698196 | DC0024-DC65 | 09/09/2003 | Dace | 14 | ESMS 2 |
| 451 | 94698222 |  | 09/09/2003 | Dace | 19 | ESMS 2 |
| 452 | 94698223 |  | 09/09/2003 | Dace | 15.2 | ESMS 2 |
| 453 | 94698224 |  | 09/09/2003 | Dace | 15.3 | ESMS 2 |
| 454 | 94698219 |  | 09/09/2003 | Roach | 13.6 | ESMS 2 |
| 455 | 94698215 |  | 09/09/2003 | Dace | 12.8 | ESMS 2 |
| 456 | 94698228 | DC0024-E25B | 09/09/2003 | Dace | 13.7 | ESMS 2 |
| 457 | 94698201 | DC0024-DB6F | 09/09/2003 | Dace |  | ESMS 2 |
| 458 | 94698204 | DC0024-E84F | 09/09/2003 | Dace | 12.5 | ESMS 2 |
| 459 | 94698213 |  | 09/09/2003 | Roach | 15.6 | ESMS 2 |
| 460 | 94698220 | DC0024-D72C | 09/09/2003 | Dace | 15.2 | ESMS 2 |
| 461 | 94698212 |  | 09/09/2003 | Dace | 14.4 | ESMS 2 |
| 462 | 94698218 |  | 09/09/2003 | Dace | 14.4 | ESMS 2 |
| 463 | 94698214 | DC0024-C03A | 09/09/2003 | Dace | 13.5 | ESMS 2 |
| 464 | 94698221 | DC0024-E12A | 09/09/2003 | Dace | 15.2 | ESMS 2 |
| 465 | 94698227 | DC0024-D05C | 09/09/2003 | Dace | 18 | ESMS 2 |
| 466 | 94698226 | DC0024-DA72 | 09/09/2003 | Dace | 16.8 | ESMS 2 |
| 467 | 94698225 | DC0024-D21B | 09/09/2003 | Dace | 14.5 | ESMS 2 |
| 468 | 94698216 | DC0024-CA5F | 09/09/2003 | Dace | 15.4 | ESMS 2 |
| 469 | 94698217 | DC0024-D913 | 09/09/2003 | Dace | 15.3 | ESMS 2 |
| 470 | 94698160 | DC001C-0540 | 09/09/2003 | Dace |  | ESMS 2 |
| 471 | 94698144 | DC001B-E1BF | 09/09/2003 | Dace | 12.2 | ESMS 2 |
| 472 | 94698147 | DC001B-F3A3 | 09/09/2003 | Dace | 15.5 | ESMS 2 |
| 473 | 94698150 | DC001B-F8A4 | 09/09/2003 | Dace | 14.5 | ESMS 2 |
| 474 | 94698156 | DC001B-E5FF | 09/09/2003 | Dace | 14 | ESMS 2 |
| 475 | 94698158 | DC001C-0CF9 | 09/09/2003 | Dace | 16.4 | ESMS 2 |
| 476 | 94698159 | DC001B-E239 | 09/09/2003 | Dace | 15.9 | ESMS 2 |
| 477 | 94698151 | DC0024-E168 | 09/09/2003 | Dace | 14.9 | ESMS 2 |
| 478 | 94698145 | DC0024-DE86 | 09/09/2003 | Dace | 15.3 | ESMS 2 |
| 479 | 94698155 | DC0024-DE05 | 09/09/2003 | Dace | 19.9 | ESMS 2 |
| 480 | 94698154 | DC0024-CA8F | 09/09/2003 | Dace | 15 | ESMS 2 |
| 481 | 94698162 | DC001B-F59E | 09/09/2003 | Dace | 14.4 | ESMS 2 |
| 482 | 94698163 | DC0024-DCD5 | 09/09/2003 | Dace | 15.5 | ESMS 2 |
| 483 | 94698164 | DC0024-CB45 | 09/09/2003 | Dace | 15.3 | ESMS 2 |
| 484 | 94698161 | DC0024-CB20 | 09/09/2003 | Dace | 14.2 | ESMS 2 |


| 485 | 94698172 | DC0024-D18B | 09/09/2003 | Dace | 14.3 | ESMS 2 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 486 | 94698129 | DC0024-D586 | 09/09/2003 | Dace | 15.5 | ESMS 2 |
| 487 | 94698132 | DC0024-E3DF | 09/09/2003 | Dace | 13.9 | ESMS 2 |
| 488 | 94698131 | DC0024-C8CF | 09/09/2003 | Dace | 15.3 | ESMS 2 |
| 489 | 94698133 | DC0024-C8A0 | 09/09/2003 | Dace | 16.4 | ESMS 2 |
| 490 | 94698136 | DC0024-CD98 | 09/09/2003 | Dace | 14.4 | ESMS 2 |
| 491 | 94698134 | DC0024-E32A | 09/09/2003 | Dace | 13.8 | ESMS 2 |
| 492 | 94698138 | DC0024-E359 | 09/09/2003 | Dace | 17.4 | ESMS 2 |
| 493 | 94698139 | DC0024-EFA0 | 09/09/2003 | Dace | 13.3 | ESMS 2 |
| 494 | 94698140 | DC0024-E58D | 09/09/2003 | Dace | 23 | ESMS 2 |
| 495 | 94698127 | DC0024-E4D2 | 09/09/2003 | Dace | 14.7 | ESMS 2 |
| 496 | 94698128 | DC0024-F0DE | 09/09/2003 | Dace | 14.2 | ESMS 2 |
| 497 | 94698141 | DC0024-F228 | 09/09/2003 | Dace | 13.8 | ESMS 2 |
| 498 | 94698146 | DC0024-D404 | 09/09/2003 | Dace | 14.5 | ESMS 2 |
| 499 | 94698130 | DC0024-E171 | 09/09/2003 | Dace | 13.2 | ESMS 2 |
| 500 | 94698024 | DC0024-C825 | 09/09/2003 | Dace | 14.7 | ESMS 2 |
| 501 | 94698142 | DC0024-D8C8 | 09/09/2003 | Dace | 14.6 | ESMS 2 |
| 502 | 94698126 | DC0024-EF87 | 09/09/2003 | Dace | 13.5 | ESMS 2 |
| 503 | 94698143 | DC0024-CCC4 | 09/09/2003 | Dace | 18.2 | ESMS 2 |
| 504 | 94698124 | DC0024-D9B3 | 09/09/2003 | Gudgeon | 12.9 | ESMS 2 |
| 505 | 94698010 | DC0024-ED3E | 09/09/2003 | Eel | 83.2 | ESMS 2 |
| 506 | 94698120 | DC0024-CB68 | 09/09/2003 | Eel | 43.8 | ESMS 2 |
| 507 | 94698123 | DC0024-DB77 | 09/09/2003 | Eel | 51.5 | ESMS 2 |
| 508 | 94698122 | DC0024-D5BA | 09/09/2003 | Dace | 16.3 | ESMS 2 |
| 509 | 94698052 |  | 10/09/2003 | Pike | 17 | Luckford Lake 2 |
| 510 | 94698051 |  | 10/09/2003 | Pike | 23.7 | Luckford Lake 2 |
| 511 | 94698048 |  | 10/09/2003 | Pike |  | Luckford Lake 4 |
| 512 | 94698720 | DC001B-F849 | 17/03/2003 | Pike | 36.9 | Flood Relief 2 |
| 513 | 94698049 |  | 10/09/2003 | Pike | 38.7 | Luckford Lake 4 |
| 514 | 113943625 |  | 05/12/2003 | Pike | 19.5 | Luckford Lake 1 |
| 515 | 113943628 |  | 05/12/2003 | Pike | 66.5 | Luckford Lake 2 |
| 516 | 113943626 |  | 05/12/2003 | Pike | 63 | Luckford Lake 2 |
| 517 |  | DC0024-F03B | 05/12/2003 | Sea trout | 38 | Luckford Lake 3 |
| 518 | 122451034 | DC0024-CA89 | 05/12/2003 | Pike | 9.9 | Luckford Lake 3 |
| 519 |  | DC001C-00CA | 05/12/2003 | Sea trout | 60 | Luckford Lake 3 |
| 520 | 94698057 |  | 05/12/2003 | Dace | 21.1 | Luckford Lake 4 |
| 521 | 113945695 |  | 05/12/2003 | Roach | 24.9 | Luckford Lake 4 |
| 522 |  |  | 05/12/2003 | Roach | 17.9 | Luckford Lake 4 |
| 523 | 113945694 |  | 05/12/2003 | Dace | 17 | Luckford Lake 4 |
| 524 | 113945692 |  | 05/12/2003 | Dace | 17.7 | Luckford Lake 4 |
| 525 | 113945691 |  | 05/12/2003 | Roach | 19.3 | Luckford Lake 4 |
| 526 | 113945690 |  | 05/12/2003 | Dace | 16.7 | Luckford Lake 4 |
| 527 | 113945689 |  | 05/12/2003 | Roach | 16.2 | Luckford Lake 4 |
| 528 | 113945688 |  | 05/12/2003 | Roach | 17.1 | Luckford Lake 4 |
| 529 | 113945687 |  | 05/12/2003 | Dace | 15.9 | Luckford Lake 4 |
| 530 | 113945686 |  | 05/12/2003 | Roach | 18.2 | Luckford Lake 4 |
| 531 | 113945685 |  | 05/12/2003 | Dace | 15 | Luckford Lake 4 |
| 532 | 113945684 |  | 05/12/2003 | Roach | 14 | Luckford Lake 4 |
| 533 | 113945683 |  | 05/12/2003 | Roach | 14.2 | Luckford Lake 4 |
| 534 | 113945679 |  | 05/12/2003 | Roach |  | Holme Bridge 3 |
| 535 | 113945678 |  | 05/12/2003 | Roach | 14.5 | Holme Bridge 3 |
| 536 | 113945677 |  | 05/12/2003 | Roach | 15.8 | Holme Bridge 3 |
| 537 | 113945676 |  | 05/12/2003 | Pike | 22.2 | Holme Bridge 4 |
| 538 | 113945675 |  | 05/12/2003 | Pike | 25.4 | Holme Bridge 4 |


| 539 | 113945671 |  | 08/12/2003 | Pike | 19.5 | Railway 3 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 540 | 113945672 |  | 08/12/2003 | Pike | 19.3 | Railway 3 |
| 541 | 113945670 |  | 08/12/2003 | Pike | 31.5 | Railway 4 |
| 542 | 113945669 |  | 08/12/2003 | Pike | 19 | Railway 4 |
| 543 | 113945668 |  | 08/12/2003 | Pike | 18.5 | Railway 4 |
| 544 | 113945665 | DC0024-E7F9 | 08/12/2003 | Pike | 371 | Flood Relief 2 |
| 545 | 113945664 | DC0024-C982 | 08/12/2003 | Pike | 45.2 | Flood Relief 2 |
| 546 | 113945667 | DC0024-8473 | 08/12/2003 | Pike | 27.4 | Flood Relief 2 |
| 547 | 113945663 |  | 08/12/2003 | Roach | 17.3 | Goldsacs |
| 548 | 113945662 |  | 08/12/2003 | Roach | 22.5 | Goldsacs 4 |
| 549 | 113945661 |  | 08/12/2003 | Dace | 20.4 | Goldsacs 4 |
| 550 | 113945659 |  | 08/12/2003 | Eel | 31.5 | ESMS 1 |
| 551 | 113945660 |  | 08/12/2003 | Eel | 31.5 | ESMS 1 |
| 552 | 94698075 | DC001B-F379 | 09/09/2003 | Trout | 23.3 | ESMS 3 |
| 553 | 113945658 |  | 08/12/2003 | Eel | 41 | ESMS 1 |
| 554 | 94698345 | DC001C-6D6D | 15/04/2003 | Pike | 62 | ESMS |
| 555 | 94609082 | DC001B-EF48 | 08/12/2003 | Trout | 19.4 | ESMS 3 |
| 556 | 113945657 |  | 08/12/2003 | Eel | 38 | ESMS 3 |
| 557 | 113945656 |  | 08/12/2003 | Eel | 37 | ESMS 3 |
| 558 | 113945655 |  | 08/12/2003 | Eel | 38.5 | ESMS 3 |
| 559 | 113945654 |  | 08/12/2003 | Eel | 36.8 | ESMS 3 |
| 560 | 113945653 |  | 08/12/2003 | Eel | 31.5 | ESMS 3 |
| 561 | 94698050 |  | 09/09/2003 | Trout | 19.5 | ESMS 4 |
| 562 |  | DC0024-EB3F | 09/12/2003 | Pike | 10.9 | Rushton 1 |
| 563 | 122451030 | DC002B-174A | 08/12/2003 | Pike | 11.5 | Rushton 1 |
| 564 | 113945652 |  | 08/12/2003 | Pike | 24 | ESMS 2 |
| 565 |  | DC0024-7A22 | 09/12/2003 | Pike | 10.4 | ESMS 2 |
| 566 | 113945651 |  | 09/12/2003 | Pike | 22.5 | ESMS 2 |
| 567 | 113945650 |  | 09/12/2003 | Pike | 17.2 | ESMS 4 |
| 568 | 113945649 |  | 09/12/2003 | Pike | 14.7 | Rushton Extra |
| 569 |  | DC0024-928A | 09/12/2003 | Pike | 122.2 | Rushton Extra |
| 570 | 35471204 | DC002A-F470 | 09/12/2003 | Pike | 12.1 | Rushton Extra |
| 571 |  | DC0024-CB21 | 09/12/2003 | Pike | 12.3 | Rushton Extra |
| 572 |  | DC0024-8516 | 09/12/2003 | Pike | 12.4 | Rushton Extra |
| 573 | 113945648 |  | 07/01/2004 | Pike | 38 | ESMS above lilypool |
| 574 | 113945647 |  | 07/01/2004 | Pike | 43.3 | DS Cattle drink |
| 575 | 113945646 |  | 07/01/2004 | Pike | 40.8 | DS Cattle drink |
| 576 | 113945645 |  | 07/01/2004 | Dace | 23.7 | ESMS above weir |
| 577 | 113945642 |  | 07/01/2004 | Dace | 23.1 | ESMS above weir |
| 578 | 113945643 |  | 07/01/2004 | Dace | 23.8 | ESMS eel trap |
| 579 | 113945644 |  | 07/01/2004 | Dace | 24.2 | ESMS eel trap |
| 580 | 113945641 |  | 07/01/2004 | Dace | 24.1 | ESMS eel trap |
| 581 | 113945639 |  | 07/01/2004 | Dace | 24.2 | ESMS eel trap |
| 582 | 113945632 |  | 07/01/2004 | Dace | 22.6 | ESMS eel trap |
| 583 | 113945633 |  | 07/01/2004 | Roach | 18 | ESMS eel trap |
| 584 | 113945631 |  | 07/01/2004 | Dace | 22.1 | ESMS eel trap |
| 585 | 113945635 |  | 07/01/2004 | Dace | 16.8 | ESMS eel trap |
| 586 | 113945636 |  | 07/01/2004 | Roach | 15.6 | ESMS eel trap |
| 587 | 113945637 |  | 07/01/2004 | Roach | 17.3 | ESMS eel trap |
| 588 | 113945634 |  | 07/01/2004 | Roach | 15.8 | ESMS eel trap |
| 589 | 113945629 |  | 07/01/2004 | Dace | 15.9 | ESMS eel trap |
| 590 | 113945630 |  | 07/01/2004 | Dace | 14.5 | ESMS eel trap |
| 591 | 113945623 |  | 07/01/2004 | Dace | 16.4 | ESMS eel trap |
| 592 | 113945625 |  | 07/01/2004 | Dace | 15.6 | ESMS eel trap |


| 593 | 113945638 |  | 07/01/2004 | Roach | 16.2 | ESMS eel trap |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 594 | 113945628 |  | 07/01/2004 | Roach | 15 | ESMS eel trap |
|  |  |  |  |  |  | Main River Frome |
| 595 | 124086110 | DC002B-O8O8 | 23/02/2004 | Pike | 73.4 | Weirpool |
|  |  |  |  |  |  | Main River Frome |
| 596 | 124086111 | DC002A-FC2D | 23/02/2004 | Pike | 59.4 | Weirpool-T4 |
|  |  |  |  |  |  | Main River Frome |
| 597 | 124086112 | DC002A-FCA2 | 23/02/2004 | Pike | 54.7 | Weirpool - T4 |
|  |  |  |  |  |  | Main River Frome |
| 598 | 94698339 | DC001C-AB91 | 16/04/2003 | Pike | 54.5 | Weirpool |
|  |  |  |  |  |  | Main River Frome |
| 599 | 113943630 | DC002A-FCE0 | 23/02/2004 | Pike | 40 | Weirpool-T4 |
| 600 | 124086114 | DC0024-EE51 | 23/02/2004 | Pike | 80 | Main River Frome T3 |
| 601 | 124086116 | DC002A-FB7D | 23/02/2004 | Pike | 91 | Main River Frome T9 Main River Frome |
| 602 | 94698340 | DC001C-85C3 |  | Pike | 57.2 | Weirpool |
| 603 | 124086117 | DC002B-O3B3 | 23/02/2004 | Pike | 69.5 | Main River Frome T9 |
| 604 | 124086118 | DC002B-1BDF | 23/02/2004 | Pike | 57.5 | Main River Frome T7- T9 |
| 605 | 124086119 | DC002B-0C0B | 23/02/2004 | Pike | 61.2 | Main River Frome T7- T9 |
| 606 | 124086120 | DC002B-12F5 | 23/02/2004 | Pike | 59.8 | Main River Frome T15 |
| 607 | 124086121 | DC002A-F201 | 23/02/2004 | Pike | 57.5 | Main River Frome T15 |
| 608 | 124086122 | DC002B-1CF7 | 23/02/2004 | Pike | 70 | Main River Frome T15 |
| 609 | 124086123 | DC002B-0297 |  | Pike | 72.8 | Main River Frome T15 |
| 610 | 124086124 | DC002B-11D6 | 23/02/2004 | Pike | 52.5 | Main River Frome T15 |
| 611 | 124086210 | DC002B-0CD8 | 24/02/2004 | Pike | 58.5 | Main River Frome T20 <br> Main River Frome T22 - |
| 612 | 124086125 | DC002A-FA15 | 24/02/2004 | Pike | 69.6 | T25 |
|  |  |  |  |  |  | Main River Frome T22- |
| 613 | 124086126 | DC002A-F50B | 24/02/2004 | Pike | 58.9 | T25 |
|  |  |  |  |  |  | Main River Frome T22- |
| 614 |  | DC002B-16FC | 24/02/2004 | Pike | 11.6 | T25 |
|  |  |  |  |  |  | Main River Frome T22- |
| 615 | 124086127 | DC002B-0D10 | 24/02/2004 | Pike | 102 | T25 |
|  |  |  |  |  |  | Main River Frome T25- |
| 616 | 124086115 |  | 24/02/2004 | Perch | 34.2 | T30 |
|  |  |  |  |  |  | Main River Frome T25- |
| 617 | 124086128 |  | 24/02/2004 | Roach | 18.4 | T30 |
|  |  |  |  |  |  | Main River Frome MS - |
| 618 | 124086129 |  |  | Pike | 67.3 | Rushton |
|  |  |  |  |  |  | Main River Frome MS - |
| 619 | 124086131 |  | 24/02/2004 | Pike | 69.2 | Rushton |
|  |  |  |  |  |  | Main River Frome MS - |
| 620 | 124086130 |  | 24/02/2004 | Pike | 90 | Rushton |
|  |  |  |  |  |  | Main River Frome MS - |
| 621 | 124086133 |  | 24/02/2004 | Pike | 64 | Rushton |
| 622 | 94698715 |  | 18/03/2003 | Pike | 48 | Rushton 4 |
|  |  |  |  |  |  | Main River Frome MS - |
| 623 | 124086134 |  | 24/02/2004 | Pike | 23.6 | Rushton |
| 624 | 124086135 |  | 24/02/2004 | Pike | 69.5 | Main River Frome Oxbow |
| 625 | 94698716 |  | 18/03/2003 | Pike | 51.5 | Rushton 4 |
| 626 | 124086136 |  | 24/02/2004 | Pike | 91.9 | Main River Frome Oxbow |
|  |  |  | 24/02/2004 | Pike | 68.5 | Main River Frome Holme Bridge |
| 628 | 94698719 |  | 18/03/2003 | Pike | 65 | ESMS 1 |
| 629 | 94698718 | DC001C-793C | 18/03/2003 | Pike | 61.5 | ESMS 3 |
| 630 | 98698118 | DC0024-DCF4 | 09/09/2003 | Gudgeon | 12.7 | ESMS 2 |
| 631 | 94698119 | DC0024-E533 | 09/09/2003 | Dace | 15.2 | ESMS 2 |
| 632 | 94698117 | DC0024-E0DD | 09/09/2003 | Dace | 16 | ESMS 2 |


| 633 | 94698113 | DC0024-EC36 | 09/09/2003 | Dace | 12.6 | ESMS 2 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 634 | 94698112 | DC0024-DCB9 | 09/09/2003 | Dace | 17.9 | ESMS 2 |
| 635 | 94698116 | DC0024-09BB | 09/09/2003 | Dace | 15.2 | ESMS 2 |
| 636 | 94698115 | DC0024-F2AF | 09/09/2003 | Dace | 15.7 | ESMS 2 |
| 637 | 94698105 | DC0024-DC59 | 09/09/2003 | Dace | 15.1 | ESMS 2 |
| 638 | 94698109 | DC0024-C8DA | 09/09/2003 | Dace | 13.5 | ESMS 2 |
| 639 | 94698106 | DC0024-DE01 | 09/09/2003 | Dace | 12.4 | ESMS 2 |
| 640 | 94698111 | DC0024-CE9A | 09/09/2003 | Dace | 13.4 | ESMS 2 |
| 641 | 94698114 | DC0024-ED0F | 09/09/2003 | Dace | 14.5 | ESMS 2 |
| 642 | 94698110 | DC0024-CA18 | 09/03/2003 | Dace | 14.8 | ESMS 2 |
| 643 | 94698104 | DC0024-DEE4 | 09/09/2003 | Dace | 14.4 | ESMS 2 |
| 644 | 94698108 | DC0024-CDEF | 09/09/2003 | Dace | 12.9 | ESMS 2 |
| 645 | 94698103 | DC001B-E952 | 09/09/2003 | Gudgeon | 13.1 | ESMS 2 |
| 646 | 94698094 | DC001C-0456 | 09/09/2003 | Pike | 16.3 | ESMS 2 |
| 647 | 94698097 | DC001C-0B51 | 09/09/2003 | Dace | 14.3 | ESMS 2 |
| 648 | 94698095 | DC001B-F171 | 09/09/2003 | Dace | 14.7 | ESMS 2 |
| 649 | 94698099 | DC001B-EBCF | 09/09/2003 | Dace | 16.2 | ESMS 2 |
| 650 | 94698101 | DC001B-F576 | 09/09/2003 | Dace | 14.7 | ESMS 2 |
| 651 | 94698096 | DC001B-F587 | 09/09/2003 | Dace | 19.2 | ESMS 2 |
| 652 | 94698098 | DC001B-E56F | 09/09/2003 | Dace | 12.3 | ESMS 2 |
| 653 | 94698102 | DC001C-0BA7 | 09/09/2003 | Eel | 50 | ESMS 2 |
| 654 | 94698093 | DC001C-0457 | 09/09/2003 | Gudgeon | 13.4 | ESMS 2 |
| 655 | 94698091 | DC001B-EE53 | 09/09/2003 | Eel | 37 | ESMS 2 |
| 656 | 94698092 | DC001C-0774 | 09/09/2003 | Eel | 48 | ESMS 2 |
| 657 | 94698089 | DC0024-D90D | 09/09/2003 | Eel | 37.5 | ESMS 2 |
| 658 | 94698084 | DC001C-0406 | 09/09/2003 | Eel | 35 | ESMS 2 |
| 659 | 94698087 | DC0024-DBE8 | 09/09/2003 | Eel | 36 | ESMS 2 |
| 660 | 94698086 | DC001B-E95C | 09/09/2003 | Eel | 35.5 | ESMS 2 |
| 661 | 94698085 | DC001B-FCB6 | 09/09/2003 | Eel | 36.8 | ESMS 2 |
| 662 | 94698090 | DC001B-E6F3 | 09/09/2003 | Eel | 37.6 | ESMS 2 |
| 663 | 94698088 | DC001C-06BA | 09/09/2003 | Eel | 36 | ESMS 2 |
| 664 | 94698073 | DC001B-FF59 | 09/09/2003 | Trout | 21.9 | ESMS 3 |
| 665 | 94698082 | DC001B-E7AE | 09/09/2003 | Trout | 19.4 | ESMS 3 |
| 666 | 94698068 | DC001C-0AF8 | 09/09/2003 | Eel | 36 | ESMS 3 |
| 667 | 94698080 | DC001B-E744 | 09/09/2003 | Dace | 13.3 | ESMS 3 |
| 668 | 94698069 | DC001C-0BC9 | 09/09/2003 | Eel | 43 | ESMS 3 |
| 669 | 94698074 | DC001C-0AEB | 09/09/2003 | Eel | 43.8 | ESMS 3 |
| 670 | 94698063 | DC001B-EFEA | 09/09/2003 | Eel | 45 | ESMS 3 |
| 671 | 94698062 | DC001B-FF22 | 09/09/2003 | Eel | 51 | ESMS 3 |
| 672 | 94698078 | DC001C-0BD1 | 09/09/2003 | Eel | 43 | ESMS 3 |
| 673 | 94698071 | DC001B-E4DD | 09/09/2003 | Eel | 42 | ESMS 3 |
| 674 | 94698070 | DC001B-E3BB | 09/09/2003 | Eel | 44 | ESMS 3 |
| 675 | 94698065 | DC0024-CD6C | 09/09/2003 | Eel | 54 | ESMS 3 |
| 676 | 94698066 | DC0024-DC82 | 09/09/2003 | Eel | 41 |  |
| 677 | 94698064 |  | 09/09/2003 | Eel | 46.5 | ESMS 3 |
| 678 | 94698059 | DC0024-0992 | 09/09/2003 | Eel | 31 | ESMS 3 |
| 679 | 94698056 | DC0024-DFFC | 09/09/2003 | Eel | 35.5 | ESMS 3 |
| 680 | 94698058 | DC0024-EDAB | 09/09/2003 | Eel | 58 | ESMS 3 |
| 681 | 94698076 |  | 09/09/2003 | Eel | 35 | ESMS 3 |
| 682 | 94698072 |  | 09/09/2003 | Eel | 33 | ESMS 3 |
| 683 | 94698060 |  | 09/09/2003 | Eel | 37.5 | ESMS 3 |
| 684 | 94698081 |  | 09/09/2003 | Eel | 42.5 | ESMS 3 |
| 685 | 94698067 |  | 09/09/2003 | Eel | 37 | ESMS 3 |
| 686 | 94698079 |  | 09/09/2003 | Eel | 37 | ESMS 3 |


| 687 | 94698077 |  | 09/09/2003 | Eel |
| :---: | :---: | :---: | :---: | :---: |
| 688 | 94698061 |  | 09/09/2003 | Trout |
| 689 | 94698055 |  | 09/09/2003 | Eel |
| 690 | 94698054 |  | 09/09/2003 | Eel |
| 694 | 124086194 | DC002B-145A | 04/03/2004 | Dace |
| 695 | 124086205 | DC002B-1CC2 | 04/03/2004 | Roach |
| 696 | 124086208 | DC002B-05D1 | 04/03/2004 | Dace |
| 697 | 124086196 | DC002A-F3A8 | 04/03/2004 | Dace |
| 698 | 124086204 | DC002B-12B7 | 04/03/2004 | Roach |
| 700 | 124086200 | DC002B-0A98 | 04/03/2004 | Roach |
| 701 | 124086193 | DC002B-0521 | 04/03/2004 | Roach |
| 702 | 124086198 | DC002B-12B6 | 04/03/2004 | Dace |
| 703 | 124086201 | DC002B-15B9 | 04/03/2004 | Dace |
| 704 | 124086206 | DC002A-FC7D | 04/03/2004 | Roach |
| 705 | 124086199 | DC002A-FA48 | 04/03/2004 | Dace |
| 706 | 124086197 | DC002B-OD63 | 04/03/2004 | Dace |
| 707 | 124086202 | DC002B-15C6 | 04/03/2004 | Pike |
| 708 | 124086209 | DC002B-1684 | 04/03/2004 | Dace |
| 709 | 124086195 | DC002B-1478 | 04/03/2004 | Dace |
| 710 | 124086186 | DC002A-F2AC | 04/03/2004 | Dace |
| 711 | 124086192 | DC002B-1AE7 | 04/03/2004 | Roach |
| 712 | 124086190 | DC002A-F43A | 04/03/2004 | Roach |
| 713 | 124086189 | DC002A-F249 | 04/03/2004 | Roach |
| 715 | 124086191 | DC002A-F355 | 04/03/2004 | Roach |
| 716 | 124086171 | DC002B-06CP | 04/03/2004 | Roach |
| 717 | 124086166 | DC002A-F6F9 | 04/03/2004 | Dace |
| 718 | 124086178 | DC002B-1846 | 04/03/2004 | Dace |
| 719 | 124086175 | DC002A-F1EA | 04/03/2004 | Dace |
| 721 | 124086177 | DC002B-1ADD | 04/03/2004 | Roach |
| 722 | 124086169 | DC002B-0C32 | 04/03/2004 | Roach |
| 724 | 124086173 | DC002A-F4F4 | 04/03/2004 | Dace |
| 725 | 124086167 | DC002A-FB06 | 04/03/2004 | Dace |
| 726 | 124086174 | DC002A-F670 | 04/03/2004 | Dace |
| 728 | 124086142 | DC002B-169B | 04/03/2004 | Dace |
| 729 | 124086138 | DC002A-FF0C | 04/03/2004 | Dace |
| 730 | 124086139 | DC002B-1B21 | 04/03/2004 | Dace |
| 731 | 124086143 | DC002B-0358 | 04/03/2004 | Dace |
| 732 | 124086140 | DC002B-060C | 04/03/2004 | Dace |
| 733 | 124086147 | DC002B-161F | 04/03/2004 | Dace |
| 735 | 124086146 | DC002B-1AEC | 04/03/2004 | Dace |
| 737 | 124086144 | DC002B-149D | 04/03/2004 | Dace |
| 738 | 124086145 | DC002B-0443 | 04/03/2004 | Dace |
| 739 | 124086148 | DC002B-008F | 04/03/2004 | Dace |
| 740 | 124086141 | DC002A-F699 | 04/03/2004 | Roach |
| 747 | 124086163 | DC0024-E8B3 |  | Roach |
| 748 | 124086161 | DC002B-0336 | 04/03/2004 | Dace |
| 749 |  | DC002A-FD22 | 04/03/2004 | Roach |
| 752 | 124086162 | DC002A-FFFA | 04/03/2004 | Dace |
| 753 | 124086159 | DC002A-FA72 | 04/03/2004 | Dace |
| 754 | 124086160 | DC002A-F51B | 04/03/2004 | Dace |
| 755 | 124086150 | DC002A-FB3A | 04/03/2004 | Roach |
| 757 | 124086149 | DC002B-0917 | 04/03/2004 | Dace |
| 758 | 124086158 | DC002A-FB55 | 04/03/2004 | Dace |
| 759 | 124086184 | DC002B-1303 | 04/03/2004 | Dace |


| 33 | ESMS 3 |
| :---: | :---: |
| 31.5 | ESMS 4 |
| 34.5 | ESMS 4 |
| 49 | ESMS 4 |
| 21.8 | ESMS ds eel poo |
| 15.6 | ESMS ds eel poo |
| 15.5 | ESMS ds eel |
| 15.6 | ESMS ds eel p |
| 13.5 | ESMS eel pool |
| 13.2 | ESMS eel pool |
| 22.5 | ESMS eel pool |
| 20.9 | ESMS eel pool |
| 18.3 | ESMS eel pool |
| 18.7 | ESMS eel pool |
| 16.6 | ESMS eel pool |
| 14.2 | ESMS eel pool |
| 41.3 | ESMS eel pool |
| 20.5 | ESMS eel pool |
| 23.9 | ESMS eel pool |
| 17 | ESMS eel pool |
| 17.2 | ESMS eel pool |
| 13.7 | ESMS eel pool |
| 16.9 | ESMS eel pool |
| 13.8 | ESMS eel pool |
| 14.2 | ESMS eel pool |
| 16 | ESMS eel pool |
| 16.5 | ESMS eel pool |
| 18.1 | ESMS eel pool |
| 14.9 | ESMS eel pool |
| 16.3 | ESMS eel pool |
| 13.5 | ESMS eel pool |
| 13.5 | ESMS eel pool |
| 21.3 | ESMS eel pool |
| 16.1 | ESMS eel pool |
| 22.8 | ESMS eel pool |
| 16.5 | ESMS eel pool |
| 16 | ESMS eel pool |
| 14.6 | ESMS eel pool |
| 15.9 | ESMS eel pool |
| 14.6 | ESMS eel pool |
| 13.5 | ESMS eel pool |
| 13.4 | ESMS eel pool |
| 12.9 | ESMS eel pool |
| 13.4 | ESMS eel pool |
| 15.4 | ESMS eel pool |
| 14.6 | ESMS eel pool |
| 12.7 | ESMS eel pool |
| 15.5 | ESMS eel pool |
| 13 | ESMS eel pool |
| 14 | ESMS eel pool |
| 14.1 | ESMS eel pool |
| 14.4 | ESMS eel pool |
| 13.1 | ESMS eel pool |
| 21.9 | ESMS eel pool |


| 760 | 124086207 | DC002A-FE06 | 04/03/2004 | Dace | 22.3 | ESMS eel pool |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 761 | 124086182 | DC002B-1D47 | 04/03/2004 | Dace | 15.7 | ESMS eel pool |
| 763 | 124086185 | DC002B-0946 | 04/03/2004 | Dace | 17.3 | ESMS eel pool |
| 764 | 124086188 | DC0020-0D28 | 04/03/2004 | Dace | 16.3 | ESMS eel pool |
| 765 | 124086180 | DC002B-0201 | 04/03/2004 | Dace | 15.8 | ESMS eel pool |
| 766 | 124086187 | DC002B-1B8E | 04/03/2004 | Dace | 14.4 | ESMS eel pool |
| 768 | 124086179 | DC002B-1268 | 04/03/2004 | Dace | 15.7 | ESMS eel pool |
| 769 | 124086181 | DC002B-05FB | 04/03/2004 | Dace | 13.2 | ESMS eel pool |
| 770 | 124086183 | DC002B-0F5B | 04/03/2004 | Roach | 15.7 | ESMS eel pool |
| 771 | 124086168 | DC002B-1560 | 04/03/2004 | Dace | 21.5 | ESMS eel pool |
| 772 | 124086176 | DC002A-FF15 | 04/03/2004 | Dace | 16.5 | ESMS eel pool |
| 773 | 124086170 | DC002B-044F | 04/03/2004 | Dace | 13.8 | ESMS eel pool |
| 775 | 124086172 | DC002B-1075 | 04/03/2004 | Dace | 15.4 | ESMS eel pool Main River Frome |
| 776 | 113943629 | DC002B-15EC | 16/03/2004 | Pike | 70.9 | Weirpool - T4 Main River Frome |
| 777 | 113945666 | DC002A-F712 | 16/03/2004 | Pike | 59.8 | Weirpool - T4 Main River Frome |
| 778 | 113943657 | DC002B-0F4C | 16/03/2004 | Pike | 65.2 | Weirpool - T4 <br> Main River Frome 150 m |
| 779 | 113945615 |  | 16/03/2004 | Dace | 19.7 | above Oxbow Main River Frome 150 m |
| 780 | 113945616 |  | 16/03/2004 | Dace | 26 | above Oxbow |
| 781 | 113945614 |  | 16/03/2004 | Dace | 26.1 | Flood relief entrance |
| 782 | 113945613 |  | 16/03/2004 | Dace | 26.5 | Flood relief entrance |
| 783 | 113945612 |  | 16/03/2004 | Dace | 22.9 | Flood relief entrance |
| 784 | 113945611 |  | 16/03/2004 | Dace | 25.5 | Flood relief entrance |
| 785 | 113945610 |  | 16/03/2004 | Dace | 21.8 | Flood relief entrance |
| 786 | 113945609 | DC002B-1C1F | 16/03/2004 | Dace | 21.1 | Flood relief entrance |
| 787 | 113945608 | DC002A-F7BA | 16/03/2004 | Dace | 21.2 | Flood relief entrance |
| 788 | 113945607 | DC002B-0C14 | 16/03/2004 | Dace | 19.4 | Flood relief entrance |
| 789 | 113945606 | DC0024-E1D7 | 16/03/2004 | Dace | 22.6 | Flood relief entrance |
| 790 | 113945605 | DC002A-F5FB | 16/03/2004 | Dace | 21.7 | Flood relief entrance |
| 791 | 113945603 | DC002B-00B7 | 16/03/2004 | Dace | 20.8 | Flood relief entrance |
| 792 | 113945602 | DC002A-F467 | 16/03/2004 | Dace | 20.5 | Flood relief entrance |
| 793 | 113945600 | DC002A-F7AE | 16/03/2004 | Dace | 22.3 | ds road bridge |
| 794 | 113945601 | DC0024-7DCE | 16/03/2004 | Dace | 21.5 | ds road bridge |
| 795 | 113945599 | DC002B-0E6A | 16/03/2004 | Dace | 22.8 | ds road bridge |
| 796 | 113945598 | DC002B-080F | 16/03/2004 | Dace | 21.7 | ds road bridge Main River Frome |
| 797 | 113945597 |  | 16/03/2004 | Dace | 23.2 | Weirpool-T4 Main River Frome |
| 798 | 113945596 |  | 16/03/2004 | Dace | 23.8 | Weirpool - T4 Main River Frome |
| 799 | 113945595 |  | 16/03/2004 | Dace | 22.9 | Weirpool - T4 Main River Frome |
| 800 | 113945594 |  | 16/03/2004 | Dace | 20.3 | Weirpool-T4 Main River Frome |
| 801 | 113945593 |  | 16/03/2004 | Dace | 20.8 | Weirpool - T4 |
| 802 | 113945592 |  | 17/03/2004 | Dace | 24.9 | ESMS 2 |
| 803 | 113945591 |  | 17/03/2004 | Dace | 27.1 | ESMS 2 |
| 804 | 113945590 |  | 17/03/2004 | Dace | 22.4 | ESMS 2 |
| 805 | 113945589 |  | 17/03/2004 | Dace | 23.2 | ESMS 2 |
| 806 | 113945588 |  | 17/03/2004 | Dace | 23.2 | ESMS 2 |
| 807 | 113943735 |  | 17/03/2004 | Dace | 24.2 | ESMS 2 |
| 808 | 113943741 |  | 17/03/2004 | Dace | 22.5 | ESMS 2 |


| 809 | 113943734 |  | 17/03/2004 | Dace | 23.9 | ESMS 2 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 810 | 113943736 |  | 17/03/2004 | Dace | 20.4 | ESMS 2 |
| 811 | 113943742 |  | 17/03/2004 | Dace | 20.6 | ESMS 2 |
| 812 | 113943738 |  | 17/03/2004 | Dace | 20.9 | ESMS 2 |
| 813 | 113943740 |  | 17/03/2004 | Dace | 25 | ESMS 2 |
| 814 | -113943743 |  | 17/03/2004 | Dace | 19.4 | ESMS 2 |
| 815 | 113943737 |  | 17/03/2004 | Dace | 20.2 | ESMS 2 |
| 816 | 113943739 |  | 17/03/2004 | Dace | 21.4 | ESMS 2 |
| 817 | 113943733 |  | 17/03/2004 | Dace | 18 | ESMS 2 |
| 818 | 113943732 |  | 17/03/2004 | Dace | 17.4 | ESMS 2 |
| 819 | 113943731 |  | 17/03/2004 | Dace | 18.5 | ESMS 2 |
| 820 | 113943730 |  | 17/03/2004 | Dace | 20.5 | ESMS 3 |
| 821 | 113943729 |  | 17/03/2004 | Dace | 23.7 | ESMS 3 |
| 822 | 113943728 |  | 17/03/2004 | Dace | 18.7 | ESMS 3 |
| 823 | 113943727 |  | 17/03/2004 | Pike | 14.8 | Rushton 1 |
| 824 | 113943726 |  | 17/03/2004 | Pike | 14.2 | Rushton 2 |
| 826 | 0 | DC002A-FC1D | 17/03/2004 | Pike | 10.3 | Rushton 2 |
| 827 | 113943725 |  | 17/03/2004 | Pike | 42.5 | Rushton 4 |
| 828 | 113943724 |  | 17/03/2004 | Pike | 13.5 | Rushton 4 |
| 829 | 35471184 | DC002A-F234 | 17/03/2004 | Pike | 12.8 | Rushton 4 |
| 830 | 35471228 | DC002B-130A | 17/03/2004 | Pike | 12.6 | Rushton 4 |
| 831 | 113943723 |  | 18/03/2004 | Dace | 22.5 | Luckford Lake 1 |
| 832 | 113943722 |  | 18/03/2004 | Perch | 25.9 | Luckford Lake 3 |
| 833 | 113943714 |  | 18/03/2004 | Dace | 25 | Luckford Lake 4 |
| 834 | 113943712 |  | 18/03/2004 | Dace | 25.8 | Luckford Lake 4 |
| 835 | 113943716 |  | 18/03/2004 | Dace | 21 | Luckford Lake 4 |
| 836 | 113943715 |  | 18/03/2004 | Roach | 18.9 | Luckford Lake 4 |
| 837 | 113943713 |  | 18/03/2004 | Roach | 16.7 | Luckford Lake 4 |
| 838 | 113943719 |  | 18/03/2004 | Roach | 18.9 | Luckford Lake 4 |
| 839 | 113943717 |  | 18/03/2004 | Roach | 17.8 | Luckford Lake 4 |
| 840 | 113943720 |  | 18/03/2004 | Roach | 21.8 | Luckford Lake 4 |
| 841 | 113943721 |  | 18/03/2004 | Dace | 18.7 | Luckford Lake 4 |
| 842 | 113943660 |  | 18/03/2004 | Roach | 17.3 | Luckford Lake 4 |
| 843 | 113943663 |  | 18/03/2004 | Pike | 22.4 | Luckford Lake 4 |
| 844 | 113943661 |  | 18/03/2004 | Perch | 22.4 | Luckford Lake 4 |
| 845 | 113943662 |  | 18/03/2004 | Roach | 20.6 | Luckford Lake 4 |
| 846 | 113943658 |  | 18/03/2004 | Dace | 17.6 | Luckford Lake 4 |
| 847 | 113943659 |  | 18/03/2004 | Roach | 16.5 | Luckford Lake 4 |
| 848 | 113943665 |  | 18/03/2004 | Dace | 15 | Luckford Lake 4 |
| 849 | 94698042 |  | 17/03/2004 | Roach | 16.8 | ESMS |
| 850 | 94698039 |  | 17/03/2004 | Pike | 16.5 | ESMS |
| 851 | 94698041 |  | 17/03/2004 | Dace | 13.4 | ESMS |
| 852 | 94698043 |  | 17/03/2004 | Dace | 16.1 | ESMS |
| 853 | 94698047 |  | 17/03/2004 | Dace | 12.5 | ESMS |
| 854 | 94698044 |  | 17/03/2004 | Dace | 14 | ESMS |
| 855 | 94698045 |  | 17/03/2004 | Roach | 20.7 | ESMS |
| 856 | 94698046 |  | 17/03/2004 | Pike | 37.8 | ESMS |
| 857 | 94698040 |  | 17/03/2004 | Roach | 18.5 | ESMS |
| 859 | 113943615 |  | 17/03/2004 | Roach | 18.9 | ESMS |
| 860 | 113943616 |  | 17/03/2004 | Roach | 19.9 | ESMS |
| 861 | 113943614 |  | 17/03/2004 | Dace | 26.5 | ESMS |
| 862 | 113943613 |  | 17/03/2004 | Roach | 18 | ESMS |
| 863 | 113943619 |  | 17/03/2004 | Dace | 23.5 | ESMS |
| 864 | 113943612 |  | 17/03/2004 | Dace | 22.3 | ESMS |


| 865 | 113943623 |  | 17/03/2004 | Roach |
| :---: | :---: | :---: | :---: | :---: |
| 866 | 113943621 |  | 17/03/2004 | Roach |
| 867 | 113943620 |  | 17/03/2004 | Roach |
| 868 | 113943622 |  | 17/03/2004 | Roach |
| 869 | 113943624 |  | 17/03/2004 | Roach |
| 870 | 113943617 |  | 17/03/2004 | Dace |
| 872 | 124086157 | DC001B-F16F | 17/03/2003 | Pike |
| 873 | 124086155 | DC001B-EBF8 | 17/03/2003 | Pike |
| 874 | 94698728 | DC001B-EF17 | 17/03/2003 | Pike |
| 875 | 124086154 | DC002A-F3AE | 12/03/2004 | Pike |
| 876 | 124086153 | DC002B-05B7 | 12/03/2004 | Pike |
| 877 | 124086151 | DC0024-CA6A | 12/03/2004 | Perch |
| 878 | 124086152 |  | 12/03/2004 | Perch |
| 880 | 113943634 |  | 15/03/2004 | Pike |
| 881 | 113943635 |  | 15/03/2004 | Dace |
| 882 | 113943636 |  | 15/03/2004 | Dace |
| 883 | 113943637 |  | 15/03/2004 | Dace |
| 884 | 113943638 |  | 15/03/2004 | Dace |
| 885 | 113943640 |  | 15/03/2004 | Dace |
| 886 | 113943641 |  | 15/03/2004 | Dace |
| 887 | 113943642 |  | 15/03/2004 | Dace |
| 888 | 113943643 |  | 15/03/2004 | Dace |
| 889 | 113943653 |  | 15/03/2004 | Dace |
| 890 | 113943618 |  | 15/03/2004 | Dace |
| 891 | 113943644 |  | 15/03/2004 | Dace |
| 892 | 113943645 |  | 15/03/2004 | Dace |
| 893 | 113943646 |  | 15/03/2004 | Dace |
| 894 | 113943647 |  | 15/03/2004 | Dace |
| 895 | 113943648 |  | 15/03/2004 | Dace |
| 896 | 113943649 |  | 15/03/2004 | Roach |
| 897 | 113943650 |  | 15/03/2004 | Dace |
| 898 | 113943651 |  | 15/03/2004 | Dace |
| 899 | 113943652 |  | 15/03/2004 | Dace |
| 900 | 113943654 |  | 15/03/2004 | Dace |
| 901 | 113943655 |  | 15/03/2004 | Dace |
| 902 | 113943656 |  | 15/03/1941 | Dace |
| 903 | 113943633 |  | 15/03/2004 | Pike |
| 904 | 113943632 |  | 15/03/2004 | Pike |
| 905 | 113943668 |  | 04/06/2003 | Dace |
| 906 | 122451025 |  | 19/07/2004 | Pike |
| 907 | 122451109 |  | 19/07/2004 | Pike |
| 910 | 122451107 |  | 19/07/2004 | Pike |
| 911 |  | DC002A-FA53 | 20/07/2004 | Pike |
| 912 |  | DC002A-FD80 | 20/07/2004 | Pike |
| 913 | 122451153 |  | 20/07/2004 | Pike |
| 914 | 35471189 |  | 21/06/2004 | Eel |
| 915 | 35471244 |  | 21/06/2004 | Eel |
| 916 | 35471239 |  | 21/06/2004 | Eel |
| 917 | 35471233 |  | 22/06/2004 | Pike |
| 918 | 35471220 |  | 22/06/2004 | Pike |
| 919 | 35471203 |  | 22/06/2004 | Pike |
| 920 |  | DC002A-F300 | 22/06/2004 | Pike |
| 921 | 35471219 |  | 22/06/2004 | Eel |

19.7 ESMS
18.3 ESMS
16.2 ESMS
19.2 ESMS
17.7 ESMS

0 ESMS Eel Pool
11.9 Railway 1
11.8 Railway 2

24 Railway 3
26.8 Railway 3
21.5 Railway 4
22.9 Flood Relief
28.4 Flood Relief
63.5 Goldsacs 1
25.1 Goldsacs 3
21.2 Goldsacs 3
15.4 Goldsacs 3
28.7 Goldsacs 4
21.9 Goldsacs 4
24.8 Goldsacs 4
23.5 Goldsacs 4
20.2 Goldsacs 4
23.4 Goldsacs 4
24.1 Goldsacs 4
25.8 Goldsacs 4
34.2 Goldsacs 4
22.6 Goldsacs 4
22.3 Goldsacs 4
23.7 Goldsacs 4
18.9 Goldsacs 4
17.5 Goldsacs 4
20.9 Goldsacs 4
22.4 Goldsacs 4
16.4 Goldsacs 4
15.8 Goldsacs 4
19.7 Goldsacs 4
21.3 Holme Bridge 3
50.1 Holme Bridge 3
20.3 Wareham Quay
22.2
14.1 Rushton 2 ESMS
9.2 Flood Relief 2

10 Flood Relief 2 Main River Frome 80m
21 DS of Rushton
29 Railway 2
43 Railway 2
31 Railway 4
32.5 Rushton 1
20.1 Rushton
20.6 Rushton
6.5 Rushton 2

43 Rushton

| 922 | 113943671 |  | 22/06/2004 | Pike | 28.8 | Rushton 2 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 923 | 35471187 |  | 22/06/2004 | Pike | 39.5 | Holme Bridge 4 |
| 924 |  | DC0029-EB76 | 22/06/2004 | Pike | 6.3 | Holme Bridge 4 |
| 925 | 113943664 |  | 22/06/2004 | Pike | 32 | Holme Bridge |
| 926 | 35471195 |  | 23/06/2004 | Dace | 16.8 | ESMS 1 |
| 927 | 35471191 |  | 23/06/2004 | Dace | 12.1 | ESMS 1 |
| 928 | 35471202 |  | 23/06/2004 | Dace | 16.9 | ESMS 1 |
| 929 | 35471237 |  | 23/06/2004 | Dace | 14.9 | ESMS 1 |
| 930 | 35471230 |  | 23/06/2004 | Eel | 39.8 | ESMS 1 |
| 931 | 35471175 |  | 23/06/2004 | Eel | 47.5 | ESMS 1 |
| 932 | 35471185 |  | 23/06/2004 | Gudgeon | 13.3 | ESMS 1 |
| 933 | 35471120 |  | 23/06/2004 | Dace | 16.6 | ESMS 1 |
| 934 | 35471121 |  | 23/06/2004 | Dace | 15.6 | ESMS 1 |
| 935 | 35471123 |  | 23/06/2004 | Roach | 11.7 | ESMS 1 |
| 936 | 35471122 |  | 23/06/2004 | Eel | 37.9 | ESMS 1 |
| 937 | 35471174 |  | 23/06/2004 | Eel | 30.5 | ESMS 1 |
| 938 | 35471173 |  | 23/06/2004 | Trout | 20.4 | ESMS 1 |
| 939 | 35471177 |  | 23/06/2004 | Eel | 52 | ESMS 1 |
| 940 | 35471176 |  | 23/06/2004 | Dace | 16.6 | ESMS 2 |
| 941 | 35471167 |  | 23/06/2004 | Trout | 19.4 | ESMS 2 |
| 942 | 35471172 |  | 23/06/2004 | Dace | 14.2 | ESMS 2 |
| 943 | 35471162 |  | 23/06/2004 | Dace | 12.4 | ESMS 2 |
| 944 | 35471169 |  | 23/06/2004 | Roach | 12.2 | ESMS 2 |
| 945 | 35471163 |  | 23/06/2004 | Roach | 13.8 | ESMS 2 |
| 946 | 35471165 |  | 23/06/2004 | Eel | 31 | ESMS 2 |
| 947 | 35471156 |  | 23/06/2004 | Eel | 41.5 | ESMS 2 |
| 948 | 35471168 |  | 23/06/2004 | Roach | 14.2 | ESMS 2 |
| 949 | 35471160 |  | 23/06/2004 | Eel | 34 | ESMS 2 |
| 950 | 35471159 |  | 23/06/2004 | Eel | 45 | ESMS 2 |
| 951 | 35471158 |  | 23/06/2004 | Eel | 40.5 | ESMS 2 |
| 952 | 35471151 |  | 23/06/2004 | Eel | 33 | ESMS 2 |
| 953 | 35471150 |  | 23/06/2004 | Eel | 46 | ESMS 2 |
| 954 | 35471152 |  | 23/06/2004 | Eel | 34.5 | ESMS 2 |
| 955 | 35471155 |  | 23/06/2004 | Eel | 46.8 | ESMS 2 |
| 956 | 35471143 |  | 23/06/2004 | Eel | 51 | ESMS 3 |
| 957 | 35471139 |  | 23/06/2004 | Eel | 28.5 | ESMS 3 |
| 958 | 35471144 |  | 23/06/2004 | Eel | 45.8 | ESMS 3 |
| 959 | 35471138 |  | 23/06/2004 | Eel | 35.6 | ESMS |
| 960 | 35471132 |  | 23/06/2004 | Eel | 40.5 | ESMS 4 |
| 962 | 35471125 |  | 23/06/2004 | Eel | 30.5 | ESMS 4 |
| 963 | 35471128 |  | 23/06/2004 | Eel | 36 | ESMS 4 |
| 964 | 35471129 |  | 23/06/2004 | Eel | 33.9 | ESMS 4 |
| 965 | 35471124 |  | 23/06/2004 | Eel | 29 | ESMS 4 |
| 966 | 35471136 |  | 23/06/2004 | Eel | 33.5 | ESMS 4 |
| 967 | 35471166 |  | 23/06/2004 | Dace | 15.5 | ESMS 4 |
| 968 | 35471192 |  | 23/06/2004 | Dace | 14.4 | ESMS 4 |
| 969 | 94698687 |  | 23/06/2004 | Dace | 34.3 | Holme Bridge 2 |
| 970 | 35471164 |  | 23/06/2004 | Dace | 16.2 | ESMS 4 |
| 971 | 35471137 |  | 23/06/2004 | Roach | 14.6 | ESMS 2 |
| 972 | 35471161 |  | 23/06/2004 | Roach | 15.8 | ESMS 2 |
| 973 | 35471145 |  | 23/06/2004 | Roach | 14.1 | ESMS 400 m US |
| 974 | 113943677 |  | 23/06/2004 | Dace | 16 | ESMS 400 m US |
| 975 | 122451103 |  | 23/06/2004 | Pike | 47.5 | ESMS 400 m US |
| 976 | 122451138 |  | 23/06/2004 | Dace | 15.3 | ESMS 400 m US |


| 977 | 122451142 | 23/06/2004 | Perch |
| :---: | :---: | :---: | :---: |
| 978 | 122451182 | 23/06/2004 | Roach |
| 979 | 122451095 | 23/06/2004 | Roach |
| 980 | 122451162 | 23/06/2004 | Dace |
| 981 | 122451128 | 23/06/2004 | Dace |
| 982 | 122451113 | 23/06/2004 | Roach |
| 983 | 122451093 | 23/06/2004 | Dace |
| 984 | 122451105 | 23/06/2004 | Dace |
| 985 | 122451102 | 23/06/2004 | Roach |
| 986 | 35471140 | 23/06/2004 | Dace |
| 987 | 122451116 | 23/06/2004 | Dace |
| 988 | 122451130 | 23/06/2004 | Roach |
| 989 | 122451155 | 23/06/2004 | Roach |
| 990 | 122451159 | 23/06/2004 | Roach |
| 991 | 122451108 | 23/06/2004 | Roach |
| 992 | 122451158 | 23/06/2004 | Dace |
| 993 | 122451134 | 23/06/2004 | Dace |
| 994 | 122451173 | 23/06/2004 | Dace |
| 995 | 122451156 | 23/06/2004 | Dace |
| 996 | 122451167 | 23/06/2004 | Dace |
| 997 | 122451148 | 23/06/2004 | Roach |
| 998 | 122451115 | 23/06/2004 | Dace |
| 999 | 122451178 | 23/06/2004 | Dace |
| 1000 | 122451123 | 23/06/2004 | Dace |
| 1001 | 35471135 | 23/06/2004 | Dace |
| 1002 | 122451160 | 23/06/2004 | Dace |
| 1003 | 122450377 | 23/06/2004 | Dace |
| 1004 | 122451097 | 23/06/2004 | Dace |
| 1005 | 122451057 | 24/06/2004 | Dace |
| 1006 | 122451179 | 24/06/2004 | Roach |
| 1007 | 122451053 | 24/06/2004 | Dace |
| 1008 | 122451145 | 24/06/2004 | Roach |
| 1009 | 122451037 | 24/06/2004 | Roach |
| 1010 | 35471126 | 24/06/2004 | Dace |
| 1011 | 122451125 | 24/06/2004 | Roach |
| 1012 | 122451099 | 24/06/2004 | Dace |
| 1013 | 122451084 | 24/06/2004 | Dace |
| 1014 | 122451066 | 24/06/2004 | Dace |
| 1015 | 122451071 | 24/06/2004 | Dace |
| 1016 | 122451072 | 24/06/2004 | Dace |
| 1017 | 122451136 | 24/06/2004 | Dace |
| 1018 | 122451129 | 24/06/2004 | Dace |
| 1019 | 122451032 | 24/06/2004 | Dace |
| 1020 | 122451074 | 24/06/2004 | Dace |
| 1021 | 122451119 | 24/06/2004 |  |
| 1022 | 122451069 | 24/06/2004 | Roach |
| 1023 | 12245069 |  | Dace |
| 1024 | 122451046 | 24/06/2004 | Dace |
| 1025 | 35471153 | 24/06/2004 | Roach |

22.4 ESMS 750 m US
17.4 ESMS 800 m US
16.1 ESMS 800m US
15.3 ESMS 800m US
15.1 ESMS 800 m US
14.8 ESMS 800 m US

17 ESMS 800m US
16.9 ESMS 800 m US
15.3 ESMS 800 m US
15.6 ESMS 800m US

16 ESMS
15.1 ESMS 800m US
15.2 ESMS 800m US
16.9 ESMS 800m US
14.7 ESMS 800 m US
16.6 ESMS 800 m US
14.9 ESMS 800 m US

18 ESMS 860m US
16.4 ESMS 860 m US
15.3 ESMS 860m US
14.6 ESMS 860m US
14.7 ESMS 860m US
17.2 ESMS 860m US
15.1 ESMS 860m US
17.1 ESMS 860m US
17.2 ESMS 860m US
16.5 ESMS 860 m US
15.5 ESMS 860m US
16.9 ESMS 3
14.4 ESMS 3
15.5 ESMS 3

17 ESMS 3
17.5 ESMS 3
22.1 ESMS 3
20.8 ESMS
18.1 ESMS 3
15.5 ESMS 3
18.5 ESMS 3
19.8 ESMS 3
17.5 ESMS 3
14.9 ESMS 3 ESMS above smolt
20.5 counter ESMS above smolt
15.8 counter ESMS above smolt
16.4 counter ESMS above smolt
16.2 counter ESMS above smolt
13.9 counter ESMS above smolt
15.8 counter ESMS above smolt
15.2 counter
17.2 ESMS above smolt

| 1026 | 122451169 | DC0029-FD0A | 24/06/2004 | Dace | 16 | ESMS above smolt counter ESMS above smolt |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  |  |  |  |
| 1027 | 122450379 |  | 24/06/2004 | Dace | 17 |  |
| 1028 |  |  | 03/06/2004 | Dace | 14.6 | Redcliff |
| 1029 | 122451141 |  | 24/06/2004 | Dace | 16.2 | ESMS above smolt |
|  |  |  |  |  |  | ESMS above smolt |
| 1030 | 122450383 |  | 24/06/2004 | Dace | 15.2 | counter |
| 1031 | 122451014 |  | 24/06/2004 | Dace | 14.7 | ESMS above smolt |
|  |  |  |  |  |  | ESMS above smolt |
| 1032 | 122451012 |  | 24/06/2004 | Dace | 15.7 | counter |
| 1033 | 35471141 |  | 24/06/2004 | Dace | 16 | ESMS above smolt counter |
|  | 122451183 |  | 24/06/2004 | Dace |  | ESMS above smolt |
| 1034 |  |  | 24/06/2004 | Dace | 15.3 | ESMS above smolt |
| 1035 | 122451004 |  | 24/06/2004 | Dace | 15.3 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1036 | 122451083 |  | 24/06/2004 | Dace | 16.2 | counter <br> ESMS above smolt |
| 1037 | 122451100 |  | 24/06/2004 | Dace | 16.04 | counter |
|  |  |  | 24/06/2004 | Roach | 25.1 | ESMS above smolt counter |
| 1038 | 122451061 |  |  |  |  | ESMS above smolt |
| 1039 | 122451039 |  | 24/06/2004 | Dace | 15.2 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1040 | 122451040 |  | 24/06/2004 | Dace | 17.8 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1041 | 35471146 |  | 24/06/2004 | Dace | 17.9 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1042 | 122451073 |  | 24/06/2004 | Dace | 23.4 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1043 | 122451033 |  | 24/06/2004 | Dace | 18.1 | counter |
| 1044 | 122451058 |  | 24/06/2004 | Dace | 16.1 |  |
|  |  |  |  |  |  | ESMS above smolt |
| 1045 | 122451059 |  | 24/06/2004 | Dace | 17.4 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1047 | 122451031 |  | 24/06/2004 | Roach | 16.9 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1048 | 122451052 |  | 24/06/2004 | Dace | 15.4 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1049 | 35471130 |  | 24/06/2004 | Roach | 18.7 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1050 | 122451175 |  | 24/06/2004 | Dace | 24 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1051 | 122451082 |  | 24/06/2004 | Dace | 14.2 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1052 | 122451042 |  | 24/06/2004 | Dace | 13.9 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1053 | 122451092 |  | 24/06/2004 | Dace | 14.5 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1054 | 122451006 |  | 24/06/2004 | Perch | 23.5 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1055 | 122451001 |  | 24/06/2004 | Dace | 17.6 | counter |
|  |  |  |  |  |  | ESMS above smolt |
| 1056 | 122451177 |  | 24/06/2004 | Roach | 19.4 | counter |
| 1057 | 122451151 |  | 24/06/2004 | Dace | 22.2 | ESMS above smolt |


| 1058 | 122451016 |  | 24/06/2004 | Dace |
| :---: | :---: | :---: | :---: | :---: |
| 1059 | 35471134 |  | 24/06/2004 | Dace |
| 1060 | 122451137 |  | 24/06/2004 | Dace |
| 1061 | 122451122 |  | 24/06/2004 | Dace |
| 1062 | 122451000 |  | 24/06/2004 | Dace |
| 1063 | 122451045 |  | 24/06/2004 | Dace |
| 1069 | 122450384 | DC0024-D72B | 17/09/2004 | Dace |
| 1070 | 122451163 | DC002B-17A6 | 17/09/2004 | Dace |
| 1071 | 122451114 | DC0024-E5C0 | 17/09/2004 | Dace |
| 1072 | 122451077 | DC002B-1376 | 17/09/2004 | Dace |
| 1073 | 122451044 | DC002A-FBAA | 17/09/2004 | Dace |
| 1074 | 122451051 | DC0024-7C89 | 17/09/2004 | Dace |
| 1075 | 122451088 | DC0024-91E4 | 17/09/2004 | Dace |
| 1076 | 122451081 | DC0024-D924 | 17/09/2004 | Dace |
| 1077 | 122451121 | DC0024-DCC2 | 17/09/2004 | Dace |
| 1078 | 122450731 |  | 17/02/2005 | Roach |
| 1079 |  | DC002B-10D8 | 17/02/2005 | Roach |
| 1080 |  | DC0029-F07F | 17/02/2005 | Roach |
| 1081 |  | DC0029-EAD7 | 17/02/2005 | Dace |
| 1082 |  | DC0024-F2CB | 17/02/2005 | Roach |
| 1083 |  | DC002B-1A9E | 17/02/2005 | Dace |
| 1084 |  | DC0024-8F42 | 17/02/2005 | Roach |
| 1085 |  | DC0024-EECD | 17/02/2005 | Dace |
| 1086 |  | DC0024-D19A | 17/02/2005 | Roach |
| 1087 |  | DC002B-0992 | 17/02/2005 | Roach |
| 1088 |  | DC0024-757F | 17/02/2005 | Dace |
| 1089 |  | DC002B-16DF | 17/02/2005 | Roach |
| 1090 |  | DC0024-CBD0 | 17/02/2005 | Roach |
| 1091 |  | DC002B-1C3D | 17/02/2005 | Roach |
| 1092 |  | DC0024-E3A4 | 17/02/2005 | Dace |
| 1093 |  | DC002B-18AF | 17/02/2005 | Roach |
| 1094 |  | DC002B-13BE | 17/02/2005 | Roach |
| 1095 |  | DC002A-F7BF | 17/02/2005 | Dace |
| 1096 |  | DC0024-7419 | 17/02/2005 | Dace |
| 1097 |  | DC002B-07CE | 17/02/2005 | Roach |
| 1098 |  | DC0024-DF8C | 17/02/2005 | Dace |
| 1099 |  | DC002A-FB9A | 17/02/2005 | Roach |
| 1100 |  | DC001B-E6D1 | 18/03/2003 | Pike |
| 1101 | 94698717 |  | 18/03/2003 | Pike |
| 1102 | 94698714 |  | 18/03/2003 | Pike |
| 1246 | 113943700 |  | 03/06/2004 | Dace |
| 1247 |  | DC0029-EF4D | 03/06/2004 | Dace |
| 1248 | 35471211 |  | 03/06/2004 | Dace |
| 1249 | 35471223 |  | 03/06/2004 | Dace |
| 1250 |  | DC002A-139F | 03/06/2004 | Dace |
| 2000 |  | DC002B-1140 | 17/02/2005 | Dace |
| 2001 |  | DC0024-E91D | 17/02/2005 | Roach |
| 2002 |  | DC0024-F0EC | 17/02/2005 | Roach |

counter
ESMS above smolt
14.7 counter

ESMS above smolt
16.4 counter

ESMS above smolt
15.1 counter

ESMS above smolt
12.9 counter

ESMS above smolt
13.4 counter

ESMS above smolt
18.2 counter

19 ESMS 2
23 ESMS 2
18.9 ESMS 2
21.8 ESMS 2
20.4 ESMS 2

19 ESMS 2
20.1 ESMS 2
19.3 ESMS 2

19 ESMS 2
19.1 Wareham Quay

21 Wareham Quay
26.7 Wareham Quay
24.2 Wareham Quay
22.4 Wareham Quay
24.6 Wareham Quay
22.5 Wareham Quay
19.8 Wareham Quay
20.1 Wareham Quay
24.4 Wareham Quay
25.5 Wareham Quay
29.9 Wareham Quay
23.1 Wareham Quay

21 Wareham Quay
25.4 Wareham Quay
20.1 Wareham Quay
24.2 Wareham Quay
25.4 Wareham Quay
19.6 Wareham Quay

36 Wareham Quay
26.4 Wareham Quay
22.2 Wareham Quay
12.7 Rushton
17.9 Rushton
44.6 Rushton
19.1 Redcliff
14.2 Redcliff
19.1 Redcliff
16.2 Redcliff
21.7 Redcliff
21.2 Wareham Quay

26 Wareham Quay
22.1 Wareham Quay

| 2003 |  | DC0024-DC07 | 17/02/2005 | Roach | 20.1 | Wareham Quay |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2004 |  | DC002B-103C | 17/02/2005 | Roach | 22.1 | Wareham Quay |
| 2005 |  | DC0024-E608 | 17/02/2005 | Dace | 22 | Wareham Quay |
| 2006 |  | DC0024-9945 | 17/02/2005 | Roach | 38.5 | Wareham Quay |
| 2007 |  | DC0024-89B3 | 17/02/2005 | Roach | 18.3 | Wareham Quay |
| 2008 |  | DC002B-0E17 | 17/02/2005 | Dace | 25.2 | Wareham Quay |
| 2009 |  | DC002A-F9F1 | 17/02/2005 | Roach | 21.6 | Wareham Quay |
| 2010 |  | DC0024-DA1B | 17/02/2005 | Roach | 21.1 | Wareham Quay |
| 2011 | 35471199 |  | 03/06/2004 | Dace | 21.1 | Redcliff |
| 2012 |  | DC0024-D718 | 17/02/2005 | Roach | 18.7 | Wareham Quay |
| 2013 |  | DC002A-FF8D | 17/02/2005 | Dace | 18 | Wareham Quay |
| 2014 |  | DC0024-74AF | 17/02/2005 | Roach | 22.2 | Wareham Quay |
| 2015 |  | DC002B-0271 | 17/02/2005 | Dace | 22.5 | Wareham Quay |
| 2016 |  | DC0024-E0B5 | 17/02/2005 | Roach | 18.2 | Wareham Quay |
| 2017 |  | DC002A-F4D0 | 17/02/2005 | Roach | 20.1 | Wareham Quay |
| 2018 |  | DC002B-06BB | 17/02/2005 | Roach | 20.5 | Wareham Quay |
| 2019 |  | DC0024-8983 | 17/02/2005 | Hybrid | 21.8 | Wareham Quay |
| 2020 |  | DC0024-82B2 | 17/02/2005 | Bream | 33.1 | Wareham Quay |
| 2021 |  | DC0024-98B9 | 17/02/2005 | Dace | 20.2 | Wareham Quay |
| 2022 |  | DC0024-D6AF | 17/02/2005 | Roach | 18.3 | Wareham Quay |
| 2023 |  | DC002B-004B | 17/02/2005 | Dace | 20.5 | Wareham Quay |
| 2024 |  | DC0024-DB9A | 17/02/2005 | Dace | 20.2 | Wareham Quay |
| 2025 |  | DC0024-E9A8 | 17/02/2005 | Roach | 19.1 | Wareham Quay |
| 2026 |  | DC0024-82C4 | 17/02/2005 | Dace | 18.5 | Wareham Quay |
| 2027 |  | DC002B-1BDB | 17/02/2005 | Dace | 22.8 | Wareham Quay |
| 2028 |  | DC002A-F65C | 17/02/2005 | Roach | 22.4 | Wareham Quay |
| 2029 |  | DC002B-0058 | 17/02/2005 | Roach | 25.6 | Wareham Quay |
| 2030 |  | DC002B-0ACE | 17/02/2005 | Roach | 18.9 | Wareham Quay |
| 2031 | 122451164 |  | 17/09/2004 | Dace | 19 | ESMS 3 |
| 2032 |  | DC0024-8225 | 17/02/2005 | Dace | 16.3 | Wareham Quay |
| 2033 |  | DC0024-7BEE | 17/02/2005 | Roach | 22.9 | Wareham Quay |
| 2034 |  | DC002B-15E1 | 17/02/2005 | Dace | 18.3 | Wareham Quay |
| 2035 | 35471245 |  | 03/06/2004 | Dace | 0 | Redcliff |
| 2036 |  | DC002A-FB16 | 17/02/2005 | Dace | 15.2 | Wareham Quay |
| 2037 |  | DC0024-DFDF | 17/02/2005 | Dace | 16.7 | Wareham Quay |
| 2038 |  | DC002B-1664 | 17/02/2005 | Dace | 15.5 | Wareham Quay |
| 2039 | 122450738 |  | 17/02/2005 | Roach | 32 | Wareham Quay |
| 2040 | 127178942 |  | 17/02/2005 | Roach | 22 | Wareham Quay |
| 2041 | 127178979 |  | 17/02/2005 | Roach | 23.5 | Wareham Quay |
| 2042 | 127178941 |  | 17/02/2005 | Roach | 23.4 | Wareham Quay |
| 2043 | 127178940 |  | 17/02/2005 | Roach | 14.1 | Wareham Quay |
| 2044 | 127178939 |  | 17/02/2005 | Dace | 26.7 | Wareham Quay |
| 2045 | 127178938 |  | 17/02/2005 | Dace | 20.1 | Wareham Quay |
| 2046 | 127178937 |  | 17/02/2005 | Roach | 18.8 | Wareham Quay |
| 2047 | 127178936 |  | 17/02/2005 | Dace | 23.5 | Wareham Quay |
| 2048 | 127178935 |  | 17/02/2005 | Dace | 19.5 | Wareham Quay |
| 2049 | 127178934 |  | 17/02/2005 | Dace | 24.1 | Wareham Quay |
| 2050 | 127178933 |  | 17/02/2005 | Roach | 32.6 | Wareham Quay |
| 2051 | 127178932 |  | 17/02/2005 | Dace | 14.6 | Wareham Quay |
| 2052 | 127178981 |  | 17/02/2005 | Roach | 24.7 | Wareham Quay |
| 2053 | 127178980 |  | 17/02/2005 | Dace | 24.1 | Wareham Quay |
| 2054 | 122450702 |  | 17/02/2005 | Roach | 21.4 | Wareham Quay |
| 2055 | 122450703 |  | 17/02/2005 | Dace | 27.1 | Wareham Quay |
| 2056 | 122450704 |  | 17/02/2005 | Dace | 26.8 | Wareham Quay |


| 2057 | 122450705 |  | 17/02/2005 | Dace |
| :---: | :---: | :---: | :---: | :---: |
| 2058 | 122450706 |  | 17/02/2005 | Roach |
| 2059 | 122450707 |  | 17/02/2005 | Roach |
| 2060 | 122450710 |  | 17/02/2005 | Dace |
| 2061 | 122450708 |  | 17/02/2005 | Roach |
| 2062 | 122450709 |  | 17/02/2005 | Dace |
| 2063 | 122450711 |  | 17/02/2005 | Roach |
| 2064 | 122450712 |  | 17/02/2005 | Dace |
| 2065 | 122450713 |  | 17/02/2005 | Roach |
| 2066 | 122450714 |  | 17/02/2005 | Dace |
| 2067 | 122450715 |  | 17/02/2005 | Roach |
| 2068 | 122450716 |  | 17/02/2005 | Roach |
| 2069 | 122450717 |  | 17/02/2005 | Bream |
| 2070 | 122450718 |  | 17/02/2005 | Roach |
| 2071 | 122450719 |  | 17/02/2005 | Dace |
| 2072 | 122450720 |  | 17/02/2005 | Roach |
| 2073 | 122450721 |  | 17/02/2005 | Dace |
| 2074 | 122450722 |  | 17/02/2005 | Roach |
| 2075 | 122450723 |  | 17/02/2005 | Dace |
| 2076 | 122450724 |  | 17/02/2005 | Roach |
| 2077 | 122450725 |  | 17/02/2005 | Roach |
| 2078 | 122450726 |  | 17/02/2005 | Roach |
| 2079 | 122450727 |  | 17/02/2005 | Roach |
| 2080 | 122450728 |  | 17/02/2005 | Roach |
| 2081 | 122450729 |  | 17/02/2005 | Roach |
| 2082 | 122450730 |  | 17/02/2005 | Roach |
| 2083 | 122450732 |  | 17/02/2005 | Roach |
| 2084 | 122450733 |  | 17/02/2005 | Roach |
| 2085 | 122450734 |  | 17/02/2005 | Roach |
| 2086 | 122450735 |  | 17/02/2005 | Dace |
| 2087 | 122450736 |  | 17/02/2005 | Dace |
| 2088 | 122450737 |  | 17/02/2005 | Dace |
| 2089 |  | DC0024-7C26 | 16/02/2005 | Dace |
| 2090 |  | DC0024-90A9 | 16/02/2005 | Dace |
| 2091 |  | DC002B-16F7 | 16/02/2005 | Dace |
| 2092 |  | DC002B-106D | 16/02/2005 | Dace |
| 2093 |  | DC0024-92F9 | 16/02/2005 | Dace |
| 2094 |  | DC0024-EB93 | 16/02/2005 | Dace |
| 2095 | 113945673 | DC001B-FA0C | 17/03/2003 | Pike |
| 2096 | 122450769 | DC001B-FF91 | 17/03/2003 | Pike |
| 2097 | 94698734 | DC001B-E824 | 17/03/2003 | Pike |
| 2098 |  | DC001B-FFF1 | 17/03/2003 | Pike |
| 2099 | 94698733 | DC001B-E7AF | 17/03/2003 | Pike |
| 2100 |  | DC001C-0907 | 17/03/2003 | Pike |
| 2101 | 94698735 | DC001B-F464 | 17/03/2003 | Pike |
| 2102 |  | DC001B-EFF7 | 17/03/2003 | Pike |
| 2103 | 94698732 | DC001B-FAC9 | 17/03/2003 | Pike |
| 2104 | 94698731 | DC001B-EA63 | 17/03/2003 | Pike |
| 2105 | 94698730 | DC001B-EDF1 | 17/03/2003 | Pike |
| 2106 | 94698729 | DC001B-F62E | 17/03/2003 | Pike |
| 2107 | 96498727 | DC001C-01F1 | 17/03/2003 | Pike |
| 2108 | 94698726 | DC001C-018C | 17/03/2003 | Pike |
| 2109 | 94698725 | DC001B-F5C4 | 17/03/2003 | Pike |
| 2110 | 94698724 | DC001C-07E6 | 17/03/2003 | Pike |

25.9 Wareham Quay
18.2 Wareham Quay

33 Wareham Quay
23.9 Wareham Quay
19.9 Wareham Quay
25.4 Wareham Quay
19.2 Wareham Quay
22.2 Wareham Quay
21.9 Wareham Quay
17.7 Wareham Quay
23.6 Wareham Quay

20 Wareham Quay
32.2 Wareham Quay
20.9 Wareham Quay
24.3 Wareham Quay
32.7 Wareham Quay
15.5 Wareham Quay
34.4 Wareham Quay
18.7 Wareham Quay
34.2 Wareham Quay

18 Wareham Quay
16.5 Wareham Quay
18.1 Wareham Quay
21.1 Wareham Quay

17 Wareham Quay
18 Wareham Quay
23.9 Wareham Quay
22.5 Wareham Quay
22.1 Wareham Quay
17.8 Wareham Quay
18.5 Wareham Quay

18 Wareham Quay
14.4 Redcliff
14.4 Redcliff
20.2 Redcliff
20.7 Redcliff
17.9 Redcliff
15.8 Redcliff

11 Railway 1
12.3 Railway 1

38 Railway 1
10.3 Railway 1
31.1 Railway
13.2 Railway 2

25 Railway 2
11.5 Railway 2

32 Railway 2
17.9 Railway 2
19.4 Railway 2
24.4 Railway 2
29.1 Railway 3
26.2 Railway 3
31.2 Railway 4
41.5 Railway 4

| 2111 | 94698723 | DC001C-025A | 17/03/2003 | Pike | 22 | Railway 4 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2112 | 94698722 | DC001B-F412 | 17/03/2003 | Pike | 45.9 | Railway 4 |
| 2113 | 122450754 |  | 18/04/2005 | Pike | 42.7 | Rushton 1 |
| 2114 |  | DC001C-02C5 | 17/03/2003 | Roach | 8.1 | Flood Relief 1 |
| 2115 |  | DC001B-FDC0 | 17/03/2003 | Pike | 14.4 | Flood Relief 2 |
| 2116 |  | DC001B-E400 | 17/03/2003 | Roach | 10.4 | Flood Relief |
| 2117 | 113945627 |  | 25/01/2004 | Grayling | 26 | ESMS |
| 2118 | 113945624 |  | 25/01/2004 | Grayling | 25.8 | ESMS |
| 2119 | 113945626 |  | 25/01/2004 | Grayling | 28.8 | ESMS |
| 2120 | 113945622 |  | 25/01/2004 | Grayling | 24.8 | ESMS |
| 2121 | 113945621 |  | 25/01/2004 | Grayling | 26.5 | ESMS |
| 2122 | 113945620 |  | 25/01/2004 | Grayling | 18 | ESMS |
| 2123 | 113945619 |  | 25/01/2004 | Grayling | 14.8 | ESMS |
| 2124 |  | DC0096-D7AB | 25/07/2000 | Pike | 27.4 | Rushton Main River Frome Holme |
| 2125 |  | DC001B-F929 | 20/03/2003 | Dace | 7.7 | Bridge |
| 2126 |  | DC001C-0C8D | 20/03/2003 | Roach | 11.5 | Holme Bridge 3 |
| 2127 | 94698686 |  | 20/03/2003 | Pike | 70.2 | Holme Bridge 4 |
| 2128 | 122451026 |  | 25/06/2004 | Eel | 35.8 | Holme Bridge |
| 2129 | 113949675 |  | 25/06/2004 | Pike | 27.1 | Holme Bridge 200-250m |
| 2130 | 122451028 |  | 25/06/2004 | Pike | 34 | Holme Bridge 400-450m |
| 2131 | 122451101 |  | 25/06/2004 | Pike | 27.5 | Holme Bridge 400-450m Main River Frome |
| 2132 | 94698338 | DC001C-A03A | 16/04/2003 | Pike | 71.5 | Weirpool Main River Frome |
| 2133 | 94698337 | DC001C-97CB | 16/04/2003 | Pike | 70.2 | Weirpool |
| 2134 | 94698336 | DC001C-68A0 | 16/04/2003 | Pike | 22.6 | Main River Frome T6-T9 <br> Main River Frome |
| 2135 | 94698335 | DC001C-933D | 16/04/2003 | Pike | 21.4 | Weirpool <br> Main River Frome |
| 2136 | 94698333 |  | 16/04/2003 | Dace | 18.4 | Weirpool |
| 2137 | 35471198 |  | 23/06/2004 | Dace | 16.8 | ESMS |
| 2139 | 35474435 |  | 23/06/2004 | Dace | 17.1 | ESMS |
| 2140 | 122491162 |  | 23/06/2004 | Dace | 15.3 | ESMS |
| 2141 | 113949636 |  | 23/06/2004 | Roach | 17.7 | ESMS |
| 2143 | 35471133 |  | 23/06/2004 | Eel | 37 | ESMS 4 |
| 2144 |  | DC002B-1769 | 23/06/2004 | Salmon | 11.5 | ESMS 300m US |
| 2145 |  | DC002A-F8AA | 21/06/2004 | Pike | 6.5 | Flood relief 1 |
| 2151 |  | DC002A-0FDC | 03/06/2004 | Dace | 20.4 | Redcliff |
| 2152 | 35471180 |  | 03/06/2004 | Dace | 14.3 | Redcliff |
| 2153 |  | DC0029-FB2F | 03/06/2004 | Dace | 22.9 | Redcliff |
| 2154 | 35471231 |  | 03/06/2004 | Dace | 15 | Redcliff |
| 2155 |  | DC002A-052D | 03/06/2004 | Dace | 15.5 | Redcliff |
| 2156 | 35471221 |  | 03/06/2004 | Dace | 24.6 | Redcliff |
| 2157 |  | DC0029-FE7F | 03/06/2004 | Dace | 22.3 | Redcliff |
| 2158 | 35471181 |  | 03/06/2004 | Dace | 18 | Redcliff |
| 2159 |  | DC002A-00F3 | 03/06/2004 | Dace | 23.8 | Redcliff |
| 2160 | 35471193 |  | 03/06/2004 | Dace | 17.4 | Redcliff |
| 2161 |  | DC002A-0FE8 | 03/06/2004 | Dace | 22.6 | Redcliff |
| 2162 | 35471226 |  | 03/06/2004 | Dace | 15.4 | Redcliff |
| 2163 |  | DC002A-0339 | 03/06/2004 | Dace | 15.2 | Redcliff |
| 2164 |  | DC002A-0DC5 | 03/06/2004 | Dace | 21.7 | Redcliff |
| 2165 |  | DC0029-FAB6 | 03/06/2004 | Dace | 15.6 | Redcliff |
| 2166 |  | DC0029-F00F | 03/06/2004 | Dace | 22.2 | Redcliff |
| 2167 |  | DC0029-EE11 | 03/06/2004 | Dace | 20.8 | Redcliff |


| 2168 |  | DC002A-14B4 | 03/06/2004 | Dace | 13 | Redcliff |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2169 |  | DC0024-F2D9 | 03/06/2004 | Dace | 19.9 | Redcliff |
| 2170 |  | DC002A-0622 | 03/06/2004 | Dace | 16.7 | Redcliff |
| 2171 |  | DC002A-0246 | 03/06/2004 | Dace | 16.5 | Redcliff |
| 2172 |  | DC0029-F4A2 | 03/06/2004 | Dace | 16 | Redcliff |
| 2173 |  | DC0029-F3CF | 03/06/2004 | Dace | 15.1 | Redcliff |
| 2174 |  | DC0029-EF10 | 03/06/2004 | Dace | 13.1 | Redcliff |
| 2175 |  | DC0029-F443 | 03/06/2004 | Dace | 21.9 | Redcliff |
| 2176 |  | DC0029-FF96 | 03/06/2004 | Dace | 0 | Redcliff |
| 2177 |  | DC0029-F848 | 03/06/2004 | Dace | 23.2 | Redcliff |
| 2178 |  | DC0029-FDF1 | 03/06/2004 | Dace | 24.2 | Redcliff |
| 2179 |  | DC002A-0352 | 03/06/2004 | Dace | 21.2 | Redcliff |
| 2180 |  | DC0029-FBA5 | 03/06/2004 | Dace | 22.2 | Redcliff |
| 2181 |  | DC002A-14FF | 03/06/2004 | Dace | 20.8 | Redcliff |
| 2182 |  | DC0029-F480 | 03/06/2004 | Dace | 16.9 | Redcliff |
| 2183 |  | DC002A-0180 | 03/06/2004 | Dace | 15.8 | Redcliff |
| 2184 |  | DC0029-ECAC | 01/03/1964 | Dace | 14.9 | Redcliff |
| 2185 |  | DC0029-F7BA | 03/06/2004 | Dace | 23.4 | Redcliff |
| 2186 |  | DC0029-EF66 | 03/06/2004 | Dace | 16.1 | Redcliff |
| 2187 |  | DC002A-0130 | 03/06/2004 | Dace | 18.3 | Redcliff |
| 2188 |  | DC002A-1472 | 03/06/2004 | Dace | 14.6 | Redcliff |
| 2189 |  | DC0029-F17A | 03/06/2004 | Dace | 14.3 | Redcliff |
| 2190 |  | DC002A-016D | 03/06/2004 | Dace | 23.3 | Redcliff |
| 2191 |  | DC002A-1427 | 03/06/2004 | Dace | 14.2 | Redcliff |
| 2192 |  | DC002A-0030 | 03/06/2004 | Dace | 15 | Redcliff |
| 2193 |  | DC0029-EB60 | 03/06/2004 | Dace | 13.7 | Redcliff |
| 2194 |  | DC002A-1141 | 03/06/2004 | Dace | 14.2 | Redcliff |
| 2195 | 35471186 |  | 03/06/2004 | Dace | 18.4 | Redcliff |
| 2196 | 35471235 |  | 03/06/2004 | Dace | 24.9 | Redcliff |
| 2197 | 35471210 |  | 03/06/2004 | Dace | 18.7 | Redcliff |
| 2198 | 35471249 |  | 03/06/2004 | Dace | 21.1 | Redcliff |
| 2199 | 35471196 |  | 03/06/2004 | Dace | 19.6 | Redcliff |
| 2200 | 35471240 |  | 03/06/2004 | Dace | 15.7 | Redcliff |
| 2201 | 35471188 |  | 03/06/2004 | Dace | 20.6 | Redcliff |
| 2202 | 35471217 |  | 03/06/2004 | Dace | 22.7 | Redcliff |
| 2203 | 35471218 |  | 03/06/2004 | Dace | 22.2 | Redcliff |
| 2204 | 35471246 |  | 03/06/2004 | Dace | 22.2 | Redcliff |
| 2205 | 35471238 |  | 03/06/2004 | Dace | 21.9 | Redcliff |
| 2206 | 35471213 |  | 03/06/2004 | Dace | 20.3 | Redcliff |
| 2207 | 35471236 |  | 03/06/2004 | Dace | 21.5 | Redcliff |
| 2208 | 35471197 |  | 03/06/2004 | Dace | 22.8 | Redcliff |
| 2209 | 35471190 |  | 03/06/2004 | Dace | 19.8 | Redcliff |
| 2210 | 35471178 |  | 03/06/2004 | Dace | 18.4 | Redcliff |
| 2211 | 35471242 |  | 03/06/2004 | Dace | 16 | Redcliff |
| 2212 | 35471208 |  | 03/06/2004 | Dace | 21.3 | Redcliff |
| 2213 | 35471212 |  | 03/06/2004 | Dace | 17.8 | Redcliff |
| 2214 | 35471229 |  | 03/06/2004 | Dace | 16 | Redcliff |
| 2215 | 35471247 |  | 03/06/2004 | Dace | 15.2 | Redcliff |
| 2216 | 35471216 |  | 03/06/2004 | Dace | 15.5 | Redcliff |
| 2217 | 35471222 |  | 03/06/2004 | Dace | 20.6 | Redcliff |
| 2218 | 35471248 |  | 03/06/2004 | Dace | 24.6 | Redcliff |
| 2219 | 35471201 |  | 03/06/2004 | Dace | 18.4 | Redcliff |
| 2220 | 35471207 |  | 03/06/2004 | Dace | 15 | Redcliff |
| 2221 | 35471205 |  | 03/06/2004 | Dace | 21.6 | Redcliff |


| 2222 | 35471243 |  | 03/06/2004 | Dace |
| :---: | :---: | :---: | :---: | :---: |
| 2223 | 35471234 |  | 03/06/2004 | Dace |
| 2224 | 35471241 |  | 03/06/2004 | Dace |
| 2225 | 35471209 |  | 03/06/2004 | Dace |
| 2226 | 35471179 |  | 03/06/2004 | Dace |
| 2227 | 35471215 |  | 03/06/2004 | Dace |
| 2228 | 113943696 |  | 03/06/2004 | Dace |
| 2229 |  | DC0029-FDBD | 03/06/2004 | Dace |
| 2230 |  | DC0029-4602 | 03/06/2004 | Dace |
| 2231 |  | DC002A-162F | 03/06/2004 | Dace |
| 2232 |  | DC0029-EF5D | 03/06/2004 | Dace |
| 2233 | 113943667 | DC0029-F25E | 03/06/2004 | Dace |
| 2234 | 113943694 | DC0029-EC2B | 03/06/2004 | Dace |
| 2235 | 113943704 | DC0029-FA4E | 03/06/2004 | Dace |
| 2236 | 113943695 | DC0029-4050 | 03/06/2004 | Dace |
| 2237 |  | DC0029-EC7A | 03/06/2004 | Dace |
| 2238 | 113943984 |  | 03/06/2004 | Dace |
| 2239 | 113943673 |  | 03/06/2004 | Dace |
| 2240 |  | DC002A-00CC | 03/06/2004 | Dace |
| 2241 | 113943690 |  | 03/06/2004 | Dace |
| 2242 |  | DC0029-F5AC | 03/06/2004 | Dace |
| 2243 | 113943693 |  | 04/06/2003 | Dace |
| 2244 | 113943691 |  | 03/06/2004 | Dace |
| 2245 | 113943703 |  | 03/06/2004 | Dace |
| 2246 | 113943698 |  | 03/06/2004 | Dace |
| 2247 | 113943681 |  | 03/06/2004 | Dace |
| 2248 | 113943682 |  | 03/06/2004 | Dace |
| 2249 | 113943701 |  | 03/06/2004 | Dace |
| 2250 | 113943702 |  | 03/06/2004 | Dace |
| 2251 | 113943676 |  | 03/06/2004 | Dace |
| 2252 | 113943672 |  | 03/06/2004 | Dace |
| 2253 | 113943686 |  | 03/06/2004 | Dace |
| 2254 | 113943675 |  | 03/06/2004 | Dace |
| 2255 | 113943678 |  | 03/06/2004 | Dace |
| 2256 | 113943689 |  | 03/06/2004 | Dace |
| 2257 | 113943674 |  | 04/06/2003 | Dace |

25.4 Redcliff
23.9 Redcliff
20.7 Redcliff
15.1 Redcliff
16.8 Redcliff

0 Redcliff
20.9 Redcliff
16.5 Redcliff
20.2 Redcliff
19.9 Redcliff
13.3 Redcliff Redcliff to Wareham
14.8 Quay

Redcliff to Wareham
21.4 Quay Redcliff to Wareham
19.2 Quay

Redcliff to Wareham
18.6 quay

Redcliff to Wareham
19 Quay Redcliff to Wareham
15.6 Quay

Redcliff to Wareham
17 Quay Redcliff to Wareham
21.3 Quay

Redcliff to Wareham
25 Quay Redcliff to Wareham
18.4 Quay

24 Wareham Quay Redcliff to Wareham
20.1 quay

Redcliff to Wareham
15.2 quay Redcliff to Wareham
21.2 quay Redcliff to Wareham
22.3 quay

Redcliff to Wareham
16.5 quay Wareham quay to
15.8 Redcliff Wareham quay to
21.8 Redcliff Wareham quay to
19.7 Redcliff Wareham quay to
23.3 Redcliff Wareham quay to
20.4 Redcliff Wareham quay to
16.2 Redcliff Wareham quay to
21.8 Redcliff Wareham quay to
20.5 Redcliff

17 Wareham Quay

| 2258 | 113943687 |  | 03/06/2004 | Dace | 15.5 | Wareham quay to Redcliff |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2259 | 113943685 |  | 04/06/2003 | Dace | 0 | Wareham Quay |
| 2260 | 113943670 |  | 03/06/2004 | Dace | 28 | Wareham quay to Redcliff |
| 2261 | 113943679 |  | 03/06/2004 | Dace | 14.7 | Wareham quay to Redcliff |
| 2262 | 113943705 |  | 03/06/2004 | Dace | 16.2 | Wareham quay to Redcliff |
| 2263 | 113943692 |  | 03/06/2004 | Dace | 19.8 | Wareham quay to Redcliff |
| 2264 | 113943683 |  | 03/06/2004 | Dace | 20.4 | Wareham quay to Redcliff |
| 2265 | 113943680 |  | 03/06/2004 | Dace | 20 | Wareham quay to Redcliff |
| 2266 | 113943666 |  | 03/06/2004 | Dace | 18 | Wareham quay to Redcliff |
| 2267 | 113943697 |  | 03/06/2004 | Dace | 16.2 | Wareham quay to Redcliff |
| 2268 | 113943699 |  | 03/06/2004 | Dace | 18 | Wareham quay to Redcliff |
| 2269 | 113943669 |  | 03/06/2004 | Dace | 17.1 | Wareham quay to Redcliff |
| 2270 | 113943688 |  | 03/06/2004 | Dace | 16.2 | Wareham quay to Redcliff |
| 2271 |  | DC0029-FFFE | 03/06/2004 | Dace | 21.4 | Wareham quay to Redcliff |
| 2272 |  | DC0029-FC8D | 03/06/2004 | Dace | 15.4 | Wareham quay to Redcliff |
| 2273 |  | DC002A-04D8 | 03/06/2004 | Dace | 23.1 | Wareham quay to Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2274 |  | DC0029-F83B | 03/06/2004 | Dace | 22.2 | Redcliff <br> Wareham quay to |
| 2275 |  | DC0029-F767 | 03/06/2004 | Dace | 21.8 | Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2276 |  | DC0029-EE37 | 03/06/2004 | Dace | 17.4 | Redcliff <br> Wareham quay to |
| 2277 |  | DC002A-1261 | 03/06/2004 | Dace | 21.8 | Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2278 |  | DC0029-F054 | 03/06/2004 | Dace | 17.2 | Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2279 |  | DC002A-045B | 03/06/2004 | Dace | 20.5 | Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2280 |  | DC002A-11BC | 03/06/2004 | Dace | 24.4 | Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2281 |  | DC002A-136D | 03/06/2004 | Dace | 16.2 | Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2282 |  | DC0029-FFA6 | 03/06/2004 | Dace | 15.2 | Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2283 |  | DC002A-129F | 03/06/2004 | Dace | 15.3 | Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2284 |  | DC002A-0822 | 03/06/2004 | Dace | 16.1 | Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2285 |  | DC002A-1678 | 03/06/2004 | Dace | 14.3 | Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2286 |  | DC002A-15C9 | 03/06/2004 | Dace | 27.3 | Redcliff |
|  |  |  |  |  |  | Wareham quay to |
| 2287 |  | DC002A-0653 | 03/06/2004 | Dace | 20.8 | Redcliff |
| 2288 |  | DC002A-1304 | 03/06/2004 | Dace | 12.8 | Wareham quay to |


| 2289 |  | DC002A-0878 | 03/06/2004 | Dace |
| :---: | :---: | :---: | :---: | :---: |
| 2290 |  | DC0029-EE2F | 03/06/2004 | Dace |
| 2291 |  | DC0029-F63E | 03/06/2004 | Dace |
| 2292 |  | DC0029-F7E8 | 03/06/2004 | Dace |
| 2293 |  | DC002A-04A8 | 03/06/2004 | Dace |
| 2294 |  | DC0029-ECF7 | 03/06/2004 | Dace |
| 2295 |  | DC002A-0335 | 03/06/2004 | Dace |
| 2296 |  | DC002A-11A2 | 03/06/2004 | Dace |
| 2297 |  | DC002A-0272 | 03/06/2004 | Dace |
| 2298 |  | DC0029-EF28 | 03/06/2004 | Dace |
| 2299 |  | DC002A-088F | 03/06/2004 | Dace |
| 2300 | 35471182 |  | 04/06/2003 | Dace |
| 2301 |  | DC0029-F6F5 | 04/06/2003 | Dace |
| 2302 | 35471227 |  | 04/06/2003 | Dace |
| 2303 |  | DC002A-1420 | 04/06/2003 | Dace |
| 2304 | 35471206 |  | 04/06/2003 | Dace |
| 2305 |  | DC002A-109A | 04/06/2003 | Dace |
| 2306 |  | DC0029-F819 | 04/06/2003 | Dace |
| 2307 | 35471232 |  | 04/06/2003 | Dace |
| 2308 |  | DC002A-1315 | 04/06/2003 | Dace |
| 2309 | 35471194 |  | 04/06/2003 | Dace |
| 2310 |  | DC002A-1660 | 04/06/2003 | Dace |
| 2311 | 35471224 |  | 04/06/2003 | Dace |
| 2312 |  | DC002A-0194 | 04/06/2003 | Dace |
| 2313 | 122451131 |  | 17/09/2004 | Dace |
| 2314 | 122451011 |  | 17/09/2004 | Dace |
| 2315 | 122451062 |  | 17/09/2004 | Dace |
| 2316 | 122451076 |  | 17/09/2004 | Dace |
| 2317 | 122451070 |  | 17/09/2004 | Dace |
| 2318 | 122451165 |  | 17/09/2004 | Dace |
| 2319 |  | DC0024-CB6A | 17/09/2004 | Dace |
| 2320 | 122451143 |  | 17/09/2004 | Dace |
| 2321 |  | DC002B-0B72 | 13/09/2004 | Pike |
| 2322 |  | DC002B-0448 | 13/09/2004 | Pike |
| 2323 | 122450618 | DC002B-110A | 13/09/2004 | Pike |
| 2324 |  | DC002B-16F2 | 13/09/2004 | Pike |
| 2325 |  | DC002B-10A1 | 13/09/2004 | Pike |
| 2326 |  | DC002A-F9F0 | 13/09/2004 | Pike |
| 2327 |  | DC002B-08B4 | 13/09/2004 | Pike |
| 2328 | 122451091 | DC0024-8883 | 13/09/2004 | Pike |
| 2329 | 122451049 |  | 14/09/2004 | Pike |
| 2340 |  | DC002B-0065 | 14/09/2004 | Dace |
| 2341 | 122451117 |  | 15/09/2004 | Pike |
| 2342 | 122451161 |  | 15/09/2004 | Pike |


|  | Redcliff |
| :---: | :---: |
|  | Wareham quay to |
| 17.8 | Redcliff |
|  | Wareham quay to |
| 27.2 | Redcliff |
|  | Wareham quay to |
| 19.8 | Redcliff |
|  | Wareham quay to |
| 20.5 | Redcliff |
|  | Wareham quay to |
| 24.2 | Redcliff |
|  | Wareham quay to |
| 17.4 | Redcliff |
|  | Wareham quay to |
| 15.4 | Redcliff |
|  | Wareham Quay to |
| 15.4 | Redcliff |
|  | Wareham quay to |
| 25 | Redcliff |
|  | Wareham quay to |
| 20.8 | Redcliff |
|  | Wareham quay to |
| 15.7 | Redcliff |
| 23.2 | Wareham Quay |
| 21.2 | Wareham Quay |
| 24.5 | Wareham Quay |
| 26.5 | Wareham Quay |
| 24 | Wareham Quay |
| 14.2 | Wareham Quay |
| 20.7 | Wareham Quay |
| 23.1 | Wareham Quay |
| 22.8 | Wareham Quay |
| 23.8 | Wareham Quay |
| 23.3 | Wareham Quay |
| 19.7 | Wareham Quay |
| 15 | Wareham Quay |
| 14 | ESMS 1 |
| 15.5 | ESMS 1 |
| 13.7 | ESMS 1 |
| 12.4 | ESMS 2 |
| 12.4 | ESMS 2 |
| 12 | ESMS 2 |
| 43 | ESMS 2 |
| 20.1 | ESMS 2 |
| 7.9 | Railway 1 |
| 9.1 | Railway 1 |
| 10.2 | Railway 1 |
| 9.5 | Railway 1 |
| 10 | Railway 1 |
| 8.9 | Railway 3 |
| 10 | Railway 3 |
| 24 | Railway 4 |
| 13.3 | Flood Relief 3 |
| 12.3 | Goldsacs 1 |
| 16.2 | Holme Bridge 1 |
| 13.9 | Holme Bridge 1 |


| 2343 |  | DC0024-D368 | 15/09/2004 | Pike |
| :---: | :---: | :---: | :---: | :---: |
| 2344 |  | DC002B-18C8 | 15/09/2004 | Pike |
| 2345 |  | DC0029-FB59 | 15/09/2004 | Pike |
| 2346 | 122451010 |  | 16/09/2004 | Pike |
| 2347 | 122451180 |  | 17/09/2004 | Dace |
| 2348 | 122451027 |  | 17/09/2004 | Dace |
| 2349 | 35471131 |  | 17/09/2004 | Dace |
| 2350 | 122451015 |  | 17/09/2004 | Dace |
| 2351 | 122451140 |  | 17/09/2004 | Dace |
| 2352 | 122451096 |  | 17/09/2004 | Dace |
| 2353 | 122451029 |  | 17/09/2004 | Dace |
| 2354 | 122451087 |  | 17/09/2004 | Dace |
| 2355 | 122451157 |  | 17/09/2004 | Dace |
| 2356 | 122451067 |  | 17/09/2004 | Dace |
| 2357 | 122451008 |  | 17/09/2004 | Dace |
| 2358 | 122451080 |  | 17/09/2004 | Dace |
| 2359 | 122451002 |  | 17/09/2004 | Dace |
| 2360 | 122451035 |  | 17/09/2004 | Dace |
| 2361 | 122451174 |  | 17/09/2004 | Dace |
| 2362 | 122451064 |  | 17/09/2004 | Dace |
| 2363 | 122451126 |  | 17/09/2004 | Dace |
| 2364 | 122450381 |  | 17/09/2004 | Dace |
| 2365 | 35471142 |  | 17/09/2004 | Dace |
| 2366 | 122451047 |  | 17/09/2004 | Dace |
| 2367 | 122451171 |  | 17/09/2004 | Dace |
| 2368 | 122451139 |  | 17/09/2004 | Dace |
| 2369 | 122451060 |  | 17/09/2004 | Dace |
| 2370 | 122451063 |  | 17/09/2004 | Dace |
| 2371 | 122451086 |  | 17/09/2004 | Eel |
| 2372 | 122451009 |  | 17/09/2004 | Eel |
| 2373 |  | DC0029-FA3C | 06/12/2004 | Pike |
| 2374 |  | DC0024-DD32 | 06/12/2004 | Pike |
| 2375 | 122450770 |  | 06/12/2004 | Pike |
| 2376 | 122450780 |  | 06/12/2004 | Pike |
| 2377 |  | DC002A-1396 | 06/12/2004 | Pike |
| 2378 |  | DC002A-0C3C | 06/12/2004 | Pike |
| 2379 | 122451166 |  | 06/12/2004 | Pike |
| 2380 | 122450995 |  | 06/12/2004 | Pike |
| 2381 | 122451135 |  | 06/12/2004 | Pike |
| 2382 | 34571148 |  | 06/12/2004 | Pike |
| 2383 | 122451017 |  | 06/12/2004 | Pike |
| 2384 | 122451144 |  | 06/12/2004 | Pike |
| 2385 | 122450999 |  |  | Pike |
| 2386 | 122451127 |  | 07/12/2004 | Pike |
| 2387 | 122451149 |  | 07/12/2004 | Pike |
| 2388 | 122451075 |  | 07/12/2004 | Pike |
| 2389 | 122451055 |  | 07/12/2004 | Pike |
| 2390 | 122451152 |  | 07/12/2004 | Pike |
| 2391 | 122451022 |  | 07/12/2004 | Perch |
| 2392 | 122450380 |  | 07/12/2004 | Pike |
| 2393 |  | DC002A-15DF | 07/12/2004 | Pike |
| 2394 | 122450771 |  | 08/12/2004 | Pike |
| 2395 | 122450774 |  | 08/12/2004 | Pike |
| 2396 | 122450773 |  | 08/12/2004 | Pike |


| 10.5 | Holme Bridge 4 |
| :---: | :---: |
| 10 | Holme bridge 4 |
| 9.6 | Holme Bridge Extra |
| 19.2 | Luckford Lake 4 |
| 13.1 | ESMS 3 |
| 15.4 | ESMS 3 |
| 17.2 | ESMS 3 |
| 18.7 | ESMS 3 |
| 15 | ESMS 3 |
| 15.6 | ESMS 3 |
| 14.2 | ESMS 3 |
| 17.5 | ESMS 3 |
| 18.1 | ESMS 3 |
| 14.8 | ESMS 3 |
| 18.2 | ESMS 3 |
| 18 | ESMS 3 |
| 19.8 | ESMS 3 |
| 18 | ESMS 3 |
| 17.6 | ESMS 3 |
| 18.5 | ESMS 3 |
| 18.3 | ESMS 3 |
| 18.9 | ESMS 3 |
| 17.3 | ESMS 3 |
| 18.3 | ESMS 3 |
| 18.4 | ESMS 3 |
| 16.5 | ESMS 3 |
| 14.5 | ESMS 3 |
| 40 | ESMS 3 |
| 47.5 | ESMS 3 |
| 34.6 | ESMS 3 |
| 11.2 | Railway 1 |
| 11.3 | Railway 1 |
| 31 | Railway 3 |
| 26.4 | Railway 3 |
| 12.1 | Railway 4 |
| 11.8 | Railway 4 |
| 27.5 | Flood Relief 1 |
| 24.5 | Flood Relief 1 |
| 20.2 | Flood Relief 1 |
| 18.3 | Flood Relief 1 |
| 13.5 | Flood Relief 1 |
| 18.5 | Flood Relief 2 |
| 13.5 | Flood Relief 2 |
| 37.4 | Rushton 1 |
| 36.4 | Rushton 1 |
| 20.5 | Rushton 1 |
| 16.7 | Rushton 1 |
| 30.5 | Rushton 1 |
| 18.9 | Rushton 2 |
| 16.4 | Rushton 4 |
| 10.5 | Rushton 4 |
| 19.2 | Holme Bridge 1 |
| 26 | Holme Bridge 1 |
| 22.3 | Holme Bridge 4 |


| 2397 | 122450776 |  | 08/12/2004 | Pike |
| :---: | :---: | :---: | :---: | :---: |
| 2398 |  | DC002B-0795 | 08/12/2004 | Pike |
| 2399 | 122450992 |  | 08/12/2004 | Pike |
| 2400 |  | DC0024-EC61 | 08/12/2004 | Pike |
| 2401 |  | DC002B-18D4 | 08/12/2004 | Pike |
| 2402 | 122451106 |  | 08/12/2004 | Pike |
| 2403 | 122450782 |  | 08/12/2004 | Pike |
| 2404 |  | DC002A-FC06 | 08/12/2004 | Pike |
| 2405 |  | DC002B-0022 | 08/12/2004 | Pike |
| 2406 | 122450778 |  | 09/12/2004 | Eel |
| 2407 | 122450987 |  | 09/12/2004 | Eel |
| 2408 | 122450781 |  | 09/12/2004 | Eel |
| 2409 | 122450988 |  | 09/12/2004 | Eel |
| 2410 | 122451068 |  | 09/12/2004 | Eel |
| 2411 | 122451036 |  | 09/12/2004 | Eel |
| 2412 | 122450984 |  | 09/12/2004 | Eel |
| 2413 | 122450989 |  | 09/12/2004 | Eel |
| 2414 | 122450779 |  | 09/12/2004 | Eel |
| 2415 | 122450777 |  | 09/12/2004 | Eel |
| 2416 | 122451054 |  | 09/12/2004 | Eel |
| 2417 | 122451089 |  | 09/12/2004 | Eel |
| 2418 | 122450772 |  | 09/12/2004 | grayling |
| 2419 | 122450775 |  | 09/12/2004 | Trout |
| 2420 | 122450985 |  | 09/12/2004 | Trout |
| 2421 | 122450986 |  | 09/12/2004 | Trout |
| 2422 | 122451098 |  | 09/12/2004 | Trout |
| 2423 | 122451147 |  | 09/12/2004 | Eel |
| 2424 | 122451043 |  | 09/12/2004 | Eel |
| 2425 | 122451065 |  | 09/12/2004 | Eel |
| 2426 | 122451133 |  | 09/12/2004 | Eel |
| 2427 | 122451018 |  | 09/12/2004 | Trout |
| 2428 | 122451172 |  | 09/12/2004 | Trout |
| 2430 | 122451079 |  | 09/12/2004 | Eel |
| 2431 | 122451050 |  | 09/12/2004 | Eel |
| 2432 | 122451007 |  | 10/12/2004 | Pike |
| 2433 | 122451048 |  | 10/12/2004 | Pike |
| 2434 | 122451094 |  | 10/12/2004 | Pike |
| 2435 | 122451019 |  | 10/12/2004 | Pike |
| 2436 | 122451020 |  | 10/12/2004 | Dace |
| 2437 | 122451176 |  | 10/12/2004 | Dace |
| 2438 | 122451168 |  | 10/12/2004 | Dace |
| 2439 | 122450378 |  | 10/12/2004 | Dace |
| 2440 | 122451181 |  | 10/12/2004 | Dace |
| 2441 | 122451104 |  | 10/12/2004 | Dace |
| 2442 | 122451038 |  | 10/12/2004 | Pike |
| 2443 | 122450996 |  | 10/12/2004 | Dace |
| 2444 | 122451078 |  | 10/12/2004 | Dace |
| 2445 | 122451041 |  | 10/12/2004 | Dace |
| 2446 | 122451112 |  | 10/12/2004 | Dace |
| 2447 | 122451085 |  | 10/12/2004 | Dace |
| 2448 | 122451146 |  | 10/12/2004 | Dace |
| 2449 | 122451111 |  | 10/12/2004 | Dace |
| 2450 | 122451170 |  | 10/12/2004 | Dace |
| 2451 | 122451118 |  | 10/12/2004 | Dace |

26.9 Holme Bridge 4
10.7 Holme Bridge 4

24 Holme Bridge 4
10.8 Holme Bridge 4
13.6 Holme Bridge 4

35 Holme Bridge Extra
30.9 Holme Bridge Extra
9.5 Holme Bridge Extra

11 Holme Bridge Extra
39.5 ESMS 1
47.5 ESMS 1
34.6 ESMS 1
39.4 ESMS 1

ESMS 1
39.4 ESMS 2
46.2 ESMS 2
49.4 ESMS 2

30 ESMS 2
39.2 ESMS 2
33.2 ESMS 2
40.3 ESMS 2

30 ESMS 3
26.3 ESMS 3
26.3 ESMS 3
20.7 ESMS 3
20.5 ESMS 3
36.6 ESMS 3
36.7 ESMS 3
45.8 ESMS 3
47.3 ESMS 3
26.7 ESMS 3
21.7 ESMS 3
33.8 ESMS 4
36.9 ESMS
49.8 Luckford Lake 1
43.5 Luckford Lake 1
42.5 Luckford Lake 1
44.4 Luckford Lake 1
20.7 Luckford Lake 2
12.2 Luckford Lake 2

19 Luckford Lake 2
13.5 Luckford Lake 2
15.2 Luckford Lake 2
13.1 Luckford Lake 2
21.7 Luckford Lake 3
18.2 Luckford Lake 3
16.2 Luckford Lake 3
12.3 Luckford Lake 3

16 Luckford Lake 3
18 Luckford Lake 3
15.1 Luckford Lake 3
13.8 Luckford Lake 3
14.9 Luckford Lake 3
13.6 Luckford Lake 3

| 2452 | 35471147 |
| ---: | ---: |
| 2453 | 122451013 |
| 2454 | 122450382 |
| 2455 | 122451124 |
| 2456 | 122451110 |
| 2457 | 122451090 |
| 2458 | 122450994 |
| 2459 | 122450998 |
| 2460 | 122450991 |
| 2461 | 35471149 |
| 2462 | 122451021 |
| 2463 | 122450997 |
| 2464 | 122450990 |
| 2465 | 113943710 |
| 2466 | 122450993 |
| 2467 | 122451023 |
| 2468 | 122451132 |
| 2469 | 127178822 |
| 2470 | 127178823 |
| 2471 | 127178824 |
| 2472 | 127178825 |
| 2473 | 127178826 |
| 2474 | 127178827 |
| 2475 | 127178828 |
| 2476 | 127178829 |
| 2477 | 127178830 |
| 2478 | 127178831 |
| 2479 | 127179182 |
| 2480 | 127179183 |
| 2481 | 127179184 |
| 2482 | 127179185 |
| 2483 | 127179186 |
| 2484 | 127178912 |
| 2485 | 127178911 |
| 2486 | 127178910 |
| 2487 | 127178909 |
| 2489 | 127178908 |
| 2490 | 127179006 |
| 2491 | 127179005 |
| 2492 | 127179004 |
| 2493 | 127179003 |
| 2495 | 127179001 |
| 2496 | 127179000 |
| 2497 | 127178999 |
| 2498 | 127178998 |
| 2499 | 127178997 |
| 2500 | 127178996 |
| 2501 | 127178995 |
| 2502 | 127178994 |
| 2503 | 127178993 |
| 2504 | 127178992 |
| 2505 | 127178991 |
| 2506 | 127178990 |
| 2507 | 127178989 |
|  |  |


| $10 / 12 / 2004$ | Dace |
| :--- | :--- |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Pike |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Dace |
| $10 / 12 / 2004$ | Dace |
| $10 / 2004$ | Dace |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Roach |
| $10 / 12 / 2004$ | Dace |
| $10 / 2 / 2004$ | Dace |
| $10 / 2004$ | Dace |
| $10 / 2004$ | Dace |
| $10 / 2$ |  |

13.9 Luckford Lake 3
14.5 Luckford Lake 3
13.7 Luckford Lake 3
15.8 Luckford Lake 3

16 Luckford Lake 3
19 Luckford Lake 3
14.7 Luckford Lake 3
18.4 Luckford Lake 3
14.8 Luckford Lake 3

15 Luckford Lake 3
13.3 Luckford Lake 3
13.2 Luckford Lake 3
12.7 Luckford Lake 3
13.9 Luckford Lake 3
13.3 Luckford Lake 3
14.5 Luckford Lake 3
14.6 Luckford Lake 3
23.6 Luckford lake 4
20.1
17.1 Luckford lake 4
16.5 Luckford lake 4
18.3 Luckford lake 4
37.8 Luckford lake 4
19.1 Luckford lake 4
17.5 Luckford lake 4
19.6 Luckford lake 4
18.5 Luckford lake 4
16.6 Luckford lake 4
20.9 Luckford lake 4
23.4 Luckford lake 4
22.4 Luckford lake 4
18.2 Luckford lake 4
20.5 Luckford lake 4
15.5 Luckford lake 4
20.9 Luckford lake 4
16.4 Luckford lake 4

22 Luckford lake 4
17.3 Luckford lake 4
15.6 Luckford lake 4
24.8 Luckford lake 4
17.6 Luckford lake 4
16.9 Luckford lake 4
18.4 Luckford lake 4
18.2 Luckford lake 4
17.7 Luckford lake 4
25.1 Luckford lake 4
20.3 Luckford Lake 4
35.7 Luckford Lake 4
21.6 Luckford Lake 4
17.3 Luckford Lake 4

19 Luckford Lake 4
24.1 Luckford Lake 4
16.6 Luckford Lake 4
20.2 Luckford Lake 4

| 2508 | 127178988 |  | 10/12/2004 | Roach |
| :---: | :---: | :---: | :---: | :---: |
| 2509 | 127178987 |  | 10/12/2004 | Roach |
| 2510 | 127178986 |  | 10/12/2004 | Dace |
| 2511 | 127178985 |  | 10/12/2004 | Dace |
| 2512 | 127178984 |  | 10/12/2004 | Roach |
| 2513 | 127178956 |  | 10/12/2004 | Dace |
| 2514 | 127178955 |  | 10/12/2004 | Dace |
| 2515 | 127178954 |  | 10/12/2004 | Roach |
| 2516 | 127178953 |  | 10/12/2004 | Roach |
| 2517 | 127178952 |  | 10/12/2004 | Roach |
| 2518 | 127178950 |  | 10/12/2004 | Dace |
| 2519 | 127178951 |  | 10/12/2004 | Dace |
| 2520 | 127178949 |  | 10/12/2004 | Roach |
| 2521 | 127178948 |  | 10/12/2004 | Roach |
| 2522 | 127178947 |  | 10/12/2004 | Dace |
| 2523 | 127178946 |  | 10/12/2004 | Roach |
| 2524 | 127178945 |  | 10/12/2004 | Dace |
| 2525 | 122450609 |  | 17/03/2005 | Pike |
| 2526 | 122450611 |  | 17/03/2005 | Pike |
| 2527 | 122450610 |  | 17/03/2005 | Dace |
| 2528 | 122450608 |  | 23/03/2005 | Pike |
| 2529 | 122450766 |  | 23/03/2005 | Pike |
| 2530 | 122450764 |  | 23/03/2005 | Pike |
| 2531 |  | DC0024-D994 | 23/03/2005 | Pike |
| 2532 |  | DC0024-EC3E | 23/03/2005 | Pike |
| 2533 |  | DC002B-00AA | 23/03/2005 | Pike |
| 2534 |  | DC002B-13A3 | 23/03/2005 | Pike |
| 2535 | 122450741 |  | 08/03/2005 | Pike |
| 2536 | 122450743 |  | 08/03/2005 | Pike |
| 2537 | 122450746 |  | 08/03/2005 | Pike |
| 2538 | 122450747 |  | 08/03/2005 | Pike |
| 2539 | 122450751 |  | 08/03/2005 | Perch |
| 2541 | 122450655 | DC002B-0C3F | 17/03/2005 | Dace |
| 2542 | 122450654 |  | 17/03/2005 | Dace |
| 2543 | 122450653 | DC002A-0768 | 17/03/2005 | Dace |
| 2544 | 122450650 |  | 16/03/2005 | Dace |
| 2545 | 122450647 |  | 17/03/2005 | Pike |
| 2546 | 122450642 | DC002A-061E | 17/03/2005 | Dace |
| 2547 | 122450646 | DC002A-0AA8 | 17/03/2005 | Dace |
| 2548 | 122450643 |  | 17/03/2005 | Dace |
| 2549 | 122450640 |  | 17/03/2005 | Dace |
| 2550 | 122450637 | DC0029-ECAO | 17/03/2005 | Dace |
| 2551 | 122450639 | DC0029-EFE7 | 17/03/2005 | Dace |
| 2552 | 122450633 |  | 17/03/2005 | Dace |
| 2553 | 122450628 |  | 17/03/2005 | Dace |
| 2554 | 122450627 |  | 17/03/2005 | Pike |
| 2555 | 122450613 |  | 17/03/2005 | Trout |
| 2556 | 122450626 |  | 17/03/2005 | Trout |
| 2557 | 122450630 |  | 17/03/2005 | Trout |
| 2558 | 122450631 |  | 17/03/2005 | Trout |
| 2559 | 122450634 |  | 17/03/2005 | Dace |
| 2560 | 122450615 |  | 17/03/2005 | Trout |
| 2561 | 122450622 |  | 17/03/2005 | Trout |

18.4 Luckford Lake 4
17.6 Luckford Lake 4
18.8 Luckford Lake 4
17.1 Luckford Lake 4
16.6 Luckford Lake 4
26.7 Luckford Lake 4
26.7 Luckford Lake
15.3 Luckford Lake 4
19.8 Luckford Lake 4
18.8 Luckford Lake 4
18.9 Luckford Lake 4
19.4 Luckford Lake 4
16.3 Luckford Lake 4
16.1 Luckford Lake 4
15.7 Luckford Lake 4
17.8 Luckford Lake 4
17.8 Luckford Lake 4
25.8 ESMS
27.7 ESMS
19.5 ESMS

26 Railway
20.4 Railway 1

18 Railway 3
11.9 Railway
11.7 Railway
11.2 Railway
10.8 Railway
25.5 River Frome T3
19.5 Main River Frome T7
46.8 Main River Frome T10-12
20.3 Main River Frome T10-12 Main River Frome Frog
39.5 T20
22.3 ESMS 1
20.5 ESMS 1
23.9 ESMS 1
18.2 Goldsacs 4
19.1 ESMS 2
27.3 ESMS 2
20.6 ESMS 2
18.5 ESMS 2
18.2 ESMS 2
22.6 ESMS 3
22.8 ESMS 3
20.1 ESMS 2
20.4 ESMS 2
24.9 ESMS 2

14 ESMS 2
13.7 ESMS 2
12.9 ESMS 2
14.4 ESMS 2
17.4 ESMS 2
15.8 ESMS 4
14.1 ESMS 4

| 2562 | 122450661 |  | 16/03/2005 | Dace |
| :---: | :---: | :---: | :---: | :---: |
| 2563 | 122450685 |  | 16/03/2005 | Dace |
| 2564 | 122450662 |  | 16/03/2005 | Dace |
| 2565 | 122450663 |  | 16/03/2005 | Dace |
| 2566 | 122450660 |  | 16/03/2005 | Dace |
| 2567 | 122450659 |  | 16/03/2005 | Dace |
| 2568 | 122450656 |  | 16/03/2005 | Dace |
| 2569 | 122450657 |  | 16/03/2005 | Dace |
| 2570 | 122450666 |  | 16/03/2005 | Dace |
| 2571 | 122450665 |  | 16/03/2005 | Dace |
| 2572 | 122450664 |  |  | Dace |
| 2573 | 113945640 |  |  | Dace |
| 2575 |  | DC0029-F363 | 15/03/2005 | Pike |
| 2576 |  | DC002A-0EE6 | 15/03/2005 | Pike |
| 2577 | 122450679 |  | 15/03/2005 | Dace |
| 2578 | 122450675 |  | 15/03/2005 | Roach |
| 2579 | 122450678 |  | 15/03/2005 | Dace |
| 2580 | 122450677 |  | 15/03/2005 | Dace |
| 2581 | 122450672 |  | 15/03/2005 | Dace |
| 2582 | 122450673 |  | 15/03/2005 | Dace |
| 2583 | 122450676 |  | 15/03/2005 | Dace |
| 2584 | 122450674 |  | 15/03/2005 | Dace |
| 2585 | 122450671 |  | 15/03/2005 | Dace |
| 2586 | 122450670 |  | 15/03/2005 | Dace |
| 2587 | 122450669 |  | 15/03/2005 | Roach |
| 2588 | 122450668 |  | 15/03/2005 | Dace |
| 2589 | 122450667 |  | 15/03/2005 | Dace |
| 2590 | 122450691 |  | 12/07/2004 | Pike |
| 2591 | 122450690 | DC002A-161B | 15/03/2005 | Dace |
| 2592 | 122450688 |  | 15/03/2005 | Pike |
| 2593 | 122450689 |  | 15/03/2005 | Pike |
| 2594 | 122450686 |  | 15/03/2005 | Pike |
| 2595 | 122450684 |  | 15/03/2005 | Pike |
| 2596 | 122450683 |  | 15/03/2005 | Pike |
| 2597 | 122450682 |  | 15/03/2005 | Pike |
| 2598 | 122450681 |  | 15/03/2005 | Pike |
| 2599 | 122450680 |  | 15/03/2005 | Pike |
| 2600 | 122450755 |  | 15/03/2005 | Pike |
| 2601 | 122450753 |  | 14/03/2005 | Perch |
| 2602 | 122450700 |  | 14/03/2005 | Pike |
| 2603 | 122450699 |  | 14/03/2005 | Pike |
| 2604 | 122450697 |  | 14/03/2005 | Pike |
| 2605 | 122450696 |  | 14/03/2005 | Pike |
| 2606 | 122450695 |  | 14/03/2005 | Pike |
| 2607 | 122450694 |  | 14/03/2005 | Pike |
| 2608 | 122450768 |  | 14/03/2005 | Pike |
| 2609 | 122450767 |  | 14/03/2005 | Pike |
| 2610 |  | DC0024-D99B | 14/03/2005 | Pike |
| 2611 |  | DC002A-0C88 | 14/03/2005 | Pike |
| 2612 | 122450765 |  | 14/03/2005 | Pike |
| 2614 | 122450763 |  | 14/03/2005 | Pike |
| 2616 | 122450762 |  | 14/03/2005 | Pike |
| 2617 | 122450761 |  | 14/03/2005 | Pike |

19.4 Goldsacs 2

18 Goldsacs 2
17.5 Goldsacs 2
23.7 Goldsacs 2
21.9 Goldsacs 2
23.2 Goldsacs 2
18.4 Goldsacs 4
19.7 Goldsacs 4
12.2 Luckford Lake 4

15 Luckford Lake 4
12.1 Luckford Lake 4

26 Luckford Lake 2
12 Holme Bridge 4
11.1 Holme Bridge 4
27.3 Luckford lake 4
21.2
17.7 Luckford lake 4
14.6 Luckford lake 4
13.1 Luckford lake 4
14.4 Luckford lake 4
16.5 Luckford lake 4
14.6 Luckford lake 4
16.1 Luckford lake 4
12.3 Luckford lake 4
23.9 Luckford lake 4
13.7
15.3 Luckford lake 4
94.5

Main River Frome us
26.4 Luckford

73 Main River Frome Oxbow
56.9 Luckford Lake 1

44 Luckford Lake 1
22.2 Holme Bridge 1
22.3 Holme Bridge 1
21.1 Holme Bridge 1
25.7 Holme Bridge 1
15.8 Holme Bridge 3
12.4 Rushton 1
19.5 Rushton 1
23.3 Rushton 2
22.2 Rushton 2
19.6 Rushton 2
16.7 Rushton 2

28 Rushton 4
22.4 Rushton 4
16.3 Railway 1
19.4 Railway 1
11.6 Railway 1
12.2 Railway 2
22.2 Railway 3
21.5 Flood Relief 1
13.8 Flood Relief 1
16.3 Flood Relief 1

| 2618 | 122450760 |  | $14 / 03 / 2005$ | Pike | 13.8 | Flood Relief 1 |
| :--- | :--- | :--- | ---: | :--- | ---: | :--- |
| 2619 | 122450759 |  | $14 / 03 / 2005$ | Pike | 16.3 | Flood Relief 2 |
| 2620 | 122450757 |  | $14 / 03 / 2005$ | Pike | 27.2 | Flood Relief 2 |
| 2621 |  | DC002B-0B76 | $14 / 03 / 2005$ | Pike | 13 | Flood Relief 2 |
| 2622 | 122450756 |  | $14 / 03 / 2005$ | Pike | 15.8 | Flood Relief 2 |
| 2944 | 127179002 |  | $10 / 2 / 2004$ | Dace | 23.2 | Luckford lake 4 |
| 2945 | 122450632 |  | $21 / 06 / 2005$ | Pike | 17.8 | Rushton 1 |
| 2946 | 122450629 |  | $21 / 06 / 2005$ | Pike | 18.1 | Luckford Lake 4 |
| 2947 | 122450621 |  | $22 / 06 / 2005$ | Dace | 12.9 | ESMS 3 |
| 2948 | 122450617 |  | $22 / 06 / 2005$ | Dace | 12.6 | ESMS 3 |
| 2949 | 122450619 |  | $22 / 06 / 2005$ | Dace | 12.8 |  |
| 2950 | 122450602 |  | $22 / 06 / 2005$ | Gudgeon | 14.3 | ESMS 3 |
| 2951 | 122450596 |  | $22 / 06 / 2005$ | Gudgeon | 12.9 | ESMS 3 |
| 2952 | 122450636 |  | $22 / 06 / 2005$ | Eel | 35 | ESMS 3 |
| 2953 | 122450595 |  | $22 / 06 / 2005$ | Pike | 66.6 | ESMS 3 |
| 2954 | 122450599 |  | $22 / 06 / 2005$ | Eel | 48.5 | ESMS 4 |
| 2955 | 122450638 |  | $22 / 06 / 2005$ | Roach | 15.4 | ESMS 5 |
| 2956 | 122450612 |  | $23 / 06 / 2005$ | Pike | 14 | Railway 1 |
| 2957 |  | DC002B-0971 | $23 / 06 / 2005$ | Pike | 13.9 | Railway 2 |
| 2958 |  | DC002B-11BC | $23 / 06 / 2005$ | Pike | 8.3 | Flood Relief 1 |
| 2959 |  | DC0024-7E72 | $23 / 06 / 2005$ | Pike | 7.5 | Flood Relief 1 |
| 2960 |  | DC002B-194D | $23 / 06 / 2005$ | Pike | 7.2 | Flood Relief 3 |
| 2961 | 122450360 |  | $12 / 07 / 2005$ | Roach | 20.4 | Wareham |
| 2962 |  | DC0029-F7F6 | $12 / 7 / 2005$ | Roach | 21.5 | Wareham |
| 2963 | 122450363 |  | $12 / 7 / 2005$ | Rooch | 20.8 | Wareham |
| 2964 |  | DC002A-04A7 | $12 / 07 / 2005$ | Roach | 21.3 | Wareham |
| 2965 | 122450371 |  | $12 / 07 / 2005$ | Roach | 24.5 | Wareham |
| 2966 |  | DC002B-0995 | $12 / 07 / 2005$ | Roach | 19.3 | Wareham |
| 2967 | 122450374 |  | $12 / 07 / 2005$ | Roach | 18.6 | Wareham |
| 2968 |  |  | DC0029-EBE9 | $12 / 07 / 2005$ | Roach | 19.3 |


| 2993 | 122450353 |  | 12/07/2005 | Roach | 22.1 | Wareham |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 2994 |  | DC002A-09D0 | 12/07/2005 | Roach | 20.1 | Wareham |
| 2995 | 122450337 |  | 12/07/2005 | Roach | 15.1 | Wareham |
| 2996 |  | DC002A-028E | 12/07/2005 | Dace | 26.6 | Wareham |
| 2997 | 122450336 |  | 12/07/2005 | Roach | 22.8 | Wareham |
| 2998 |  | DC002A-0B76 | 12/07/2005 | Roach | 26.1 | Wareham |
| 2999 | 122450351 |  | 12/07/2005 | Roach | 13.8 | Wareham |
| 3000 |  | DC002A-0EB0 | 12/07/2005 | Roach | 23.8 | Wareham |
| 3001 | 122450341 |  | 12/07/2005 | Roach | 24.8 | Wareham |
| 3002 |  | DC002B-16EE | 12/07/2005 | Roach | 15.7 | Wareham |
| 3003 | 122450347 |  | 12/07/2005 | Roach | 19.8 | Wareham |
| 3004 |  | DC0029-F26D | 12/07/2005 | Roach | 26.3 | Wareham |
| 3005 | 122450340 |  | 12/07/2005 | Roach | 25.1 | Wareham |
| 3006 |  | DC0029-FAAO | 12/07/2005 | Roach | 19.4 | Wareham |
| 3007 | 122450339 |  | 12/07/2005 | Roach | 21.9 | Wareham |
| 3008 |  | DC002A-F293 | 12/07/2005 | Roach | 16.5 | Wareham |
| 3009 | 122450346 |  | 12/07/2005 | Roach | 12.9 | Wareham |
| 3010 |  | DC002A-0DF5 | 12/07/2005 | Roach | 20.5 | Wareham |
| 3011 | 122450355 |  | 12/07/2005 | Dace | 12.8 | Wareham |
| 3012 |  | DC002A-F6EE | 12/07/2005 | Dace | 13.1 | Wareham |
| 3013 |  | DC002B-1172 | 12/07/2005 | Dace | 12.3 | Wareham |
| 3014 | 122450345 |  | 12/07/2005 | Roach | 37.3 | Wareham |
| 3015 |  | DC0029-F4D7 | 12/07/2005 | Roach | 32.5 | Wareham |
| 3016 | 122450343 |  | 12/07/2005 | Roach | 22.1 | Wareham |
| 3017 |  | DC0024-7274 | 12/07/2005 | Dace | 12.8 | Wareham |
| 3018 |  | DC0029-CF35 | 12/07/2005 | Dace | 24.4 | Wareham |
| 3019 | 122450338 |  | 12/07/2005 | Roach | 25 | Wareham |
| 3020 |  | DC002A-0B29 | 12/07/2005 | Roach | 18.1 | Wareham |
| 3021 | 122450332 |  | 12/07/2005 | Dace | 24.2 | Wareham |
| 3022 |  | DC002A-0EB8 | 12/07/2005 | Roach | 35.8 | Wareham |
| 3023 | 122450328 |  | 12/07/2005 | Roach | 20.7 | Wareham |
| 3024 |  | DC002A-F80E | 12/07/2005 | Dace | 18.8 | Wareham |
| 3025 | 122450331 |  | 12/07/2005 | Roach | 12.9 | Wareham |
| 3026 |  | DC002A-12F4 | 12/07/2005 | Roach | 18 | Wareham |
| 3027 | 122450334 |  | 12/07/2005 | Dace | 17.6 | Wareham |
| 3028 |  | DC002A-086F | 13/07/2005 | Dace | 21.1 | Wareham |
| 3029 | 122450329 |  | 13/07/2005 | Dace | 19.8 | Wareham |
| 3030 |  | DC002A-0CE5 | 13/07/2005 | Roach | 17.1 | Wareham |
| 3031 | 122450330 |  | 13/07/2005 | Dace | 23.1 | Wareham |
| 3032 |  | DC002A-F613 | 13/07/2005 | Dace | 13.6 | Wareham |
| 3033 | 122450316 |  | 13/07/2005 | Roach | 35.7 | Wareham |
| 3034 |  | DC0029-EF35 | 13/07/2005 | Dace | 24.2 | Wareham |
| 3035 |  | DC0029-F8CA | 13/07/2005 | Dace | 13.6 | Wareham |
| 3036 | 122450327 |  | 13/07/2005 | Roach | 13 | Wareham |
| 3037 |  | DC002A-088C | 13/07/2005 | Dace | 26.6 | Wareham |
| 3038 | 122450322 |  | 13/07/2005 | Roach | 23.1 | Wareham |
| 3039 |  | DC002B-000D | 13/07/2005 | Roach | 34.4 | Wareham |
| 3040 | 122450344 |  | 13/07/2005 | Roach | 26.4 | Wareham |
| 3041 |  | DC0029-EF3A | 13/07/2005 | Dace | 21.9 | Wareham |
| 3042 | 122450321 |  | 13/07/2005 | Dace | 19.7 | Wareham |
| 3043 |  | DC002B-1C07 | 13/07/2005 | Roach | 35.3 | Wareham |
| 3044 | 122450333 |  | 13/07/2005 | Dace | 12.4 | Wareham |
| 3045 |  | DC0029-F086 | 13/07/2005 | Dace | 24.5 | Wareham |
| 3046 | 122450320 |  | 13/07/2005 | Roach | 24.1 | Wareham |


| 3047 |  | DC002B-0D6A | 13/07/2005 | Roach | 30.4 | Wareham |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 3048 | 122450317 |  | 13/07/2005 | Roach | 15.4 | Wareham |
| 3049 |  | DC0024-CC3E | 13/07/2005 | Dace | 25.1 | Wareham |
| 3050 | 122450342 |  | 13/07/2005 | Dace | 26.8 | Wareham |
| 3051 | 122450323 |  | 13/07/2005 | Roach | 18.3 | Wareham |
| 3052 |  | DC0029-F755 | 13/07/2005 | Roach | 27.5 | Wareham |
| 3053 | 122450326 |  | 13/07/2005 | Dace | 24.5 | Wareham |
| 3054 |  | DC002A-01F5 | 13/07/2005 | Roach | 21.2 | Wareham |
| 3055 | 122450319 |  | 13/07/2005 | Dace | 16.4 | Wareham |
| 3056 |  | DC002A-10F2 | 13/07/2005 | Dace | 20.5 | Wareham |
| 3057 | 122450310 |  | 13/07/2005 | Roach | 27.3 | Wareham |
| 3058 |  | DC0029-F5B2 | 13/07/2005 | Roach | 21.2 | Wareham |
| 3059 | 122450315 |  | 13/07/2005 | Dace | 24.5 | Wareham |
| 3060 |  | DC0029-EE36 | 13/07/2005 | Dace | 23 | Wareham |
| 3061 | 122450313 |  | 13/07/2005 | Roach | 28.2 | Wareham |
| 3062 |  | DC002A-F631 | 13/07/2005 | Roach | 13.1 | Wareham |
| 3063 | 122450308 |  | 13/07/2005 | Dace | 14 | Wareham |
| 3064 |  | DC002A-FFA8 | 13/07/2005 | Dace | 21.9 | Wareham |
| 3065 | 122450312 |  | 13/07/2005 | Dace | 12.2 | Wareham |
| 3066 |  | DC002A-1423 | 13/07/2005 | Dace | 12 | Wareham |
| 3067 | 122450309 |  | 13/07/2005 | Dace | 13.9 | Wareham |
| 3068 |  | DC0029-F762 | 13/07/2005 | Roach | 32.3 | Wareham |
| 3069 |  | DC002B-01C4 | 13/07/2005 | Dace | 20.7 | Wareham |
| 3070 | 122450324 |  | 13/07/2005 | Dace | 19.9 | Wareham |
| 3071 |  | DC002A-085D | 13/07/2005 | Dace | 23.2 | Wareham |
| 3072 | 122450311 |  | 13/07/2005 | Dace | 17.2 | Wareham |
| 3073 |  | DC002A-048E | 13/07/2005 | Dace | 12.5 | Wareham |
| 3074 | 122450302 |  | 13/07/2005 | Roach | 12.5 | Wareham |
| 3075 |  | DC002A-0346 | 13/07/2005 | Dace | 13.1 | Wareham |
| 3076 | 122450277 |  | 13/07/2005 | Dace | 21.7 | Wareham |
| 3077 |  | DC002A-0BDD | 13/07/2005 | Dace | 25.1 | Wareham |
| 3078 | 122450297 |  | 13/07/2005 | Roach |  | Wareham |
| 3079 |  | DC002B-17C5 | 13/07/2005 | Roach | 24.3 | Wareham |
| 3080 | 122450299 |  | 13/07/2005 | Roach | 19.7 | Wareham |
| 3081 |  | DC002A-0B61 | 13/07/2005 | Dace | 16.1 | Wareham |
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| 3083 |  | DC002A-0900 | 13/07/2005 | Roach | 12.5 | Wareham |
| 3084 | 122450306 |  | 13/07/2005 | Dace | 12.4 | Wareham |
| 3085 |  | DC0029-EEE6 | 13/07/2005 | Roach | 12.3 | Wareham |
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| 3090 |  | DC002B-14EF | 13/07/2005 | Dace | 12.7 | Wareham |
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| 3092 |  | DC0029-EC5F | 13/07/2005 | Dace | 12.4 | Wareham |
| 3093 | 122450314 |  | 13/07/2005 | Dace | 13.7 | Wareham |
| 3094 |  | DC002A-FD07 | 13/07/2005 | Dace | 12.6 | Wareham |
| 3095 | 122450296 |  | 13/07/2005 | Dace | 14.5 | Wareham |
| 3096 |  | DC002A-0DA7 | 13/07/2005 | Roach | 12.5 | Wareham |
| 3097 | 122450301 |  | 13/07/2005 | Dace | 12.5 | Wareham |
| 3098 |  | DC002A-FF43 | 13/07/2005 | Roach | 12.7 | Wareham |
| 3099 | 122450300 |  | 13/07/2005 | Dace | 13.1 | Wareham |
| 3100 |  | DC002A-0199 | 13/07/2005 | Dace | 12.3 | Wareham |


| 3101 | 122450305 |  | 13/07/2005 | Dace | 12.2 | Wareham |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 3102 |  | DC002A-FBE9 | 13/07/2005 | Dace | 12.2 | Wareham |
| 3103 | 122450616 |  | 12/07/2005 | Dace | 12.8 | Wareham |
| 3104 | 122450635 |  | 12/07/2005 | Dace | 17.9 | Wareham |
| 3105 | 122450625 |  | 12/07/2005 | Dace | 13.2 | Wareham |
| 3106 | 122450601 |  | 12/07/2005 | Dace | 20.1 | Wareham |
| 3107 | 122450624 |  | 12/07/2005 | Roach | 13.3 | Wareham |
| 3108 | 122450600 |  | 12/07/2005 | Dace | 21.8 | Wareham |
| 3109 |  | DC002A-016E | 12/07/2005 | Roach | 16.1 | Wareham |
| 3110 |  | DC002A-148E | 12/07/2005 | Rudd | 17.6 | Wareham |
| 3111 | 122450593 |  | 12/07/2005 | Roach | 15.8 | Wareham |
| 3112 |  | DC002B-1475 | 12/07/2005 | Roach | 12.7 | Wareham |
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| 3115 | 122450592 |  | 12/07/2005 | Dace | 16.8 | Wareham |
| 3116 |  | DC0029-FAA9 | 12/07/2005 | Dace | 16.9 | Wareham |
| 3117 | 122450589 |  | 13/07/2005 | Roach | 23.1 |  |
| 3118 |  | DC0024-8BCD | 12/07/2005 | Dace | 12.5 | Wareham |
| 3119 | 122450591 |  | 12/07/2005 | Dace | 12.5 | Wareham |
| 3120 |  | DC002A-060F | 12/07/2005 | Roach | 15.7 | Wareham |
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| 3122 |  | DC0029-FE3A | 12/07/2005 | Roach | 16 | Wareham |
| 3123 | 122450590 |  | 12/07/2005 | Roach | 35.8 | Wareham |
| 3124 |  | DC002A-06A9 | 12/07/2005 | Roach | 17.3 | Wareham |
| 3125 | 122450588 |  | 12/07/2005 | Roach | 23.9 | Wareham |
| 3126 |  | DC0024-8CBB | 12/07/2005 | Dace | 20.8 | Wareham |
| 3127 | 122450584 |  | 12/07/2005 | Roach | 18.3 | Wareham |
| 3128 |  | DC0029-F34D | 12/07/2005 | Roach | 20.3 | Wareham |
| 3129 | 122450585 |  | 12/07/2005 | Roach | 29.2 | Wareham |
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| 3131 | 122450586 |  | 12/07/2005 | Dace | 25.7 | Wareham |
| 3132 |  | DC002A-00DO | 12/07/2005 | Dace | 12.6 | Wareham |
| 3133 | 122450376 |  | 12/07/2005 | Dace | 16.7 | Wareham |
| 3134 |  | DC002A-0F2C | 12/07/2005 | Roach | 13.5 | Wareham |
| 3135 | 122450375 |  | 12/07/2005 | Dace | 19.2 | Wareham |
| 3136 |  | DC002B-1444 | 12/07/2005 | Dace | 16.5 | Wareham |
| 3137 | 122450583 |  | 12/07/2005 | Roach | 24.7 | Wareham |
| 3138 |  | DC002A-064C | 12/07/2005 | Dace | 18 | Wareham |
| 3139 |  | DC002B-130C | 12/07/2005 | Roach | 13 | Wareham |
| 3140 |  | DC002B-186A | 12/07/2005 | Dace | 18.2 | Wareham |
| 3141 |  | DC002B-0246 | 12/07/2005 | Roach | 14 | Wareham |
| 3142 |  | DC002B-09A7 | 12/07/2005 | Roach | 14 | Wareham |
| 3143 | 122450372 |  | 12/07/2005 | Roach | 13.4 | Wareham |
| 3144 |  | DC002A-0919 | 12/07/2005 | Dace | 21.1 | Wareham |
| 3145 | 122450373 |  | 12/07/2005 | Dace | 25.3 | Wareham |
| 3146 |  | DC0029-FA18 | 12/07/2005 | Dace | 26.7 | Wareham |
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| 3149 | 122450364 |  | 12/07/2005 | Roach | 12.6 | Wareham |
| 3150 |  | DC0024-EDFE | 12/07/2005 | Roach | 12.3 | Wareham |
| 3151 | 122450362 |  | 12/07/2005 | Dace | 12.3 | Wareham |
| 3152 |  | DC002A-0F15 | 12/07/2005 | Dace | 12.6 | Wareham |
| 3153 | 122450368 |  | 12/07/2005 | Dace | 18.5 | Wareham |
| 3154 |  | DC002B-0B46 | 12/07/2005 | Dace | 15.5 | Wareham |


| 3155 | 122450370 |  | 12/07/2005 | Dace |
| :---: | :---: | :---: | :---: | :---: |
| 3156 |  | DC002A-0808 | 12/07/2005 | Dace |
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| 3158 |  | DC002A-F204 | 12/07/2005 | Dace |
| 3159 | 122450303 |  | 12/09/2005 | Pike |
| 3160 | 133913381 |  | 12/09/2005 | Dace |
| 3161 | 122450307 |  | 12/09/2005 | Roach |
| 3162 | 133913380 |  | 12/09/2005 | Dace |
| 3163 | 133913379 |  | 12/09/2005 | Dace |
| 3164 | 133913378 |  | 12/09/2005 | Dace |
| 3165 | 133913373 |  | 12/09/2005 | Dace |
| 3166 | 133913374 |  | 12/09/2005 | Dace |
| 3167 | 133913376 |  | 12/09/2005 | Dace |
| 3168 | 133913375 |  | 12/09/2005 | Dace |
| 3169 | 133913377 |  | 12/09/2005 | Roach |
| 3170 | 133913370 |  | 12/09/2005 | Dace |
| 3171 | 133913372 |  | 12/09/2005 | Dace |
| 3172 | 133913371 |  | 12/09/2005 | Dace |
| 3173 | 133912214 |  | 12/09/2005 | Dace |
| 3174 | 133912213 |  | 12/09/2005 | Dace |
| 3175 | 133912212 |  | 12/09/2005 | Dace |
| 3176 | 133912211 |  | 12/09/2005 | Dace |
| 3177 | 133912210 |  | 12/09/2005 | Dace |
| 3178 | 133912259 |  | 12/09/2005 | Dace |
| 3179 | 133912258 |  | 12/09/2005 | Dace |
| 3180 | 133912256 |  | 12/09/2005 | Roach |
| 3181 | 133912255 |  | 12/09/2005 | Dace |
| 3182 | 133912254 |  | 12/09/2005 | Dace |
| 3183 | 133912253 |  | 12/09/2005 | Roach |
| 3184 | 133912252 |  | 12/09/2005 | Dace |
| 3185 | 133912251 |  | 12/09/2005 | Dace |
| 3186 | 133912250 |  | 12/09/2005 | Dace |
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| 3188 | 133912247 |  | 12/09/2005 | Roach |
| 3189 | 133912246 |  | 12/09/2005 | Dace |
| 3190 | 133912248 |  | 12/09/2005 | Roach |
| 3191 | 133912245 |  | 12/09/2005 | Roach |
| 3192 | 133912244 |  | 12/09/2005 | Dace |
| 3193 | 133912243 |  | 12/09/2005 | Dace |
| 3194 | 133912242 |  | 12/09/2005 | Dace |
| 3195 | 133912241 |  | 12/09/2005 | Dace |
| 3196 | 133912240 |  | 12/09/2005 | Dace |
| 3197 | 133912239 |  | 12/09/2005 | Dace |
| 3198 | 133912238 |  | 12/09/2005 | Dace |
| 3199 | 133912237 |  | 12/09/2005 | Dace |
| 3200 | 133912273 |  | 12/09/2005 | Eel |
| 3201 | 133912277 |  | 12/09/2005 | Eel |
| 3202 | 133912274 |  | 12/09/2005 | Eel |
| 3203 | 133912275 |  | 12/09/2005 | Eel |
| 3204 | 133912236 |  | 12/09/2005 | Dace |
| 3205 | 133912235 |  | 12/09/2005 | Dace |
| 3206 | 133912284 |  | 12/09/2005 | Dace |
| 3207 | 133912283 |  | 12/09/2005 | Dace |
| 3208 | 133912282 |  | 12/09/2005 | Dace |

16.5 Wareham
12.1 Wareham
12.1 Wareham

13 Wareham
15.3 Luckford Lake 4
16.1 ESMS Extra Shock 1
16.3 ESMS Extra Shock 1
16.7 ESMS Extra Shock 1
13.1 ESMS Extra Shock 1
13.8 ESMS Extra Shock 1
14.6 ESMS Extra Shock 1
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13.8 ESMS Extra Shock 1
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19.2 ESMS Extra Shock 1
14.8 ESMS Extra Shock 1
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15.7 ESMS Extra Shock 1
44.8 ESMS Extra Shock 1
32.8 ESMS Extra Shock 1
54.3 ESMS Extra Shock 1
41.5 ESMS Extra Shock 1
13.6 ESMS Extra Shock 1
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15.6 ESMS Extra Shock 1
15.8 ESMS Extra Shock 1
15.9 ESMS Extra Shock 1

| 3209 | 133912281 |
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| 3220 | 133912266 |
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| 3240 | 133911695 |
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| 3247 | 133911688 |
| 3248 | 133911683 |
| 3249 | 133911684 |
| 3250 | 133911685 |
| 3251 | 133911682 |
| 3252 | 133911681 |
| 3253 | 133911730 |
| 3254 | 133911731 |
| 3255 | 133911729 |
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| 3257 | 133911727 |
| 3258 | 133911726 |
| 3259 | 133911725 |
| 3260 | 133911723 |
| 3261 | 133911722 |
| 3262 | 133911721 |
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| $12 / 09 / 2005$ | Dace |
| :--- | :--- |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Eel |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Eel |
| $12 / 09 / 2005$ | Eel |
| $12 / 09 / 2005$ | Eel |
| $12 / 09 / 2005$ | Eel |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Eel |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Dace |
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| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Roach |
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| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Roach |
| 12/09 | Dace |
| 12005 | Dach |
| 12005 |  |

14.4 ESMS Extra Shock 1
15.5 ESMS Extra Shock 1
14.9 ESMS Extra Shock 1
12.6 ESMS Extra Shock 1
44.5 ESMS Extra Shock 1
13.3 ESMS Extra Shock 2

15 ESMS Extra Shock 2
13.7 ESMS Extra Shock 2
13.2 ESMS Extra Shock 2
13.5 ESMS Extra Shock 2
12.7 ESMS Extra Shock 2
16.6 ESMS Extra Shock 2
31.6 ESMS Extra Shock 2
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38.7 ESMS Extra Shock 2
35.5 ESMS Extra Shock 2
12.5 ESMS Extra Shock 3
32.5 ESMS 1
12.9 ESMS 1

15 ESMS 1
19.4 ESMS 1
17.3 ESMS 2
16.2 ESMS 2
16.5 ESMS 2
20.9 ESMS 2
16.9 ESMS 2
14.9 ESMS 2
17.1 ESMS 2
15.7 ESMS 2
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16 ESMS 2
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17.3 ESMS 2
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15.3 ESMS 2
16.5 ESMS 2

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| 3311 | 133911773 |
| 3312 | 133911772 |
| 3313 | 133911771 |
| 3314 | 133911770 |
| 3315 | 133911769 |
| 3316 | 133911768 |
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| 12/09/2005 | Roach | 16 | ESMS 2 |
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| 12/09/2005 | Roach | 15.1 | ESMS 2 |
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| 12/09/2005 | Roach | 14.5 | ESMS 2 |
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| 12/09/2005 | Roach | 21.8 | ESMS 2 |
| 12/09/2005 | Roach | 18.4 | ESMS 2 |
| 12/09/2005 | Roach | 16.8 | ESMS 2 |
| 12/09/2005 | Roach | 16.9 | ESMS 2 |
| 12/09/2005 | Roach | 18.4 | ESMS 2 |
| 12/09/2005 | Dace | 13.6 | ESMS 2 |
| 12/09/2005 | Dace | 13.8 | ESMS 2 |
| 12/09/2005 | Dace | 15.9 | ESMS 2 |
| 12/09/2005 | Dace | 15.4 | ESMS 2 |
| 12/09/2005 | Dace | 14.1 | ESMS 2 |
| 12/09/2005 | Roach | 18.2 | ESMS 2 |
| 12/09/2005 | Roach | 15.6 | ESMS 2 |
| 12/09/2005 | Roach | 14.4 | ESMS 2 |
| 12/09/2005 | Roach | 16.2 | ESMS 2 |
| 12/09/2005 | Roach | 16.8 | ESMS 2 |
| 12/09/2005 | Roach | 15.2 | ESMS 2 |
| 12/09/2005 | Dace | 20.6 | ESMS 2 |
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| 12/09/2005 | Dace | 15.3 | ESMS 2 |
| 12/09/2005 | Dace | 14.9 | ESMS 2 |
| 12/09/2005 | Dace | 14.4 | ESMS 2 |
| 12/09/2005 | Dace | 14.6 | ESMS 2 |
| 12/09/2005 | Dace | 15.3 | ESMS 2 |
| 12/09/2005 | Dace | 17.6 | ESMS 2 |
| 12/09/2005 | Roach | 16.8 | ESMS 2 |
| 12/09/2005 | Roach | 18.3 | ESMS 2 |
| 12/09/2005 | Roach | 16.2 | ESMS 2 |
| 12/09/2005 | Dace | 20.7 | ESMS 2 |
| 12/09/2005 | Dace | 16.6 | ESMS 2 |
| 12/09/2005 | Roach | 14.9 | ESMS 2 |
| 12/09/2005 | Dace | 17.5 | ESMS 2 |
| 12/09/2005 | Roach | 16.4 | ESMS 2 |
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| 12/09/2005 | Roach | 17 | ESMS 2 |
| 12/09/2005 | Roach | 16.6 | ESMS 2 |
| 12/09/2005 | Dace | 15.7 | ESMS 2 |
| 12/09/2005 | Dace | 17.4 | ESMS 2 |
| 12/09/2005 | Roach | 17.4 | ESMS 2 |
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| 12/09/2005 | Dace | 16.6 | ESMS 2 |
| 12/09/2005 | Dace | 15.4 | ESMS 2 |
| 12/09/2005 | Dace | 15.6 | ESMS 2 |
| 12/09/2005 | Dace | 14.3 | ESMS 2 |
| 12/09/2005 | Dace | 15.4 | ESMS 2 |
| 12/09/2005 | Dace | 15 | ESMS 2 |
| 12/09/2005 | Dace | 15 | ESMS 2 |
| 12/09/2005 | Dace | 15.4 | ESMS 2 |


| 3317 | 133911767 | 12/09/2005 | Dace | 16.1 | ESMS 2 |
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| 3331 | 133911803 | 12/09/2005 | Dace | 18.4 | ESMS 2 |
| 3332 | 133911801 | 12/09/2005 | Dace | 17.3 | ESMS 2 |
| 3333 | 133911800 | 12/09/2005 | Dace | 16.2 | ESMS 2 |
| 3334 | 133911799 | 12/09/2005 | Dace | 17.2 | ESMS 2 |
| 3335 | 133911798 | 12/09/2005 | Dace | 15.2 | ESMS 2 |
| 3336 | 133911797 | 12/09/2005 | Dace | 17.5 | ESMS 2 |
| 3337 | 133911796 | 12/09/2005 | Dace | 14.3 | ESMS 2 |
| 3338 | 133911795 | 12/09/2005 | Dace | 13.1 | ESMS 2 |
| 3339 | 133911794 | 12/09/2005 | Dace | 13.5 | ESMS 2 |
| 3340 | 133911793 | 12/09/2005 | Dace | 16.7 | ESMS 2 |
| 3341 | 133911792 | 12/09/2005 | Dace | 15 | ESMS 2 |
| 3342 | 133911791 | 12/09/2005 | Dace | 16.7 | ESMS 2 |
| 3343 | 133911790 | 12/09/2005 | Dace | 14.9 | ESMS 2 |
| 3344 | 133911789 | 12/09/2005 | Dace | 15.2 | ESMS 2 |
| 3345 | 133911788 | 12/09/2005 | Dace | 15.7 | ESMS 2 |
| 3346 | 133911787 | 12/09/2005 | Dace | 15.3 | ESMS 2 |
| 3347 | 133911786 | 12/09/2005 | Dace | 18.3 | ESMS 2 |
| 3348 | 133911785 | 12/09/2005 | Roach | 19.6 | ESMS 2 |
| 3349 | 133911784 | 12/09/2005 | Roach | 15.5 | ESMS 2 |
| 3350 | 133911783 | 12/09/2005 | Roach | 19.3 | ESMS 2 |
| 3351 | 133911782 | 12/09/2005 | Dace | 17.3 | ESMS 2 |
| 3352 | 133911831 | 12/09/2005 | Dace | 15.2 | ESMS 2 |
| 3353 | 133911830 | 12/09/2005 | Dace | 16.2 | ESMS 2 |
| 3354 | 133911829 | 12/09/2005 | Dace | 15 | ESMS 2 |
| 3355 | 133911828 | 12/09/2005 | Dace | 16.8 | ESMS 2 |
| 3356 | 133911827 | 12/09/2005 | Roach | 17.7 | ESMS 2 |
| 3357 | 133911826 | 12/09/2005 | Dace | 15.8 | ESMS 2 |
| 3358 | 133911825 | 12/09/2005 | Roach | 16.5 | ESMS 2 |
| 3359 | 133911824 | 12/09/2005 | Dace | 17 | ESMS 2 |
| 3360 | 133911823 | 12/09/2005 | Dace | 19.8 | ESMS 2 |
| 3361 | 133911822 | 12/09/2005 | Dace | 14.8 | ESMS 2 |
| 3362 | 133911821 | 12/09/2005 | Dace | 16.4 | ESMS 2 |
| 3363 | 133911820 | 12/09/2005 | Dace | 16.5 | ESMS 2 |
| 3364 | 133911819 | 12/09/2005 | Dace | 17.5 | ESMS 2 |
| 3365 | 133911818 | 12/09/2005 | Dace | 16.1 | ESMS 2 |
| 3366 | 133911817 | 12/09/2005 | Dace | 21.1 | ESMS 2 |
| 3367 | 133911816 | 12/09/2005 | Dace | 15.1 | ESMS 2 |
| 3368 | 133911815 | 12/09/2005 | Dace | 14.6 | ESMS 2 |
| 3369 | 133911814 | 12/09/2005 | Dace | 15 | ESMS 2 |
| 3370 | 133911813 | 12/09/2005 | Dace | 16.3 |  |


| 3371 | 133911812 |
| :--- | :--- |
| 3372 | 133911811 |
| 3373 | 133911810 |
| 3374 | 133911809 |
| 3375 | 133911808 |
| 3376 | 133911807 |
| 3377 | 133911855 |
| 3378 | 133911856 |
| 3379 | 133911854 |
| 3380 | 133911852 |
| 3381 | 133911851 |
| 3382 | 133911850 |
| 3383 | 133911849 |
| 3384 | 133911848 |
| 3385 | 133911847 |
| 3386 | 133911846 |
| 3387 | 133911845 |
| 3388 | 133911844 |
| 3389 | 133911841 |
| 3390 | 133911840 |
| 3391 | 133911865 |
| 3392 | 133911842 |
| 3393 | 133911839 |
| 3394 | 133911838 |
| 3395 | 133911837 |
| 3396 | 133911836 |
| 3397 | 133911835 |
| 3398 | 133911834 |
| 3399 | 133911882 |
| 3400 | 133911881 |
| 3401 | 133911833 |
| 3402 | 133911880 |
| 3403 | 133911879 |
| 3404 | 133911878 |
| 3405 | 133911877 |
| 3406 | 133911876 |
| 3407 | 133911874 |
| 3408 | 133911873 |
| 3409 | 133911872 |
| 3410 | 133911871 |
| 3411 | 133911870 |
| 3412 | 133911869 |
| 3413 | 133911867 |
| 3414 | 133911866 |
| 3415 | 133911864 |
| 3416 | 133911863 |
| 3417 | 133911862 |
| 3418 | 133911861 |
| 3419 | 133911860 |
| 3420 | 133911857 |
| 3421 | 133911858 |
| 3422 | 133911832 |
| 3423 | 133911859 |
| 3424 | 133911503 |
|  |  |


| $12 / 09 / 2005$ | Dace |
| :--- | :--- |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 209 / 2005$ | Dace |
| $12 / 209 / 2005$ | Eel |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Roach |
| $12 / 2005$ | Dace |
| $12 / 09 / 2005$ | Roach |
| $12 / 09 / 2005$ | Dace |
| $12 / 09 / 2005$ | Dace |
| 12005 | Dace |
| $12 / 2005$ | Dace |
| $12 / 205$ |  |

15.1 ESMS 2
15.7 ESMS 2
16.3 ESMS 2
14.7 ESMS 2
12.5 ESMS 2
16.4 ESMS 2
14.8 ESMS 2
18.5 ESMS 2
15.3 ESMS 2
16.8 ESMS 2
16.6 ESMS 2
14.7 ESMS 2

16 ESMS 2
16.4 ESMS 2
15.5 ESMS 2
18.8 ESMS 2
14.3 ESMS 2
14.2 ESMS 2
17.3 ESMS 2

15 ESMS 2
15.9 ESMS 2
15.7 ESMS 2
22.4 ESMS 2
16.9 ESMS 2
14.8 ESMS 2
15.7 ESMS 2

17 ESMS 2
19.1 ESMS 2
15.1 ESMS 2
14.5 ESMS 2
16.4 ESMS 2
16.3 ESMS 2
13.2 ESMS 2
16.6 ESMS 2
16.3 ESMS 2
18.2 ESMS 2
16.5 ESMS 2

16 ESMS 2
15.7 ESMS 2

16 ESMS 2
16.1 ESMS 2

15 ESMS 2
16 ESMS 2
20.7 ESMS 2
18.9 ESMS 2
17.3 ESMS 2
14.2 ESMS 2
17.5 ESMS 2
15.6 ESMS 2
15.7 ESMS 2
17.1 ESMS 2
17.2 ESMS 2
14.9 ESMS 2
15.4 ESMS 2

| 3425 | 133911502 | 12/09/2005 | Dace | 17 | ESMS 2 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 3426 | 133911501 | 12/09/2005 | Dace | 15.8 | ESMS 2 |
| 3427 | 133911500 | 12/09/2005 | Dace | 14.6 | ESMS 2 |
| 3428 | 133911499 | 12/09/2005 | Dace | 17.6 | ESMS 2 |
| 3429 | 133911497 | 12/09/2005 | Dace | 16 | ESMS 2 |
| 3430 | 133911496 | 12/09/2005 | Dace | 18.3 | ESMS 2 |
| 3431 | 133911493 | 12/09/2005 | Eel | 32 | ESMS 2 |
| 3432 | 133911494 | 12/09/2005 | Eel | 40 | ESMS 2 |
| 3433 | 133911495 | 12/09/2005 | Eel | 34.8 | ESMS 2 |
| 3434 | 133911491 | 12/09/2005 | Roach | 23.5 | ESMS 2 |
| 3435 | 133911492 | 12/09/2005 | Dace | 19.9 | ESMS 2 |
| 3436 | 133911490 | 12/09/2005 | Dace | 16.5 | ESMS 2 |
| 3437 | 133911489 | 12/09/2005 | Roach | 15.2 | ESMS 2 |
| 3438 | 133911488 | 12/09/2005 | Dace | 16 | ESMS 2 |
| 3439 | 133911481 | 12/09/2005 | Dace | 16 | ESMS 2 |
| 3440 | 133911483 | 12/09/2005 | Dace | 18 | ESMS 2 |
| 3441 | 133911480 | 12/09/2005 | Dace | 14.4 | ESMS 2 |
| 3442 | 133911482 | 12/09/2005 | Dace | 13.7 | ESMS 2 |
| 3443 | 133911486 | 12/09/2005 | Dace | 15.2 | ESMS 2 |
| 3444 | 133911485 | 12/09/2005 | Dace | 17.4 | ESMS 2 |
| 3445 | 133911484 | 12/09/2005 | Dace | 16.4 | ESMS 2 |
| 3446 | 133911479 | 12/09/2005 | Dace | 17.1 | ESMS 2 |
| 3447 | 133911487 | 12/09/2005 | Roach | 17.7 | ESMS 2 |
| 3448 | 133911528 | 12/09/2005 | Dace | 16.2 | ESMS 2 |
| 3449 | 133911526 | 12/09/2005 | Dace | 15.6 | ESMS 2 |
| 3450 | 133911527 | 12/09/2005 | Dace | 19.7 | ESMS 2 |
| 3451 | 133911525 | 12/09/2005 | Dace | 13.7 | ESMS 2 |
| 3452 | 133911524 | 12/09/2005 | Roach | 15.8 | ESMS 2 |
| 3453 | 133911523 | 12/09/2005 | Roach | 17.4 | ESMS 2 |
| 3454 | 133911521 | 12/09/2005 | Dace | 16.5 | ESMS 2 |
| 3455 | 133911522 | 12/09/2005 | Roach | 15.6 | ESMS 2 |
| 3456 | 133911519 | 12/09/2005 | Roach | 14.8 | ESMS 2 |
| 3457 | 133911520 | 12/09/2005 | Dace | 14 | ESMS 2 |
| 3458 | 133911518 | 12/09/2005 | Dace | 15.7 | ESMS 2 |
| 3459 | 133911516 | 12/09/2005 | Dace | 14.7 | ESMS 2 |
| 3460 | 133911517 | 12/09/2005 | Roach | 17.6 | ESMS 2 |
| 3461 | 133911514 | 12/09/2005 | Dace | 19.7 | ESMS 2 |
| 3462 | 133911511 | 12/09/2005 | Roach | 15.7 | ESMS 2 |
| 3463 | 133911513 | 12/09/2005 | Dace | 15.1 | ESMS 2 |
| 3464 | 133911512 | 12/09/2005 | Roach | 15.8 | ESMS 2 |
| 3465 | 133911515 | 12/09/2005 | Dace | 15.3 | ESMS 2 |
| 3466 | 133911509 | 12/09/2005 | Dace | 14.5 | ESMS 2 |
| 3467 | 133911510 | 12/09/2005 | Roach | 16.1 | ESMS 2 |
| 3468 | 133911507 | 12/09/2005 | Roach | 17.5 | ESMS 2 |
| 3469 | 133911505 | 12/09/2005 | Dace | 17.4 | ESMS 2 |
| 3470 | 133911508 | 12/09/2005 | Dace | 14.6 | ESMS 2 |
| 3471 | 133911506 | 12/09/2005 | Roach | 17.2 | ESMS 2 |
| 3472 | 133911554 | 12/09/2005 | Roach | 17.5 | ESMS 2 |
| 3473 | 133911504 | 12/09/2005 | Dace | 17 | ESMS 2 |
| 3474 | 133911553 | 12/09/2005 | Roach | 16.1 | ESMS 2 |
| 3475 | 133911551 | 12/09/2005 | Dace | 15.2 | ESMS 2 |
| 3476 | 133911550 | 12/09/2005 | Dace | 15.8 | ESMS 2 |
| 3477 | 133911549 | 12/09/2005 | Dace | 17.7 | ESMS 2 |
| 3478 | 133911547 | 12/09/2005 | Dace | 17.5 | ESMS 2 |


| 3479 | 133911548 |  | 12/09/2005 | Dace |
| :---: | :---: | :---: | :---: | :---: |
| 3480 | 133911552 |  | 12/09/2005 | Dace |
| 3481 | 133911546 |  | 12/09/2005 | Dace |
| 3482 | 133911545 |  | 12/09/2005 | Dace |
| 3483 | 133911538 |  | 12/09/2005 | Dace |
| 3484 | 133911537 |  | 12/09/2005 | Dace |
| 3485 | 133911539 |  | 12/09/2005 | Dace |
| 3486 | 133911543 |  | 12/09/2005 | Dace |
| 3487 | 133911544 |  | 12/09/2005 | Dace |
| 3488 | 133911542 |  | 12/09/2005 | Roach |
| 3489 | 133911541 |  | 12/09/2005 | Roach |
| 3490 | 133911536 |  | 12/09/2005 | Dace |
| 3491 | 133911535 |  | 12/09/2005 | Dace |
| 3492 | 133911533 |  | 12/09/2005 | Dace |
| 3493 | 133912257 |  | 12/09/2005 | Eel |
| 3494 | 133911593 |  | 12/09/2005 | Eel |
| 3495 | 133911532 |  | 12/09/2005 | Eel |
| 3496 | 133911853 |  | 12/09/2005 | Pike |
| 3497 | 133911875 |  | 14/09/2005 | Pike |
| 3498 | 133911843 |  | 14/09/2005 | Pike |
| 3498 | 133911843 |  | 14/09/2005 | Pike |
| 3498 | 133911843 |  | 14/09/2005 | Pike |
| 3498 | 133911843 |  | 14/09/2005 | Pike |
| 3499 | 133911802 |  | 14/09/2005 | Pike |
| 3500 |  | DC002B-1BC6 | 14/09/2005 | Pike |
| 3501 |  | DC002A-0819 | 14/09/2005 | Pike |
| 3502 | 127178671 |  | 14/09/2005 | Pike |
| 3503 | 127178668 |  | 14/09/2005 | Pike |
| 3504 | 127178669 |  | 14/09/2005 | Pike |
| 3505 | 127178672 |  | 14/09/2005 | Pike |
| 3506 | 127178673 |  | 14/09/2005 | Pike |
| 3507 |  | DC0029-ED9E | 14/09/2005 | Pike |
| 3508 |  | DC0029-EBA5 | 14/09/2005 | Pike |
| 3509 |  | DC002A-F4F5 | 14/09/2005 | Pike |
| 3510 | 133911594 |  | 14/09/2005 | Pike |
| 3511 |  | DC002A-0EEC | 14/09/2005 | Pike |
| 3512 |  | DC002A-03AF | 14/09/2005 | Pike |
| 3513 |  | DC002A-FB61 | 14/09/2005 | Pike |
| 3514 |  | DC002A-02D0 | 14/09/2005 | Pike |
| 3515 |  | DC002B-0298 | 14/09/2005 | Pike |
| 3516 |  | DC002A-0E64 | 14/09/2005 | Pike |
| 3517 |  | DC0029-F169 | 14/09/2005 | Pike |
| 3518 |  | DC0029-EDA7 | 14/09/2005 | Pike |
| 3519 | 127178675 |  | 14/09/2005 | Pike |
| 3520 | 127178676 |  | 14/09/2005 | Pike |
| 3521 | 127178677 |  | 14/09/2005 | Pike |
| 3522 | 127178678 |  | 14/09/2005 | Pike |
| 3523 | 127178679 |  | 14/09/2005 | Pike |
| 3524 | 127178629 |  | 14/09/2005 | Pike |
| 3525 | 127178713 |  | 05/12/2005 | Pike |
| 3526 | 133911597 |  | 05/12/2005 | Pike |
| 3527 | 133911596 |  | 05/12/2005 | Pike |
| 3528 | 127178745 |  | 05/12/2005 | Pike |
| 3529 | 127178730 |  | 05/12/2005 | Pike |

16.9 ESMS 2
16.8 ESMS 2
14.1 ESMS 2
19.5 ESMS 2
15.8 ESMS 2
16.6 ESMS 2

15 ESMS 2
14.5 ESMS 2

17 ESMS 2
17.6 ESMS 2
17.7 ESMS 2
17.7 ESMS 2
15.5 ESMS 2
13.8 ESMS 2
50.2 ESMS 2
37.5 ESMS 2

35 ESMS 2
21.5 ESMS 3
19.4 Railway 1
16.9 Railway 1
16.9 Railway 1
18.4 Railway 1
18.4 Railway 1
19.4 Railway 1
10.4 Railway 1
9.7 Railway 1

18 Railway 2
16.5 Railway 2
16.2 Railway 2
13.4 Railway 2
16.8 Railway 2
9.3 Railway 3
9.2 Flood Relief 2
10.2 Flood Relief 2
25.1 Rushton 1
10.4 Rushton 1
8.7 Rushton 1
9.6 Rushton 1
10.2 Rushton 1
8.8 Rushton 2
8.9 Rushton 2
9.4 Rushton 2
9.7 Rushton 3

34 ESMS 1
36.5 ESMS 1
36.5 ESMS 1
21.2 ESMS 1
21.2 ESMS 1
20.6 ESMS 1
23.3 Railway 1

22 Railway 1
21.2 Railway 2
25.9 Railway 2
24.5 Railway 4

| 3530 | 127178968 | $05 / 12 / 2005$ | Pike | 21.4 | Railway 4 |
| :--- | ---: | ---: | :--- | :--- | :--- |
| 3531 | 113945680 | $05 / 12 / 2003$ | Pike | 39.2 | Holme Bridge 3 |
| 3532 | 127178746 | $05 / 12 / 2005$ | Pike | 25.9 | Railway 1 |
| 3533 | 113945682 |  | $05 / 12 / 2003$ | Pike | 41.8 |
| 3534 | 94698295 | DC0024-D2EC |  | Pike |  |

## Appendix 2 Periods of PIT Detector Operation



Appendix 2 Percentage of each week that each PIT detector was functioning. Note that Luckford was not installed until March 2005.

## Appendix 3 Sample of Standardised PIT data

FlinkaFiskar UKAC CF-recorder ver 1.00

## CONSOLIDATED PIT TAG DATA

Location MH
Run start/end
Gate PIT no.
194698150
194698212
294698212
194698242
194698253
1
24698254
1946982542
1946982573
2946982573

17/12/2003 16:32:05 06/01/2004 10:55:59
Species Time in
25/12/2003 15:28:49 $\quad 25 / 12 / 2003 \quad 15: 28: 49$
18/12/2003 $11: 51: 34 \quad 18 / 12 / 2003 \quad 11: 51: 34$
18/12/2003 11:51:45 $\quad 18 / 12 / 2003 \quad 11: 51: 46$
25/12/2003 12:36:09 $\quad 25 / 12 / 2003 \quad 12: 36: 10$
25/12/2003 $15: 28: 50 \quad 25 / 12 / 2003 \quad 15: 28: 50$
18/12/2003 $11: 51: 29 \quad 18 / 12 / 2003 \quad 11: 51: 30$
18/12/2003 12:11:22 $\quad 18 / 12 / 2003 \quad 12: 11: 22$
18/12/2003 12:11:31 $\quad 18 / 12 / 2003 \quad 12: 11: 31$
18/12/2003 16:06:09 18/12/2003 16:06:10
18/12/2003 16:06:30 18/12/2003 16:10:16

Appendix 3. Second step in the management of PIT data presenting data having been standardised and compressed.

## Appendix 4 Sample of Combined

## Positions Record


Appendix 4. A sample of the Combined Positions Record, which for each fish brings together all detections from any of the
TIRIS PIT readers. It is the main PIT data library, and the starting point for fish centred analysis.

## Appendix 5 Sample of PIT Data Overview


Appendix 5. A sample of the summary of all daily movements measured with PIT telemetry for individual fish created by the Overview macro.

## Appendix 6 Details of Radio Tracked Pike

| Frequency | Name | Start date | End date | No. Fixes | Sex | Size Range* (cm) | Weight Range* <br> (g) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 746 | Alice | 25/05/2000 | 11/09/2002 | 689 | F | 86 | 5216 |
| 788 | Anabelle | 25/02/2004 | 28/07/2005 | 127 | F | 52-65 | 2410 |
| 561/950 | Anakin | 01/03/2004 | 24/06/2005 | 222 | M | 72-73 | 3100-4300 |
| 622 | Bertie | 25/02/2004 | 28/07/2005 | 189 | M | 55 | 2150 |
| 730/930 | Boris | 25/05/2000 | 29/07/2005 | 1222 | M | 71-78 | 3300-5330 |
| 833 | Chica | 09/06/2000 | 05/07/2002 | 1023 | F | 61-72 | 2500-3900 |
| 856 | Chris | 02/02/2003 | 29/10/2004 | 66 | M | 56-65 | 1600-1730 |
| 995 | Elizabeth | 25/02/2004 | 28/07/2005 | 194 | F | 65 | 2550 |
| 522 | Emma | 05/07/2000 | 18/01/2001 | 52 | F | 83 | 4500 |
| 893 | Fiona | 25/07/2000 | 21/11/2001 | 103 | F | 56-63 | 2000 |
| 653 | Frances | 25/02/2004 | 28/07/2005 | 151 | F | 69.5 | 2920 |
| 682 | Fred | 03/03/2004 | 06/01/2005 | 68 | M | 55-61 | 1420-2840 |
| 904 | Gabby | 01/08/2000 | 07/06/2001 | 223 | F | 62-66 | 2700 |
| 778 | Gertie | 25/02/2004 | 28/07/2005 | 150 | F | 58-73 | 1790 |
| 862 | Hannah | 25/08/2000 | 06/10/2003 | 696 | F | 58-87 | 1560-3060 |
| 735 | Helonn | 25/02/2004 | 28/07/2005 | 153 | F | 91 | 7910 |
| 698 | Henry | 16/01/2004 | 01/12/2004 | 50 | M | 64-67 | 2330-2640 |
| 972 | Isaac | 16/10/2000 | 20/04/2003 | 384 | M | 81-87 | 4540-5900 |
| 338 | Janey | 09/03/2005 | 28/07/2005 | 92 | F | 74 | 3120 |
| 959 | Julia | 01/12/2000 | 24/07/2002 | 331 | F | 87-93 | 6900-8160 |
| 947 | Kate | 18/01/2001 | 19/04/2002 | 120 | F | 95 | 8160 |
| 376 | Kathleen | 09/03/2005 | 28/07/2005 | 91 | F | 68 | 2580 |
| 850 | Kin | 16/01/2004 | 28/07/2005 | 146 | F | 101 | 9700-10600 |
| 541 | Lisa | 18/01/2001 | 10/03/2004 | 1386 | F | 77-81 | 3600-4700 |
| 323 | Luke | 09/03/2005 | 28/07/2005 | 93 | M | 50-54 | 1105 |
| 988 | Mark | 18/01/2001 | 28/07/2005 | 737 | M | 64-72 | 2800-3800 |
| 561 | Nicola | 26/02/2001 | 04/07/2001 | 157 | F | 65 | 1900 |
| 581 | Orla | 02/05/2001 | 28/05/2004 | 451 | F | 60-80 | 1800-5200 |
| 530 | Pete | 23/05/2001 | 15/07/2004 | 625 | M | 58-66 | 1700-3060 |
| 514 | Quentin | 23/05/2001 | 14/10/2002 | 238 | M | 69 | 3200 |
| 590 | Rob | 23/05/2001 | 06/07/2001 | 13 | M | 60 | 2000 |
| 505 | Willow | 30/08/2001 | 29/10/2001 | 59 | F | 63 | 1800 |
| 591 | Xena | 06/09/2001 | 22/07/2004 | 413 | F | 54-73 | 1400-3900 |
| 921 | Yoda | 14/11/2001 | 20/04/2003 | 205 | M | 64 | 2250 |
| 756 | Yul | 25/02/2004 | 28/07/2005 | 164 | M | 73 | 4370 |
| 613 | Zac | 25/02/2004 | 28/07/2005 | 195 | M | 59 | 2690 |
| 900 | Zebedee | 28/01/2002 | 04/06/2005 | 224 | M | 55-67 | 1500-3000 |
| 906 | Sian | 31/07/2001 | 26/09/2001 | 74 | F | 52-55 | 1200 |
| 975 | Una | 31/07/2001 | 10/08/2001 | 20 | F | 44 | 1800 |
| 602 | Victoria | 31/07/2001 | 10/02/2003 | 323 | F | 52 | 1700 |
| 065 | Mikey | 09/03/2005 | 15/03/2005 | 7 |  | 26 | 114 |
| 164 |  | 17/03/2005 | 28/04/2005 | 38 | M | 26 | - |
| 188 |  | 18/03/2005 | 20/05/2005 | 61 |  | 26 | 11-178 |
| 404 |  | 18/03/2005 | 24/05/2005 | 60 | M | 28 | - |


| 015 |  | $17 / 03 / 2005$ | $24 / 05 / 2005$ | 66 | M | 25 | 109 |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| 026 |  | $14 / 03 / 2005$ | $21 / 03 / 2005$ | 8 |  | 25 | - |
| $35 / 002$ |  | $14 / 03 / 2005$ | $28 / 04 / 2005$ | 41 | F | 26 | $60-135$ |
| 044 |  | $14 / 03 / 2005$ | $22 / 04 / 2005$ | 30 | M | 27 | 162 |
| 054 |  | $09 / 03 / 2005$ | $07 / 05 / 2005$ | 54 |  | 25 | - |
| 072 |  | $14 / 03 / 2005$ | $06 / 06 / 2005$ | 72 | M | 32 | 284 |
| 084 |  | $14 / 03 / 2005$ | $25 / 03 / 2005$ | 12 | M | 27 | 150 |
| 093 |  | $14 / 03 / 2005$ | $30 / 04 / 2005$ | 42 |  | 25 | - |
| 006 |  | $14 / 03 / 2005$ | $06 / 04 / 2005$ | 20 | M | 26 | 135 |
| 875 | MJ | $17 / 03 / 2004$ | $21 / 04 / 2004$ | 75 |  | 25 | $86-116$ |
|  | Paddy | $17 / 03 / 2004$ | $21 / 04 / 2004$ | 75 |  | 25 | $108-162$ |
| 307 |  | $23 / 06 / 2005$ | $15 / 07 / 2005$ | 12 |  | 25 | - |
| 326 |  | $23 / 06 / 2005$ | $01 / 08 / 2005$ | 19 |  | 25 | - |
| 347 |  | $23 / 06 / 2005$ | $28 / 07 / 2005$ | 17 |  | 26 | $30-142$ |
| 304 | Maddy | $20 / 07 / 2004$ | $06 / 08 / 2004$ | 28 |  | 22 | 97 |
| 854 | Magnus | $20 / 07 / 2004$ | $10 / 08 / 2004$ | 29 |  | 17 | $11-106$ |
| 192 | Mandy | $20 / 07 / 2004$ | $22 / 07 / 2004$ | 3 |  | 14 | 29 |
| 270 | Martha | $20 / 07 / 2004$ | $10 / 08 / 2004$ | 21 |  | 21 | 80 |
| 728 | Maximus | $21 / 07 / 2004$ | $10 / 08 / 2004$ | 29 |  | 9.2 | 8 |
| 793 | Mildred | $20 / 07 / 2004$ | $30 / 07 / 2004$ | 12 |  | 16 | 53 |
| 837 | Morris | $21 / 07 / 2004$ | $10 / 08 / 2004$ | 29 |  | 10 | $8-16$ |
| 764 | Maggie | $17 / 03 / 2004$ | $21 / 04 / 2004$ | 60 |  | 15 | $21-235$ |
|  | Mitch | $17 / 03 / 2004$ | $09 / 04 / 2004$ | 68 |  | 14 | 22 |
|  | Morgan | $17 / 03 / 2004$ | $21 / 04 / 2004$ | 76 |  | 14 | 19 |

Appendix 6 Details of the pike (Esox lucius) radio tracked during the study. Start and end date give the tagging date and the date the fish was last located respectively.

* Size and weight range indicates the size/weight of pike at each capture and thus the known size/weight range during the study.


## Appendix 7 Details of Radio Tracked Dace

| Frequency | Start date | End of tracking | Size (cm) | Weight $(\mathrm{g})$ |
| :---: | :---: | :---: | :---: | :---: |
| 173.801 | $07 / 01 / 2004$ | $17 / 03 / 2004$ | 24.2 | 219 |
| 173.912 | $07 / 01 / 2004$ | $27 / 01 / 2004$ | 24.1 | 206 |
| 173.885 | $07 / 01 / 2004$ | $06 / 02 / 2004$ | 25.7 | 262 |
| 173.615 | $07 / 01 / 2004$ | $12 / 02 / 2004$ | 24.2 | 214 |
| 173.735 | $07 / 01 / 2004$ | $12 / 02 / 2004$ | 23.8 | 230 |
| 173.584 | $07 / 01 / 2004$ | $29 / 01 / 2004$ | 23.1 | 199 |
| 173.765 | $07 / 01 / 2004$ | $12 / 02 / 2004$ | 23.7 | 186 |
| 173.810 | $16 / 03 / 2004$ | $17 / 03 / 2004$ | 19.7 | 112 |
| 173.557 | $16 / 03 / 2004$ | $21 / 03 / 2004$ | 26.0 | 220 |
| 173.713 | $16 / 03 / 2004$ | $19 / 03 / 2004$ | 26.1 | 288 |
| 173.595 | $16 / 03 / 2004$ | $02 / 04 / 2004$ | 26.5 | 284 |
| 173.726 | $16 / 03 / 2004$ | $16 / 03 / 2004$ | 22.9 | 165 |
| 173.625 | $16 / 03 / 2004$ | $19 / 03 / 2004$ | 25.5 | 219 |
| 173.841 | $16 / 03 / 2004$ | $22 / 03 / 2004$ | 21.8 | 174 |
| 173.745 | $16 / 03 / 2004$ | $17 / 03 / 2004$ | 23.2 | 185 |
| 173.894 | $16 / 03 / 2004$ | $01 / 04 / 2004$ | 23.8 | 180 |
| 173.952 | $16 / 03 / 2004$ | $13 / 04 / 2004$ | 20.3 | 120 |
| 173.718 | $16 / 03 / 2004$ | $07 / 04 / 2004$ | 20.8 | 135 |
| 173.761 | $17 / 03 / 2004$ | $15 / 04 / 2004$ | 24.9 | 263 |
| 173.744 | $17 / 03 / 2004$ | $08 / 04 / 2004$ | 27.1 | 333 |
| 173.764 | $17 / 03 / 2004$ | $19 / 03 / 2004$ | 22.4 | 159 |
| 173.616 | $17 / 03 / 2004$ | $01 / 04 / 2004$ | 23.2 | 179 |
| 173.759 | $17 / 03 / 2004$ | $23 / 03 / 2004$ | 23.2 | 196 |
| 173.413 | $17 / 09 / 2004$ | $18 / 10 / 2004$ | 20.7 | - |
| 173.425 | $17 / 09 / 2004$ | $18 / 10 / 2004$ | 20.1 | - |
| 173.485 | $17 / 09 / 2004$ | $02 / 0 / 2004$ | 21.8 | - |
| 173.476 | $17 / 09 / 2004$ | $01 / 12 / 2004$ | 20.4 | - |
| 173.496 | $17 / 09 / 2004$ | $14 / 10 / 2004$ | 22.1 | - |
| 173.504 | $17 / 09 / 2004$ | $01 / 12 / 2004$ | 19.0 | - |
| 173.453 | $17 / 09 / 2004$ | $01 / 12 / 2004$ | 23.0 | - |
| 173.943 | $17 / 09 / 2004$ | $17 / 11 / 2004$ | 19.0 | - |
| 173.436 | $17 / 09 / 2004$ | $01 / 12 / 2004$ | 19.3 | - |
| 173.445 | $17 / 09 / 2004$ | $01 / 12 / 2004$ | 18.9 | - |
| 173.115 | $17 / 03 / 2005$ | $19 / 03 / 2005$ | 22.3 | 181 |
| 173.130 | $17 / 03 / 2005$ | $27 / 03 / 2005$ | 20.6 | 130 |
| 173.142 | $17 / 03 / 2005$ | $21 / 03 / 2005$ | 22.6 | 174 |
| 173.153 | $17 / 03 / 2005$ | $25 / 03 / 2005$ | 22.8 | 194 |
| 173.173 | $17 / 03 / 2005$ | $21 / 03 / 2005$ | 26.4 | - |
| 173.121 | $17 / 03 / 2005$ | $21 / 03 / 2005$ | 22.7 | - |
| 173.106 | $17 / 03 / 2005$ | $21 / 03 / 2005$ | 21.3 | - |

Appendix 7 Details of the dace (Leuciscus leuciscus) radio tracked during the study.
Start and end date give the tagging date and the date the fish was last located respectively. * Size range indicates the size of pike at each capture and thus the known size/weight range during the study.

## Appendix 8 Seasonal Habitat Variability

Abiotic data were single measurements with hand-meters made in each side-channel on the day of each quarterly electric
fishing sampling. Some variability may arise as measurements were made on different days according to when each channel
was sampled. Data presented are the mean over 2003-2005.

| TEMPERATURE $\left({ }^{\circ} \mathrm{C}\right)$ | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 4 | 0.2 | 4 | 4 | 14 | 2 | 16 | 11 | 14 | 2 | 16 | 12 | 4 | 1 | 5 | 4 |
| Millstream | 9 | 0.1 | 10 | 9 | 18 | 1 | 20 | 17 | 15 | 1 | 17 | 14 | 7 | 1 | 10 | 5 |
| Rushton | x | x | x | x | 14 | 2 | 20 | 12 | 14 | 2 | 16 | 11 | 7 | 1 | 9 | 6 |
| Luckford | 9 | 0.1 | 9 | 9 | 14 | 2 | 19 | 12 | 8 | 4 | 13 | 3 | 7 | 2 | 9 | 5 |
| Goldsacs | 9 | 0.1 | 9 | 9 | 14 | 1 | 15 | 13 | 14 | 1 | 15 | 14 | 7 | 1 | 9 | 6 |
| Holme Bridge | 2 | 0.1 | 2 | 1 | 15 | 1 | 18 | 14 | 14 | 1 | 16 | 12 | 7 | 1 | 8 | 5 |
| Flood Relief | 7 | 0.6 | 7 | 6 | 18 | 4 | 24 | 13 | 15 | 2 | 17 | 7 | 7 | 1 | 9 | 5 |


| DISSOLVEDOXYGEN $(\mathrm{mg} / \mathrm{l})$ | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 2 | 0 | 2 | 2 | 6 | 7 | 31 | 1 | 5 | 2 | 8 | 2 | 6 | 1 | 7 | 5 |
| Millstream | 13 | 4 | 17 | 9 | 11 | 3 | 14 | 8 | 13 | 2 | 16 | 10 | 10 | 1 | 14 | 8 |
| Rushton | x | x | x | x | 5 | 2 | 9 | 1 | 8 | 3 | 12 | 5 | 5 | 1 | 6 | 4 |
| Luckford | 2 | 0.8 | 18 | 15 | 4 | 2 | 10 | 2 | 6 | 3 | 12 | 4 | 9 | 2 | 12 | 8 |
| Goldsacs | 5 | 3 | 9 | 2 | 8 | 2 | 10 | 1 | 10 | 2 | 12 | 7 | 10 | 0 | 10 | 10 |
| Holme Bridge | 4 | 3 | 8 | 1 | 5 | 4 | 11 | 1 | 4 | 2 | 6 | 1 | 5 | 1 | 7 | 5 |
| Flood Relicf | 1 | 0.2 | 2 | 1 | 8 | 4 | 14 | 4 | 7 | 4 | 16 | 3 | 6 | 2 | 8 | 4 |


| DEPTH (cm) | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 36 | 17 | 82 | 7 | 29 | 15 | 70 | 5 | 27 | 12 | 58 | 5 | 33 | 13 | 63 | 9 |
| Millstream | 43 | 22 | 100 | 4 | 50 | 23 | 120 | 18 | 37 | 22 | 99 | 4 | 31 | 17 | 78 | 3 |
| Rushton | 50 | 18 | 83 | 16 | 53 | 25 | 115 | 15 | 47 | 20 | 100 | 2 | 50 | 19 | 100 | 6 |
| Luckford | 59 | 26 | 130 | 10 | 53 | 28 | 120 | 8 | 50 | 26 | 119 | 12 | 50 | 26 | 119 | 12 |
| Goldsacs | 30 | 19 | 100 | 8 | 22 | 12 | 55 | 2 | 26 | 17 | 79 | 1 | 37 | 19 | 95 | 7 |
| Holme Bridge | 49 | 29 | 100 | 10 | 52 | 25 | 105 | 15 | 43 | 27 | 100 | 0 | 49 | 24 | 100 | 10 |
| Flood Relief | 39 | 22 | 90 | 1 | 45 | 30 | 120 | 2 | 38 | 22 | 100 | 5 | 40 | 21 | 79 | 5 |
| WIDTH (m) |  |  | rch |  |  |  | ne |  |  | Sept | mber |  |  | Dec | mber |  |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 4 | 2 | 7 | 2 | 3 | 1 | 5 | 1 | 3 | 1 | 4 | 2 | 3 | 1 | 4 | 2 |
| Millstream | 5 | 1 | 8 | 4 | 6 | 1 | 8 | 5 | 5 | 1 | 7 | 3 | 7 | 3 | 14 | 4 |
| Rushton | 3 | 1 | 4 | 2 | 3 | 1 | 4 | 2 | 3 | 1 | 4 | 2 | 3 | 1 | 4 | 1 |
| Luckford | 5 | 1 | 8 | 2 | 6 | 1 | 8 | 2 | 5 | 1 | 9 | 3 | 5 | 1 | 8 | 4 |
| Goldsacs | 2 | 1 | 4 | 1 | 2 | 1 | 4 | 1 | 2 | 1 | 4 | 1 | 2 | 1 | 4 | 1 |
| Holme Bridge | 5 | 3 | 13 | 2 | 4 | 2 | 10 | 2 | 4 | 2 | 10 | 0 | 4 | 2 | 9 | 1 |
| Flood Relief | 3 | 1 | 6 | 1 | 4 | 1 | 6 | 2 | 3 | 1 | 6 | 1 | 3 | 1 | 6 | 2 |


| $\begin{aligned} & \hline \text { TURBULENT } \\ & (\%) \end{aligned}$ | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Millstream | 1 | 2 | 5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Rushton | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Luckford | 1 | 2 | 5 | 0 | 1 | 2 | 5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Goldsacs | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 5 | 0 |
| Holme Bridge | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Flood Relief | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |  | 0 | 0 | 0 |


| GLIDE (\%) | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 10 | 28 | 80 | 0 | 1 | 2 | 5 | 0 | 0 | 0 | 0 | 0 | 8 | 7 | 20 | 0 |
| Millstream | 98 | 5 | 100 | 80 | 93 | 10 | 100 | 60 | 94 | 6 | 100 | 80 | 99 | 4 | 100 | 80 |
| Rushton | 4 | 8 | 20 | 0 | 1 | 4 | 10 | 0 | 0.9 | 2 | 5 | 0 | 2 | 3 | 10 | 0 |
| Luckford | 7 | 14 | 50 | 0 | 1 | 2 | 5 | 0 | 14 | 21 | 60 | 0 | 5 | 0 | 5 | 5 |
| Goldsacs | 100 | 0 | 100 | 100 | 98 | 4 | 100 | 90 | 68 | 32 | 100 | 10 | 99 | 4 | 100 | 90 |
| Holme Bridge | 6 | 20 | 80 | 0 | 54 | 49 | 100 | 0 | 0 | 0 | 0 | 0 | 3 | 4 | 10 | 0 |
| Flood Relief | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 10 | 24 | 80 | 0 | 5 | 15 | 60 | 0 |


| SLACK (\%) | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 95 | 20 | 100 | 20 | 100 | 1 | 100 | 95 | 100 | 0 | 100 | 100 | 95 | 7 | 100 | 80 |
| Millstream | 2 | 5 | 20 | 0 | 7 | 10 | 40 | 0 | 6 | 6 | 20 | 0 | 3 | 7 | 20 | 0 |
| Rushton | 98 | 5 | 100 | 80 | 100 | 2 | 100 | 90 | 99 | 2 | 100 | 95 | 99 | 3 | 100 | 90 |
| Luckford | 95 | 12 | 100 | 50 | 100 | 1 | 100 | 95 | 91 | 18 | 100 | 40 | 97 | 2 | 100 | 95 |
| Goldsacs | 100 | 0 | 100 | 100 | 3 | 5 | 10 | 0 | 32 | 32 | 90 | 0 | 4 | 5 | 10 | 0 |
| Holme Bridge | 94 | 20 | 100 | 20 | 64 | 47 | 100 | 0 | 100 | 0 | 100 | 100 | 98 | 4 | 100 | 90 |
| Flood Relief | 100 | 0 | 100 | 100 | 100 | 0 | 100 | 100 | 93 | 20 | 100 | 20 | 96 | 12 | 100 | 40 |
|  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |
| CLAY (\%) |  |  | arch |  |  |  | ne |  |  | Septe | mber |  |  | Dece | nber |  |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 0 | 0 | 0 | 0 | 17 | 24 | 50 | 0 | 19 | 20 | 80 | 0 | 3 | 4 | 10 | 0 |
| Millstream | 1 | 3 | 10 | 0 | 18 | 33 | 80 | 0 | 8 | 9 | 30 | 0 | 5 | 6 | 30 | 0 |
| Rushton | 0 | 0 | 0 | 0 | 28 | 32 | 90 | 0 | 23 | 31 | 80 | 0 | 27 | 22 | 70 | 0 |
| Luckford | 0 | 0 | 0 | 0 | 30 | 45 | 100 | 0 | 20 | 25 | 100 | 0 | 14 | 22 | 80 | 0 |
| Goldsacs | 4 | 7 | 20 | 0 | 0 | 0 | 0 | 0 | 6 | 10 | 40 | 0 | 3 | 5 | 15 | 0 |
| Holme Bridge | 0 | 0 | 0 | 0 | 5 | 12 | 40 | 0 | 13 | 12 | 40 | 0 | 7 | 16 | 50 | 0 |
| Flood Relief | 0 | 0 | 0 | 0 | 2 | 5 | 20 | 0 | 30 | 37 | 90 | 0 | 8 | 10 | 30 | 0 |


| SILT (\%) | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 33 | 45 | 100 | 0 | 83 | 24 | 100 | 50 | 78 | 21 | 100 | 20 | 91 | 18 | 100 | 10 |
| Millstream | 6 | 4 | 10 | 0 | 4 | 5 | 10 | 0 | 99 | 6 | 20 | 0 | 2 | 3 | 10 | 0 |
| Rushton | 12 | 26 | 80 | 0 | 54 | 33 | 100 | 10 | 67 | 39 | 100 | 0 | 61 | 26 | 100 | 10 |
| Luckford | 73 | 29 | 100 | 20 | 58 | 44 | 100 | 0 | 61 | 29 | 100 | 0 | 75 | 30 | 100 | 10 |
| Goldsacs | 7 | 5 | 10 | 0 | 15 | 22 | 90 | 0 | 19 | 13 | 40 | 10 | 14 | 13 | 40 | 0 |
| Holme Bridge | 64 | 41 | 100 | 5 | 72 | 37 | 100 | 0 | 77 | 29 | 100 | 10 | 78 | 33 | 100 | 0 |
| Flood Relief | 35 | 47 | 100 | 0 | 80 | 29 | 100 | 10 | 40 | 33 | 100 | 0 | 79 | 23 | 100 | 20 |
| SAND (\%) |  |  |  |  |  |  | ne |  |  | Sept | mber |  |  | Dece | nber |  |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 33 | 47 | 100 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Millstream | 22 | 21 | 50 | 0 | 14 | 16 | 60 | 0 | 13 | 18 | 80 | 0 | 12 | 10 | 50 | 0 |
| Rushton | 5 | 20 | 100 | 0 | 9 | 15 | 50 | 0 | 0 | 0 | 0 | 0 | , | 0 | 0 | 0 |
| Luckford | 31 | 43 | 100 | 0 | 1 | 3 | 15 | 0 | 2 | 4 | 10 | 0 | 2 | 5 | 20 | 0 |
| Goldsacs | 57 | 35 | 90 | 10 | 78 | 23 | 100 | 10 | 73 | 23 | 90 | 20 | 76 | 14 | 100 | 50 |
| Holme Bridge | 35 | 43 | 100 | 0 | 17 | 38 | 100 | 0 | 0 | 0 | 0 | 0 | 3 | 7 | 30 | 0 |
| Flood Relief | 33 | 48 | 100 | 0 | 3 | 6 | 20 | 0 | 2 | 8 | 40 | 0 | 0 | 1 | 5 | 0 |


| GRAVEL (\%) | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 0 | 1 | 5 | 0 | 0.2 | 1 | 5 | 0 | 3 | 4 | 10 | 0 | 2 | 5 | 20 | 0 |
| Millstream | 57 | 13 | 80 | 45 | 64 | 35 | 100 | 5 | 62 | 27 | 90 | 0 | 77 | 1 | 90 | 40 |
| Rushton | 33 | 47 | 100 | 0 | 9 | 20 | 90 | 0 | 1 | 21 | 70 | 0 | 13 | 25 | 85 | 0 |
| Luckford | 9 | 18 | 70 | 0 | 12 | 24 | 75 | 0 | 17 | 25 | 80 | 0 | 10 | 22 | 80 | 0 |
| Goldsacs | 32 | 42 | 90 | 0 | 6 | 9 | 30 | 0 | 2 | 4 | 10 | 0 | 6 | 9 | 35 | 0 |
| Holme Bridge | 5 | 15 | 70 | 0 | 6 | 11 | 40 | 0 | 10 | 24 | 80 | 0 | 13 | 26 | 80 | 0 |
| Flood Relief | 0 | 1 | 5 | 0 | 15 | 25 | 80 | 0 | 20 | 27 | 80 | 0 | 10 | 19 | 60 | 0 |


| COBBLES (\%) | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 2 | 10 | 0 |
| Millstream | 16 | 23 | 50 | 0 | 0 | 0 | 0 | 0 | 1 | 2 | 5 | 0 | 4 | 4 | 10 | 0 |
| Rushton | 29 | 46 | 100 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 1 | 5 | 0 |
| Luckford | 3 | 1 | 40 | 0 | 0 | 1 | 5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Goldsacs | 0 | 0 | 0 | 0 | 0 | 1 | 5 | 0 | 0 | 1 | 5 | 0 | 1 | 2 | 5 | 0 |
| Holme Bridge | 0 | 2 | 10 | 0 | 1 | 2 | 5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Flood Relief | 0 | 1 | 5 | 0 | 0 | 1 | 5 | 0 | 1 | 3 | 15 | 0 | 3 | 7 | 30 | 0 |


| $\begin{aligned} & \text { TREE ROOTS } \\ & (\%) \end{aligned}$ | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 1 | 5 | 20 | 0 | 2 | 4 | 10 | 0 | 6 | 9 | 40 | 0 | 7 | 15 | 65 | 0 |
| Millstream | 6 | 12 | 40 | 0 | 1 | 3 | 10 | 0 | 4 | 6 | 25 | 0 | 5 | 8 | 30 | 0 |
| Rushton | 0 | 0 | 0 | 0 | 0 | 1 | 5 | 0 | 3 | 5 | 20 | 0 | 2 | 4 | 10 | 0 |
| Luckford | 21 | 29 | 80 | 0 | 6 | 6 | 20 | 0 | 11 | 14 | 50 | 0 | 9 | 14 | 50 | 0 |
| Goldsacs | 9 | 16 | 50 | 0 | 2 | 6 | 30 | 0 | 14 | 23 | 90 | 0 | 3 | 4 | 10 | 0 |
| Holme Bridge | 7 | 17 | 70 | 0 | 2 | 4 | 15 | 0 | 4 | 4 | 10 | 0 | 7 | 11 | 50 | 0 |
| Flood Relief | 3 | 10 | 50 | 0 | 1 | 2 | 5 | 0 | 2 | 4 | 20 | 0 | 0 | 1 | 5 | 0 |


| BRANCHES/ LOGS (\%) | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 30 | 43 | 100 | 0 | 0 | 2 | 10 | 0 | 4 | 7 | 30 | 0 | 3 | 8 | 40 | 0 |
| Millstream | 3 | 6 | 20 | 0 | 1 | 4 | 20 | 0 | 1 | 3 | 10 | 0 | 0 | 2 | 5 | 0 |
| Rushton | 12 | 21 | 70 | 0 | 2 | 10 | 50 | 0 | 1 | 2 | 10 | 0 | 2 | 5 | 20 | 0 |
| Luckford | 20 | 25 | 80 | 0 | 22 | 23 | 80 | 0 | 37 | 28 | 90 | 0 | 39 | 62 | 300 | 0 |
| Goldsacs | 2 | 5 | 20 | 0 | 2 | 4 | 10 | 0 | 0 | 0 | 0 | 0 | 2 | 4 | 15 | 0 |
| Holme Bridge | 17 | 28 | 80 | 0 | 10 | 17 | 60 | 0 | 15 | 19 | 60 | 0 | 30 | 33 | 100 | 0 |
| Flood Relief | 6 | 21 | 100 | 0 | 2 | 4 | 20 | 0 | 1 | 2 | 10 | 0 | 3 | 7 | 30 | 0 |


| SUBMERGED SPARSE (\%) | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 1 | 5 | 20 | 0 | 0 | 0 | 0 | 0 | 9 | 11 | 25 | 0 | 0 | 0 | 0 | 0 |
| Millstream | 5 | 11 | 40 | 0 | 2 | 5 | 20 | 0 | 2 | 4 | 10 | 0 | 4 | 6 | 25 | 0 |
| Rushton | 0 | 0 | 0 | 0 | 13 | 18 | 70 | 0 | 4 | 11 | 40 | 0 | 7 | 7 | 20 | 0 |
| Luckford | 0 | 0 | 0 | 0 | 0 | 2 | 10 | 0 | 0 | 1 | 5 | 0 | 0 | 0 | 0 | 0 |
| Goldsacs | 0 | 0 | 0 | 0 | 0 | 2 | 10 | 0 | 0 | 1 | 5 | 0 | 1 | 2 | 5 | 0 |
| Holme Bridge | 5 | 17 | 70 | 0 | 3 | 6 | 25 | 0 | 1 | 4 | 20 | 0 | 3 | 6 | 20 | 0 |
| Flood Relief | 3 | 10 | 50 | 0 | 3 | 7 | 30 | 0 | 12 | 32 | 95 | 0 | 1 | 2 | 5 | 0 |


| EMERGENT | March |  |  |  | June |  |  |  |  |  |  |  |  |  | September |  |  |  |  | December |  |  |  |
| :--- | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | ---: | :---: | :---: | :---: | :---: | :---: | :---: |
| DENSE (\%) | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |  |  |  |  |  |  |  |
| Railway | 29 | 44 | 100 | 0 | 86 | 19 | 100 | 20 | 86 | 27 | 100 | 0 | 64 | 32 | 100 | 0 |  |  |  |  |  |  |  |
| Millstream | 0 | 0 | 0 | 0 | 2 | 5 | 20 | 0 | 5 | 9 | 40 | 0 | 0 | 1 | 5 | 0 |  |  |  |  |  |  |  |
| Rushton | 12 | 21 | 70 | 0 | 21 | 30 | 80 | 0 | 46 | 36 | 100 | 0 | 33 | 26 | 80 | 0 |  |  |  |  |  |  |  |
| Luckford | 0 | 2 | 10 | 0 | 1 | 4 | 15 | 0 | 4 | 10 | 30 | 0 | 0.6 | 2 | 10 | 0 |  |  |  |  |  |  |  |
| Goldsacs | 0 | 1 | 3 | 0 | 11 | 19 | 50 | 0 | 16 | 23 | 80 | 0 | 1 | 3 | 15 | 0 |  |  |  |  |  |  |  |
| Holme Bridge | 1 | 2 | 10 | 0 | 6 | 11 | 40 | 0 | 9 | 16 | 60 | 0 | 4 | 10 | 35 | 0 |  |  |  |  |  |  |  |
| Flood Relief | 6 | 21 | 100 | 0 | 1 | 3 | 10 | 0 | 14 | 32 | 100 | 0 | 2 | 4 | 10 | 0 |  |  |  |  |  |  |  |


| SUBMERGED DENSE (\%) | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 0 | 0 | 0 | 0 | 0 | 0 | 2 | 0 | 0 | 0 | 0 | 0 | 1 | 3 | 15 | 0 |
| Millstream | 16 | 29 | 80 | 0 | 36 | 37 | 100 | 0 | 2 | 4 | 15 | 0 | 13 | 23 | 70 | 0 |
| Rushton | 8 | 23 | 100 | 0 | 7 | 15 | 60 | 0 | 21 | 23 | 80 | 0 | 14 | 18 | 60 | 0 |
| Luckford | 1 | 4 | 20 | 0 | 0 | 0 | 0 | 0 | 0 |  | 0 | 0 | 0 | 0 | 0 | 0 |
| Goldsacs | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.4 | 1 | 5 | 0 | 0 | 0 | 0 | 0 |
| Holme Bridge | 12 | 25 | 80 | 0 | 3 | 9 | 40 | 0 | 12 | 26 | 90 | 0 | 0 | 0 | 0 | 0 |
| Flood Relief | 6 | 22 | 100 | 0 | 35 | 37 | 90 | 0 | 20 | 35 | 100 | 0 | 17 | 33 | 100 | 0 |


| $\begin{aligned} & \text { OVERHUNG } \\ & \text { TREES (\%) } \end{aligned}$ | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 0 | 0 | 0 | 0 | 0 | 0 | 2 | 0 | 0 | 0 | 0 | 0 | 1 | 3 | 15 | 0 |
| Millstream | 6 | 12 | 50 | 0 | 6 | 12 | 50 | 0 | 17 | 16 | 40 | 0 | 10 | 11 | 40 | 0 |
| Rushton | 11 | 24 | 100 | 0 | 25 | 30 | 100 | 0 | 28 | 30 | 100 | 0 | 21 | 24 | 90 | 0 |
| Luckford | 32 | 37 | 100 | 0 | 84 | 24 | 100 | 40 | 90 | 22 | 100 | 0 | 37 | 31 | 90 | 0 |
| Goldsacs | 13 | 24 | 80 | 0 | 48 | 37 | 100 | 0 | 55 | 40 | 100 | 0 | 39 | 28 | 90 | 5 |
| Holme Bridge | 49 | 40 | 100 | 0 | 77 | 30 | 100 | 0 | 75 | 38 | 100 | 0 | 65 | 37 | 100 | 0 |
| Flood Relief | 13 | 24 | 100 | 0 | 44 | 41 | 100 | 0 | 36 | 42 | 100 | 0 | 23 |  | 70 | 0 |


| EMERGENT SPARSE (\%) | March |  |  |  | June |  |  |  | September |  |  |  | December |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 4 | 11 | 50 | 0 | 0 | 4 | 15 | 0 | 4 | 13 | 60 | 0 | 9 | 19 | 80 | 0 |
| Millstream | 0 | 0 | 0 | 0 | 4 | 5 | 20 | 0 | 2 | 4 | 10 | 0 | 5 | 11 | 40 | 0 |
| Rushton | 3 | 10 | 50 | 0 | 30 | 30 | 90 | 0 | 7 | 11 | 40 | 0 | 12 | 9 | 40 | 0 |
| Luckford | 7 | 15 | 50 | 0 | 0 | 1 | 5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Goldsacs | 3 | 7 | 30 | 0 | 2 | 5 | 20 | 0 | 3 | 7 | 30 | 0 | 1 | 2 | 5 | 0 |
| Holme Bridge | 14 | 29 | 95 | 0 | 5 | 7 | 25 | 0 | 3 | 5 | 20 | 0 | 3 | 5 | 15 | 0 |
| Flood Relief | 7 | 14 | 50 | 0 | 5 | 6 | 20 | 0 | 6 | 8 | 30 | 0 | 4 | 8 | 30 | 0 |

## Appendix 9 Annual Habitat Variation

Abiotic data were single measurements with hand-meters made in each side-channel on the day of each quarterly electric fishing sampling. Some variability may arise as measurements were made on different days according to when each channel was sampled. Data presented are the mean of all seasons in a given year.


| TURBULENT (\%) | $\begin{aligned} & 2003 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | 2004 <br> Mean | SD | Max | Min | $\begin{aligned} & 2005 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Railway | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Millstream | 0.6 | 2 | 5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Rushton | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Luckford | 0 | 0 | 0 | 0 | 1 | 2 | 5 | 0 | 0 | 0 | 0 | 0 |
| Goldsacs | 1 | 2 | 5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Holme Bridge | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| Flood Relief | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| GLIDE (\%) | 2003 <br> Mean | SD | Max | Min | $\begin{aligned} & 2004 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | $\begin{aligned} & 2005 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min |
|  |  |  |  |  |  |  |  |  |  |  |  |  |
| Railway | 11 | 20 | 80 | 0 | 5 | 0 | 5 | 5 | 0.6 | 2 | 5 | 0 |
| Millstream | 98 | 4 | 100 | 80 | 93 | 10 | 100 | 60 | 96 | 6 | 100 | 80 |
| Rushton | 1 | 2 | 5 | 0 | 30 | 6 | 20 | 0 | 1 | 2 | 5 | 0 |
| Luckford | 16 | 20 | 60 | 0 | 6 | 14 | 50 | 0 | 3 | 4 | 20 | 0 |
| Goldsacs | 89 | 22 | 100 | 10 | 89 | 26 | 100 | 10 | 92 | 18 | 100 | 20 |
| Holme Bridge | 6 | 20 | 80 | 0 | 12 | 27 | 80 | 0 | 45 | 50 | 100 | 0 |
| Flood Relief | 13 | 27 | 80 | 0 | 2 | 3 | 5 | 0 | 3 | 4 | 10 | 0 |
| SLACK (\%) | $\begin{aligned} & \hline 2003 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | $\begin{aligned} & 2004 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | 2005 <br> Mean | SD | Max | Min |
|  |  |  |  |  |  |  |  |  |  |  |  |  |
| Railway | 95 | 15 | 100 | 20 | 99 | 2 | 100 | 95 | 100 | 1 | 100 | 95 |
| Millstream | 3 | 5 | 20 | 0 | 7 | 10 | 40 | 0 | 5 | 6 | 20 | 0 |
| Rushton | 100 | 2 | 100 | 95 | 98 | 4 | 100 | 80 | 100 | 1 | 100 | 95 |
| Luckford | 92 | 16 | 100 | 40 | 97 | 10 | 100 | 50 | 98 | 4 | 100 | 80 |
| Goldsacs | 66 | 41 | 100 | 10 | 15 | 28 | 90 | 0 | 15 | 23 | 80 | 0 |
| Holme Bridge | 97 | 14 | 100 | 20 | 92 | 23 | 100 | 20 | 78 | 42 | 100 | 0 |
| Flood Relief | 94 | 20 | 100 | 20 | 100 | 2 | 100 | 95 | 99 | 2 | 100 | 90 |
| $\begin{aligned} & \text { COBBLES } \\ & \text { (\%) } \end{aligned}$ | $2003$ <br> Mean | SD | Max | Min | $\begin{aligned} & \hline 2004 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | $\begin{aligned} & 2005 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min |
|  |  |  |  |  |  |  |  |  |  |  |  |  |
| Railway | 0 | 0 | 0 | 0 | 0 | 1 | 5 | 0 | 0 | 2 | 10 | 0 |
| Millstream | 2 | 3 | 10 | 0 | 18 | 30 | 80 | 0 | 0 | 1 | 5 | 0 |
| Rushton | 0 | 1 | 5 | 0 | 0 | 1 | 5 | 0 | 0 | 0 | 0 | 0 |
| Luckford | 0 | 0 | 0 | 0 | 1 | 3 | 15 | 0 | 0 | 0 | 0 | 0 |
| Goldsacs | 0 | 1 | 5 | 0 | 3 | 7 | 30 | 0 | 0 | 1 | 5 | 0 |
| Holme Bridge | 0 | 0 | 0 | 0 | 2 | 6 | 25 | 0 | 0 | 2 | 10 | 0 |
| Flood Relief | 0 | 0 | 0 | 0 | 2 | 5 | 20 | 0 | 2 | 6 | 30 | 0 |
| GRAVEL (\%) | $\begin{aligned} & 2003 \\ & \text { Mean } \\ & \hline \end{aligned}$ | SD | Max | Min | $\begin{aligned} & 2004 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | $\begin{aligned} & \hline 2005 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min |
|  |  |  |  |  |  |  |  |  |  |  |  |  |
| Railway | 1 | 3 | 10 | 0 | 1 | 4 | 20 | 0 | 1 | 3 | 10 | 0 |
| Millstream | 73 | 18 | 100 | 45 | 56 | 28 | 90 | 0 | 60 | 30 | 90 | 5 |
| Rushton | 32 | 42 | 100 | 0 | 9 | 21 | 70 | 0 | 6 | 12 | 50 | 0 |
| Luckford | 17 | 27 | 80 | 0 | 7 | 18 | 80 | 0 | 11 | 21 | 70 | 0 |
| Goldsacs | 2 | 7 | 35 | 0 | 24 | 30 | 80 | 0 | 5 | 7 | 20 | 0 |
| Holme Bridge | 5 | 14 | 70 | 0 | 10 | 21 | 80 | 0 | 10 | 24 | 80 | 0 |
| Flood Relief | 15 | 27 | 80 | 0 | 10 | 20 | 80 | 0 | 10 | 16 | 50 | 0 |


| SAND (\%) | $\begin{aligned} & 2003 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | $\begin{aligned} & 2004 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | $\begin{aligned} & 2005 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Railway | 0 | 1 | 5 | 0 | 25 | 44 | 100 | 0 | 0 | 0 | 0 | 0 |
| Millstream | 19 | 19 | 50 | 0 | 14 | 16 | 60 | 0 | 13 | 15 | 80 | 0 |
| Rushton | 4 | 18 | 100 | 0 | 24 | 39 | 100 | 0 | 24 | 39 | 100 | 0 |
| Luckford | 2 | 4 | 10 | 0 | 26 | 42 | 100 | 0 | 0.9 | 3 | 15 | 0 |
| Goldsacs | 92 | 8 | 100 | 65 | 49 | 24 | 80 | 10 | 74 | 17 | 90 | 10 |
| Holme Bridge | 15 | 33 | 100 | 0 | 23 | 41 | 100 | 0 | 3 | 5 | 20 | 0 |
| Flood Relief | 3 | 9 | 40 | 0 | 23 | 41 | 100 | 0 | 0 | 0 | 0 | 0 |
| CLAY (\%) | $\begin{aligned} & 2003 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | $\begin{aligned} & 2004 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | $\begin{aligned} & 2005 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min |
|  |  |  |  |  |  |  |  |  |  |  |  |  |
| Railway | 15 | 21 | 50 | 0 | 2 | 6 | 30 | 0 | 12 | 20 | 80 | 0 |
| Millstream | 3 | 4 | 10 | 0 | 2 | 5 | 25 | 0 | 19 | 28 | 80 | 0 |
| Rushton | 32 | 34 | 90 | 0 | 13 | 21 | 80 | 0 | 14 | 19 | 50 | 0 |
| Luckford | 13 | 24 | 100 | 0 | 7 | 20 | 80 | 0 | 28 | 38 | 100 | 0 |
| Goldsacs | 0 | 0 | 0 | 0 | 4 | 9 | 40 | 0 | 6 | 7 | 20 | 0 |
| Holme Bridge | 8 | 15 | 50 | 0 | 7 | 12 | 40 | 0 | 4 | 11 | 40 | 0 |
| Flood Relief | 5 | 9 | 40 | 0 | 8 | 24 | 80 | 0 | 16 | 29 | 90 | 0 |
| SILT (\%) | $\begin{aligned} & 2003 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | $\begin{aligned} & \hline 2004 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min | $\begin{aligned} & \hline 2005 \\ & \text { Mean } \end{aligned}$ | SD | Max | Min |
|  |  |  |  |  |  |  |  |  |  |  |  |  |
| Railway | 83 | 21 | 100 | 50 | 72 | 43 | 100 | 0 | 59 | 42 | 100 | 0 |
| Millstream | 3 | 4 | 10 | 0 | 4 | 5 | 20 | 0 | 8 | 5 | 20 | 0 |
| Rushton | 32 | 34 | 100 | 0 | 54 | 43 | 100 | 0 | 50 | 36 | 100 | 0 |
| Luckford | 68 | 32 | 100 | 0 | 60 | 43 | 100 | 0 | 60 | 38 | 100 | 0 |
| Goldsacs | 6 | 6 | 20 | 0 | 23 | 11 | 40 | 10 | 15 | 17 | 90 | 0 |
| Holme Bridge | 73 | 35 | 100 | 0 | 58 | 43 | 100 | 0 | 83 | 29 | 100 | 10 |
| Flood Relief | 83 | 29 | 100 | 10 | 53 | 43 | 100 | 0 | 47 | 39 | 100 | 0 |
| BRANCHES/ LOGS (\%) | 2003 <br> Mean | SD | Max | Min | 2004 |  | Max | Min | $\begin{aligned} & 2005 \\ & \text { Mean } \\ & \hline \end{aligned}$ | SD | Max | Min |
|  |  |  |  |  | Mean | SD |  |  |  |  |  |  |
| Railway | 0.9 | 3 | 10 | 0 | 24 | 39 | 100 | 0 | 3 | 6 | 30 | 0 |
| Millstream | 0.6 | 2 | 10 | 0 | 1 | 4 | 20 | 0 | 2 | 5 | 20 | 0 |
| Rushton | 2 | 9 | 50 | 0 | 10 | 22 | 80 | 0 | 1 | 4 | 20 | 0 |
| Luckford | 24 | 26 | 80 | 0 | 20 | 25 | 80 | 0 | 44 | 54 | 300 | 0 |
| Goldsacs | 1 | 3 | 10 | 0 | 0.8 | 3 | 15 | 0 | 2 | 5 | 20 | 0 |
| Holme Bridge | 6 | 11 | 50 | 0 | 13 | 19 | 70 | 0 | 35 | 32 | 100 | 0 |
| Flood Relief | 0 | 1 | 5 | 0 | 2 | 4 | 10 | 0 | 3 | 7 | 30 | 0 |
| TREE ROOT |  |  |  |  | 2004 |  |  |  | 2005 |  |  |  |
| SYSTEMS (\%) | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 7 | 13 | 65 | 0 | 2 | 5 | 20 | 0 | 2 | 7 | 40 | 0 |
| Millstream | 6 | 9 | 40 | 0 | 4 | 8 | 30 | 0 | 4 | 8 | 40 | 0 |
| Rushton | 2 | 4 | 20 | 0 | 2 | 3 | 10 | 0 | 0.6 | 2 | 10 | 0 |
| Luckford | 6 | 10 | 50 | 0 | 15 | 22 | 80 | 0 | 9 | 10 | 40 | 0 |
| Goldsacs | 8 | 19 | 90 | 0 | 2 | 5 | 20 | 0 | 12 | 16 | 50 | 0 |
| Holme Bridge | 6 | 15 | 70 | 0 | 15 | 24 | 80 | 0 | 6 | 9 | 50 | 0 |
| Flood Relief | 3 | 10 | 50 | 0 | 0.4 |  | 5 | 0 | 0.5 | 2 | 5 | 0 |


| OVERHUNG | 2003 |  |  | 2004 |  |  | 2005 |  |  |  |  | Min |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| TREES (\%) | Me an | SD | Max | Min | Me <br> an | SD | Max | Min | Me <br> an | SD | Max |  |
| Railway | 15 | 16 | 50 | 0 | 9 | 13 | 40 | 0 | 11 | 18 | 60 | 0 |
| Millstream | 15 | 18 | 70 | 0 | 21 | 27 | 90 | 0 | 19 | 18 | 75 | 0 |
| Rushton | 22 | 28 | 100 | 0 | 19 | 27 | 100 | 0 | 17 | 24 | 100 | 0 |
| Luckford | 55 | 42 | 100 | 0 | 49 | 41 | 100 | 0 | 77 | 30 | 100 | 0 |
| Goldsacs | 19 | 23 | 100 | 0 | 41 | 39 | 100 | 0 | 56 | 35 | 100 | 0 |
| Holme Bridge | 65 | 36 | 100 | 0 | 61 | 46 | 100 | 0 | 66 | 39 | 100 | 0 |
| Flood Relief | 36 | 39 | 100 | 0 | 22 | 27 | 70 | 0 | 31 | 38 | 100 | 0 |
| EMERGENT | 2003 |  |  |  | 2004 |  |  |  | 2005 |  |  |  |
| SPARSE (\%) | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 5 | 10 | 25 | 0 | 1 | 3 | 10 | 0 | 1 | 5 | 20 | 0 |
| Millstream | 4 | 6 | 25 | 0 | 1 | 3 | 10 | 0 | 3 | 6 | 30 | 0 |
| Rushton | 8 | 13 | 50 | 0 | 7 | 15 | 70 | 0 | 3 | 6 | 20 | 0 |
| Luckford | 0 | 0 | 0 | 0 | 2 | 5 | 20 | 0 | 0 | 0 | 0 | 0 |
| Goldsacs | 0.6 | 2 | 10 | 0 | 1 | 2 | 5 | 0 | 0 | 0 | 0 | 0 |
| Holme Bridge | 2 | 5 | 25 | 0 | 1 | 3 | 15 | 0 | 3 | 6 | 20 | 0 |
| Flood Relief | 0 | 1 | 5 | 0 | 10 | 28 | 95 | 0 | 2 | 6 | 30 | 0 |
| EMERGENT | 2003 |  |  |  | 2004 |  |  |  | 2005 |  |  |  |
| DENSE (\%) | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 88 | 22 | 100 | 0 | 57 | 40 | 100 | 0 | 55 | 43 | 100 | 0 |
| Millstream | 1 | 3 | 10 | 0 | 1 | 2 | 10 | 0 | 3 | 9 | 40 | 0 |
| Rushton | 26 | 29 | 90 | 0 | 31 | 30 | 95 | 0 | 28 | 34 | 100 | 0 |
| Luckford | 3 | 8 | 30 | 0 | 2 | 5 | 20 | 0 | 0 | 2 | 10 | 0 |
| Goldsacs | 10 | 19 | 80 | 0 | 6 | 13 | 50 | 0 | 8 | 16 | 50 | 0 |
| Holme Bridge | 4 | 9 | 30 | 0 | 5 | 7 | 20 | 0 | 8 | 15 | 60 | 0 |
| Flood Relief | 15 | 32 | 100 | 0 | 2 | 4 | 20 | 0 | 2 | 4 | 10 | 0 |
| SUBMERGED | 2003 |  |  |  | 2004 |  |  |  | 2005 |  |  |  |
| SPARSE (\%) | Mean | SD | Max | Min | Mean | SD | Max | Min | Mean | SD | Max | Min |
| Railway | 5 | 10 | 25 | 0 | 1 | 3 | 10 | 0 | 1 | 5 | 20 | 0 |
| Millstream | 4 | 6 | 25 | 0 | 1 | 2 | 10 | 0 | 3 | 6 | 30 | 0 |
| Rushton | 8 | 13 | 50 | 0 | 7 | 15 | 70 | 0 | 3 | 6 | 20 | 0 |
| Luckford | 0 | 0 | 0 | 0 | 2 | 5 | 20 | 0 | 0 | 0 | 0 | 0 |
| Goldsacs | 1 | 2 | 10 | 0 | 1 | 12 | 5 | 0 | 0 | 0 | 0 | 0 |
| Holme Bridge | 2 | 5 | 25 | 0 | 1 | 3 | 15 | 0 | 3 | 6 | 20 | 0 |
| Flood Relief | 0 | 0.9 | 5 | 0 | 10 | 28 | 95 | 0 | 2 | 6 | 30 | 0 |
| SUBMERGED | 2003 |  |  |  | $2004$ |  |  |  | $2005$ |  |  |  |
| DENSE (\%) | Mean | SD | Max | Min | Mean | SD | Max | Min |  | SD | Max | Min |
| Railway | 0 | 0 | 0 | 0 | 1 | 3 | 15 | 0 | 0 | 0 | 0 | 0 |
| Millstream | 0 | 1 | 3 | 0 | 21 | 28 | 90 | 0 | 28 | 36 | 100 | 0 |
| Rushton | 16 | 25 | 100 | 0 | 18 | 21 | 70 | 0 | 9 | 17 | 60 | 0 |
| Luckford | 1 | 4 | 20 | 0 | 0 | 2 | 10 | 0 | 0 | 0 | 0 | 0 |
| Goldsacs | 0 | 0 | 0 | 0 | 0 | 1 | 5 | 0 | 0 | 0 | 0 | 0 |
| Holme Bridge | 9 | 22 | 80 | 0 | 3 | 13 | 70 | 0 | 7 | 21 | 90 | 0 |
| Flood Relief | 16 | 32 | 100 | 0 | 15 | 30 | 100 | 0 | 26 | 39 | 100 | 0 |

## Appendix 10 Two-way ANOVA results for differences between side-channel sections

Two-way $A N O V A$ s were used to elucidate any differences between catch and species diversity in the four 50 m sections of each side channel. As a large number of tests were undertaken to examine this, Bonferroni corrections were applied. Using the Bonferroni correction $P$-values $<0.00029$ were significant.

|  | ESMS |  | Flood relief |  | Goldsacs |  | Holme Bridge |  | Luckford |  | Railway |  | Rushton |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  | $F$ | $P$ | $F$ | $P$ | $F$ | $P$ | $F$ | $P$ | $F$ | $P$ | $F$ | $P$ | $F$ | $P$ |
| 3ullhead | 2.93 | 0.048 | 0.83 | 0.488 | 0.69 | 0.565 | 1.78 | 0.169 | 0.66 | 0.58 | 1 | 0.405 | 5.74 | 0.003 |
| lace | 2.5 | 0.077 | 0.96 | 0.425 | 0.3 | 0.828 | 1 | 0.405 | 1.32 | 0.28 | 1 | 0.405 |  | 0.405 |
| Jel | 3.74 | 0.02 | 1.16 | 0.341 | 1.85 | 0.158 | 1.08 | 0.37 | 2.7 | 0.06 | 3.77 | 0.02 | 0.81 | 0.497 |
| ${ }^{\text {Flounder }}$ | 2 | 0.133 | 1 | 1 | 2.73 | 0.059 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | $\backslash$ |
| Jrayling | 1.1 | 0.362 | 1 | 1 | 0.65 | 0.59 | 1 | 1 | 1 | 1 | 1 | 1 | 1 | 1 |
| judgeon | 1.44 | 0.248 | 1 | 0.405 | 1 | 1 | 1 | 0.405 | 3.1 | 0.04 | 1 | 1 | 0.72 | 0.55 |
| amprey | 1.17 | 0.334 | 2.2 | 0.107 | 0.41 | 0.745 | 2.2 | 0.107 | 1.89 | 0.15 | 0.79 | 0.51 | 1.34 | 0.277 |
| Minnow | 1.8 | 0.166 | 1.19 | 0.328 | 0.85 | 0.476 | 0.95 | 0.427 | 1.11 | 0.36 | 1.08 | 0.372 | 0.58 | 0.632 |
| ıerch | 1 | 0.405 | 1 | 1 | 1 | 1 | 1 | 0.405 | 1.66 | 0.2 | 1 | 1 | 1.37 | 0.268 |
| 'ike | 1.04 | 0.388 | 3.26 | 0.034 | 2.2 | 0.107 | 10.3 | 0 | 2.79 | 0.06 | 2.12 | 0.117 | 3.18 | 0.037 |
| 2oach | 1.13 | 0.352 | 2.63 | 0.067 | 0.2 | 0.898 | 0.89 | 0.455 | 1.62 | 0.2 | 1 | 1 | 1.34 | 0.278 |
| almon | 3.67 | 0.022 | 1 | 1 | 0.88 | 0.459 | 1 | 0.405 | 1.54 | 0.22 | 1 | , | 1 | 1 |
| itickleback | 1.06 | 0.378 | 0.84 | 0.479 | 2.55 | 0.072 | 0.65 | 0.59 |  | 0.41 | 1.37 | 0.268 | 1.25 | 0.308 |
| itone loach | 1.38 | 0.267 | 4.21 | 0.013 | 1.6 | 0.208 | 2.37 | 0.089 | 3.16 | 0.04 | 0.53 | 0.662 | 1.01 | 0.401 |
| rout | 1.23 | 0.315 | 1 | 1 | 5.26 | 0.004 | 1 | 0.405 | 1.24 | 0.31 | 1 | 1 | 1 | 1 |
| 'hannon's H' | 3.63 | 0.023 | 0.91 | 0.447 | 1.03 | 0.393 | 2.13 | 0.115 | 0.72 | 0.55 | 1.59 | 0.211 | 8.1 | 0 |
| hannon's E | 2.08 | 0.122 | 0.91 | 0.447 | 0.64 | 0.595 | 6.34 | 0.002 | 0.06 | 0.98 | 1.03 | 0.392 | 3.81 | 0.019 |
| Lichness | 5.77 | 0.003 | 0.55 | 0.652 | 0.42 | 0.739 | 4.11 | 0.014 | 10.8 | 0 | 1.76 | 0.174 | 5.18 | 0.005 |

## Appendix 11 Habitat Stability Index

Appendix 11. Habitat stability index for each side-channel studied (2003-2005). Rows give the relative stability of each abiotic factor between streams during different seasons over the three years. The last row gives the average stability (the 'Habitat Stability Index') for each stream. Variability in the stability of side-channels is not due to any one factor strongly weighting the average i.e. different abiotic factors are the strongest influence on the total index for different channels. (FR, Flood Relief; GS, Goldsacs; HB, Holme Bridge; LL, Luckford; MS, Millstream; RU, Rushton; RW, Railway).

|  | FR |  | GS | HB | LL | MS | RU |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| RW |  |  |  |  |  |  |  |
| Depth (cm) | 1.11 | 0.77 | 1.10 | 0.87 | 0.89 | 0.75 | 0.59 |
| Width (m) | 0.58 | 0.27 | 1.07 | 0.60 | 0.89 | 0.31 | 0.53 |
| DO (mg/l) | 0.86 | 0.79 | 0.65 | 1.18 | 0.73 | 0.65 | 0.72 |
| Temperature ( ${ }^{\circ} \mathrm{C}$ ) | 1.18 | 0.73 | 1.06 | 0.76 | 1.06 | 0.75 | 1.10 |
| Glide (\%) | 0.32 | 0.74 | 0.70 | 0.24 | 0.14 | 0.28 | 0.29 |
| Slack (\%) | 0.26 | 0.75 | 0.64 | 0.24 | 0.19 | 0.03 | 0.17 |
| Clay (\%) | 1.00 | 0.32 | 0.62 | 1.33 | 0.90 | 1.30 | 0.85 |
| Gravel (\%) | 0.80 | 0.26 | 0.66 | 0.78 | 0.72 | 1.14 | 0.16 |
| Sand (\%) | 0.25 | 0.63 | 0.75 | 0.17 | 0.54 | 0.40 | 0.03 |
| Silt (\%) | 0.74 | 0.37 | 0.76 | 0.82 | 0.14 | 0.88 | 0.53 |
| Branches/logs (\%) | 0.28 | 0.24 | 1.29 | 1.54 | 0.27 | 0.43 | 0.27 |
| Emergent dense (\%) | 0.76 | 0.53 | 0.40 | 0.25 | 0.19 | 0.87 | 1.04 |
| Emergent Sparse (\%) | 0.59 | 0.48 | 0.48 | 0.17 | 0.51 | 1.88 | 1.09 |
| Overhanging trees (\%) | 0.99 | 0.95 | 0.75 | 0.95 | 0.39 | 0.59 | 0.42 |
| Submerged Dense (\%) | 1.56 | 0.05 | 1.00 | 0.14 | 1.24 | 0.90 | 0.11 |
| Submerged sparse (\%) | 2.02 | 0.22 | 0.58 | 0.17 | 0.47 | 1.17 | 0.70 |
| Tree root systems (\%) | 0.77 | 1.63 | 0.95 | 0.94 | 0.95 | 0.34 | 0.82 |
| TOTAL | $\mathbf{1 4}$ | $\mathbf{9 . 7}$ | $\mathbf{1 3}$ | $\mathbf{1 1}$ | $\mathbf{1 0}$ | $\mathbf{1 3}$ | $\mathbf{9 . 4}$ |

Appendix 12 Abiotic factors measured in sidechannels



Appendix 12 Abiotic factors (side-channel water depth, rainfall at East Stoke and Discharge at the East Stoke gauging weir) measured during the study and used for regression analysis with fish movements (see Appendix 13). Rainfall data was not available for the full study period.

# Appendix 13 Regression results for abiotic factors on fish movements 

Table 13.1 Regression results for eel (Anguilla anguilla) movements measured by PIT telemetry according to abiotic factors. See section 5.2.1 for description of methods used to collect abiotic data and section 3.6 for methods on PIT telemetry.

| Season | Location | Abiotic factor | R2 | P-value | n | Coefficient |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Autumn | Millstream | Channel depth | $52.3 \%$ | 0.000 | 24 | 14.24 |
| Autumn | Millstream | Channel depth change | $24.9 \%$ | 0.008 | 24 | 13.89 |
| Autumn | Millstream | River discharge | $40.0 \%$ | 0.001 | 24 | 1.47 |
| Autumn | Millstream | River discharge change | $24.5 \%$ | 0.008 | 24 | 1.55 |
| Spring | Millstream | Daily temperature | $10.9 \%$ | 0.000 | 108 | 1.47 |

Table 13.2 Regression results for roach (Rutilus rutilus) movements measured by PIT telemetry according to abiotic factors.

| Season | Location | Abiotic factor | R2 | P-value | n | Coefficient |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Spring | Millhead | Daily temperature | $15.4 \%$ | 0.015 | 32 | 0.44 |
| Summer | Millstream | Channel depth | $43.7 \%$ | 0.022 | 15 | -7.59 |
| Summer | Millstream | Rainfall | $81.3 \%$ | 0.000 | 11 | 11.13 |
| Summer | Millstream | River discharge | $43.3 \%$ | 0.023 | 10 | -1.76 |
| Summer | Millstream | Max daily light | 53.5 | 0.024 | 8 | -0.0014 |

Table 13.3 Regression results for dace (Leuciscus leuciscus) movements measured by PIT telemetry according to abiotic factors.

| Season | Location | Abiotic factor | R2 | P-value | n | Coefficient |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Spring | Millhead | Daily temperature | $9 \%$ | 0.006 | 71 | 1.46 |
| Winter | Millhead | Daily temperature | $6.4 \%$ | 0.025 | 63 | 0.26 |
| Winter | Millhead | Channel depth | $12.7 \%$ | 0.002 | 63 | 4.38 |
| Winter | Millhead | River discharge | $11.5 \%$ | 0.006 | 54 | 0.44 |
| Autumn | Millstream | Rainfall | 69.4 | 0.002 | 10 | 3.19 |

Table 13.4 Regression results for pike (Esox lucius) movements measured by PIT telemetry according to abiotic factors.

| Season | Location | Abiotic factor | R2 | P-value | n | Coefficient |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Winter | Railway | Channel depth | $24.4 \%$ | 0.030 | 16 | 9.12 |
| Autumn | Millhead | Daily temperature | $12.3 \%$ | 0.048 | 25 | -0.35 |
| Autumn | Millhead | Rainfall | $10.3 \%$ | 0.065 | 25 | 2.89 |
| Spring | Millhead | Daily temperature | $37.2 \%$ | 0.036 | 25 | -0.0081 |
| Autumn | Millstream | Rainfall | $65.5 \%$ | 0.000 | 25 | 7.72 |
| Spring | Millstream | Channel depth | $27.2 \%$ | 0.000 | 51 | 17.66 |
| Spring | Millstream | River discharge | $48.6 \%$ | 0.000 | 51 | 2.18 |
| Summer | Millstream | Rainfall | $15.7 \%$ | 0.007 | 40 | 9.13 |
| Winter | Millstream | Channel depth | $25.4 \%$ | 0.078 | 10 | 3.14 |
| Winter | Millstream | River discharge | $39.8 \%$ | 0.055 | 8 | 0.44 |
| Winter | Millstream | Max daily light | $37.2 \%$ | 0.081 | 8 | 0.02 |
| Spring | Rushton | Rainfall | $15.3 \%$ | 0.013 | 34 | 33.38 |
| Summer | Luckford | River discharge | $38.0 \%$ | 0.034 | 10 | -28.08 |

# Appendix 14 Large Scale High Flow Experiment 

## INTRODUCTION

Floods influence the behaviour of most fish species as they respond to the increased discharge or newly available areas (David and Closs 2002). Some species take advantage of the feeding opportunities of invertebrates that have been washed downstream or the protection from visual predators in cloudy, sediment-loaded water, while others must shelter from the high flows (Valdez et al. 2001). Understanding the spatial scale of fishes response to flood events is important in enabling efficient implementation of river management for habitat restoration and maintenance. This investigation used radio-tracking of pike and dace to investigate fish behavioural responses to an artificial flood.

## MATERIALS AND METHODS

It was possible to control the flow of the millstream downstream of the fluvarium by adjusting hatches both in the fluvarium and of channels separating just upstream and bypassing the fluvarium. Standard autumn low flow conditions were maintained from $1^{\text {st }}$ October 2003 onwards by leaving hatches low, thus preventing increases in flow during natural high flow periods. This also caused water to back up in the Millhead behind the fluvarium and into the main river, creating a head of water that could be used to generate high flow conditions.

Five pike (Mean FL $\pm$ SD: $44.9 \pm 8 \mathrm{~cm}$ ) and seven dace (Mean FL $\pm$ SD: $24.1 \pm 0.8$ cm ) captured in the millstream by electric fishing were externally tagged on $7^{\text {th }}$ January 2004 with TW-4 tags as described in section 3.8. Three previously internally tagged pike were present in the millstream at this time (Mean FL $\pm$ SD: $74.3 \pm 23.2$ cm ). These fish were tracked at dawn, midday and dusk daily for 3 weeks prior to the experimental flood. This enabled a short-term home range to be estimated.

The experimental flood was carried out on $27^{\text {th }}$ January. By this time three pike and five dace remained in the millstream between the fluvarium and the millstream exit. Fish were tracked every 10 minutes from 09.00 until 14.00 . Hatch opening began at 10.00 and one hatch was opened $10 \%$ every 15 minutes until 11.30. After this two hatches were opened $10 \%$ every 15 minutes to simulate natural increasing flow. Peak flow (all hatches fully open) occurred at 13.30 at the fluvarium and 10 minutes later at the millstream exit. While hatches stayed open for the remainder of the experiment, the water flow reduced after 14.00 once the head of water in the Millhead had passed through.

Fish were tracked hourly from 14.00 until an hour after dark (20.00) to monitor post flood movements. Fish were then once again tracked at dawn, midday and dusk for two weeks following the flood in order to calculate post-flood short-term home ranges.

## RESULTS

The response of pike and dace to discharge was investigated at a fine scale during the large scale experimental flood. While the sample size was small, with only three pike and five dace, the results can be used to suggest trends in activity following a flood event. The two species were combined during the analysis because no differences were apparent between species in distance or direction moved upon inspection of the data. Activity was compared before and after the onset of the flood event. It was not possible to make direct comparisons with activity and movements on the day of the flood as fish locations were collected more frequently than during the pre- and post flood periods. Despite this differences in activity were apparent before and after the flood event. The mean distance travelled by fish between subsequent relocations was significantly higher after the flood event ( $t$-test, $P<0.05$ ) (Figure 14.1). This increase in activity was not due to fish being washed downstream on the first increase in flow and later recovering their original positions as upstream and downstream movement was evenly distributed throughout the experiment (Figure 14.2).


Figure 14.1 Mean distance travelled by pike and dace (combined $n=8$ ) two weeks before and two weeks after the flood event.


Figure 14.2 Up- and downstream movements (positive and negative values respectively) of pike and dace before, during and after the flood event. Red line denotes the period of the high flow.

In addition to increasing activity, the structure of fish's home ranges also changed during the higher discharges post-flood. The number of home range cores increased
significantly during the post-flood period, despite a high level of variability ( $t$-test, $P$ $<0.05$ ) (Figure 14.3). However, the size of the home range did not and tended towards a decrease in size after the flood event ( $t$-test, $P>0.05$ ) (Figure 14.4). This suggests that fish concentrated their time in smaller, but more widely interspersed focal areas and spent less time in the river between these areas.


Figure 14.3 Number of cores in the home range of pike and dace (combined $n=8$ ) before and after the flood event.


Figure 14.4 Home range area of pike and dace (combined $n=8$ ) before and after the flood event.

## DISCUSSION

Dace and pike responded to the experimental flood with an increase in activity with time focussed in a larger number of smaller areas (cores). Fish were not displaced downstream during the flood, however this was a small scale flood in terms of the fish and during a larger natural flood in the River Thames pike were displaced downstream (Langford 1979). This post-flood increased activity may be due to fish taking advantage of feeding opportunities from aquatic and terrestrial invertebrates washed downstream. However, the reduction in core ranges indicated that fish focussed their time in a smaller area suggesting that they were sheltering from the increased flow. Pike in the River Frome were previously found to visit side-channels during high flow events (Masters et al. 2002), in fact the distance the fish was found up the side-channel was related to the discharge. Masters et al (2002) argued that this use of side-channels was for feeding, however a simpler explanation would be that they were used for sheltering from high flows. It is likely that as flows increase more pike move to the side-channels for shelter and so fish distribute further from the main river. Pike could probably maintain their position in the main river but it may be energetically more efficient to shelter in a side-channel when possible because of the lower locomotor costs.

While small scale floods do not seem to have a large effect on the fish behaviour it is possible that floods have long term beneficial effects on available habitat. Experimental flooding on a regulated river was found to enhance the habitat of trout without influencing their behaviour during the flood events (Ortlepp and Murle 2003). This lead to a three-fold increase in the number of redds created since the initiation of the flooding.

References
2000. Directive 2000/60/EC of the European Parliament and of the Council of 23 October 2000 establishing a framework for Community action in the field of water policy In Official Journal of the European Community. pp. 1-72.
Aebischer, N.J., Robertson, P.A., and Kenward, R.E. 1993. Compositional analysis of habitat use from animal radio-tracking data. Ecology 74(5): 1313-1325.

Alabaster, J. S., and Lloyd, R. 1982 Water quality criteria for freshwater fish - Second edition. Food and Agriculture Organisation of the United Nations. London: Butterworth Scientific. pp 278.
Allouche, S., Thevenet, A., and Gaudin, P. 1999. Habitat use by chub (Leuciscus cephalus L. 1766) in a large river, the French Upper Rhone, as determined by radiotelemetry. Archiv für Hydrobiologie 145(2): 219-236.

Amoros, C., and Roux, A.L. 1988. Interaction between water bodies within the floodplains of large rivers: function and development of connectivity. 2nd International Seminar of the International Association for Landscape Ecology, pp. 125-130.

Anonymous. 2005. The Frome, Piddle and Purbeck catchment abstraction management strategy, Environment Agency, Exeter.

Armitage, P.D., Szoszkiewicz, K., Blackburn, J.H., and Nesbitt, I. 2003. Ditch communities: a major contributor to floodplain biodiversity. Aquatic Conservation: Marine and Freshwater Ecosystems 13(2): 165-185.

Baade, U., and Fredrich, F. 1998. Movement and pattern of activity of the roach in the River Spree, Germany. Journal of Fish Biology 52.
Bagenal, T. 1978. Methods for Assessment of Fish Production in Fresh Waters. Blackwell Scientific Publications.
Bagenal, T. 1973. Identification of British Fishes. Hulton Educational Publications Ltd. pp 199.
Bahr, M.A., and Shrimpton, J.M. 2004. Spatial and quantitative patterns of movement in large bull trout (Salvelinus confluentus) from a watershed in north-western British Columbia, Canada, are due to habitat selection and not differences in life history. Ecology of Freshwater Fish 13(4): 294-304.
Baras, E. 1997. Environmental determinants of residence area selection by Barbus barbus in the River Ourthe. Aquatic Living Resources 10(4): 195-206.

Baras, E., Jeandrain, D., Serouge, B., and Philippart, J.C. 1998. Seasonal variations in time and space utilization by radio-tagged yellow eels Anguilla anguilla (L.) in a small stream. Hydrobiologia 371/372: 187-198.

Baras, E., and Nindaba, J. 1999. Seasonal and diel utilisation of inshore microhabitats by larvae and juveniles of Leuciscus cephalus and Leuciscus leuciscus. Environmental Biology of Fishes 56: 183-197.

Barreto, G.R., and MacDonald, D.W. 2000. The decline and local extinction of a population of water voles, Arvicola terrestris, in southern England. International Journal of Mammalian Biology 65(2): 110-120.

Bartozova, S., and Jurajda, P. 2001. A comparison of $0+$ fish communities in borrow pits under different flooding regime. Folia Zoologica 50(4): 305-315.

Bayley, P.B. 1995. Understanding large river-floodplain ecosystems. BioScience 45(3): 153-158.

Bayley, P.B., O'Hara, K., and Steel, R. 2000. Defining and achieving fish habitat rehabilitation in large, low-gradient rivers. In Management and ecology of river fisheries. Edited by I.G. Cowx. Fishing News Books, Oxford.

Beaumont, W.R.C., Clough, S., Ladle, M., and Welton, J.S. 1996. A method for the attachment of miniature radio tags to small fish. Fisheries Management and Ecology 3: 201-207.
Beaumont, W.R.C., Cresswell, B., Hodder, K.H., Masters, J.E.G., and Welton, J.S. 2002. A simple activity monitoring radio tag for fish. Hydrobiologia 483(1-3): 219-224.

Beaumont, W.R.C., Hodder, K.H., Masters, J.E.G., Scott, L.J., and Welton, J.S. 2005. Activity patterns in pike (Esox Lucius), as determined by motion-sensing telemetry. In Aquatic telemetry: advances and applications. Proceedings of the Fifth Conference on Fish Telemetry held in Europe. Ustica, Italy, 9-13 June 2003. Edited by M.T. Spedicato, G. Marmulla and G. Lembo. FAO/COISPA, Rome. pp. 231-243.
Behrmann-Godel, J., and Eckmann, R. 2003. A preliminary telemetry study of the migration of silver European eel (Anguilla anguilla L.) in the River Mosel, Germany. Ecology of Freshwater Fish 12: 196-202.
Bell, E., Duffy, W.G., and Roelofs, T.D. 2001. Fidelity and survival of juvenile coho salmon in response to a flood. Transactions of the American Fisheries Society 130: 450-458.

Billard, R. 1996. Reproduction of pike: gametogenesis, gamete biology and early development. In Pike Biology and Exploitation. Edited by J. Craig. Chapman \& Hall, London. p. 298.

Blundell, G.M., Maier, J.A.K., and Debevec, E.M. 2001. Linear home ranges: Effects of smoothing, sample size and autocorrelation on kernel estimates. Ecological Monographs 71(3): 469-489.

Borcherding, J., Bauerfeld, M., Hintzen, D., and Neuman, D. 2002. Lateral migrations of fishes between floodplain lakes and their drainage channels in the Lower Rhine: diel and seasonal aspects. Journal of Fish Biology 61: 1154-1170.
Börger, L., N. Franconi, F. Ferretti, F. Meschi, G. De Michele, A. Gantz, , and T. Coulson. (2006) An integrated approach to identify spatiotemporal and individual-level determinants of animal home range size. The American Naturalist, 168, 471-485.

Bowman, J. 2003. Is dispersal distance of birds proportional to territory size? Canadian Journal of Zoology 81: 195-202.

Bowman, J., Jaeger, J.A.G., and Fahrig, L. 2002. Dispersal distance of mammals is proportional to home range size. Ecology 83(7): 2049-2055.

Bregazzi, P. R. and Kennedy, C. R. (1980) The biology of pike, Esox lucius (L.), in a southern eutrophic lake. Journal of Fish Biology. 17. 91-122.

Bridcut, E.E., and Giller, P.S. 1993. Movement and site fidelity in young brown trout Salmo trutta populations in a southern Irish stream. Journal of Fish Biology 43(6): 889-899.
Brook, A. J. and Bromage, N. R. 1989. Photoperiod - the principal environmental cue for reproduction in the dace, Leuciscus leuciscus (L.). Journal of Interdisciplinary Cyclical Research. 19. 165-166.

Broomhall, L.S., Mills, M.G.L., and du Toit, J.T. 2003. Home range and habitat use by cheetahs (Acinonyx jubatus) in the Kruger National Park. Journal of the Zoological Society of London 261: 116-128.
Brosse, S., and Lek, S. 2000. Modelling roach (Rutilus rutilus) microhabitat using linear and nonlinear techniques. Freshwater Biology 44: 441-452.
Brown, J.H. 1984. On the relationship between abundance and distribution of species. American Naturalist 124: 225-279.

Brown, R.S., Power, G., and Beltoas, S. 2001. Winter movements and habitat use of riverine brown trout, white sucker and common carp in relation to flooding and ice break-up. Journal of Fish Biology 59: 1126-1141.

Bryce, J., Johnson, P.J., and MacDonald, D.W. 2002. Can niche use in red and grey squirrels offer clues for their apparent coexistence? Journal of Applied Ecology 39: 875-887.

Buijse, A.D., Coops, H., Staras, M., Jans, L.H., Van Geest, G.J., Grifts, R.E., Ibelings, B.W., Oosterberg, W., and Roozen, F.C.J.M. 2002. Restoration strategies for river floodplains along large lowland rivers in Europe. Freshwater Biology 47: 889-907.

Bunn, S.E., and Arthington, A.H. 2002. Basic principles and ecological consequences of altered flow regimes for aquatic biodiversity Environmental Management 30(4): 492-507

Burgman, M. A. and Fox, J. C. (2003) Bias in species range estimates from minimum convex polygons: implications for conservation and options for improved planning. Animal Conservation, 6, 19-28
Burt, W.H. 1943. Territoriality and home range concepts as applied to mammals. Journal of Mammology 24: 346-352.
Calhoun, J.B., and Casby, J.U. 1958. Calculation of home range and density of small mammals. In Public Health Monograph 55. United States Public Health Service.

Carbine, W.F., and Applegate, W.C. 1946. The movement of marked pike in Houghton Lake and Muskegon River. Papers of the Michigan Academy of Science, Arts and Letters 32: 215-238.
Carle, F.L., and Strub, M.R. 1978. New method for estimating population size from removal data Biometrics 34(4): 621-630.
Casey, H. 1969. The chemical composition of some Southern English chalk streams and its relation to discharge. River Authorities Association Yearbook: 100113.

Cassleman, J.M., and Lewis, C.A. 1996. Habitat requirements of northern pike (Esox lucius). Canadian Journal of Fisheries and Aquatic Sciences 53(Suppl. 1): 161174.

Cattaneo, F. 2005. Does hydrology constrain the structure of fish assemblages in French streams? Local scale analysis. Archiv für Hydrobiologie 164(3): 345365.

Chapman, D.W. 1968. Production. In Fish production in fresh waters. Edited by W.E. Ricker. Blackwell Scientific Publications, Oxford and Edinburgh.
Chapman, L. J., Mackay, W. C. and Wilkinson, C. W. 1989 Feeding flexibility in nothern pike (Esox lucius) - Fish versus invertebrate prey. Canadian Journal of Fisheries and Aquatic Sciences. 46. 666-669
Chorley, R.J. 1962. Geomorphology and general systems theory. U.S Geological Survey Professional Paper 500-B: 10p.
Clough, S. 1997. Diel migration and site fidelity in a stream dwelling cyprinid, Leuciscus leuciscus. Journal of Fish Biology 50: 1117-1119.
Clough, S. 1998. Migration and habitat use of the dace (Leuciscus leuciscus (L.)) in an English chalk stream, PhD thesis, University of St Andrews.
Clough, S., Garner, P., Deans, D., and Ladle, M. 1998. Post-spawning movements and habitat selection of dace (Leuciscus leuciscus (L.)) in the River Frome, Dorset. Journal of Fish Biology 53(5): 1060-1070.
Cook, M.F., and Bergersen, E.P. 1988. Movements, habitat selection, and activity periods of northern pike in Eleven Mile Reservoir, Colorado. Transactions of the American Fisheries Society 117: 495-502.
Copp, G.H. 1997. Importance of marinas and off-channel water bodies as refuges for young fishes in a regulated lowland river. Regulated Rivers: Research \& Management 13: 303-307.
Cowx, I. G. 2001 Factors influencing coarse fish populations in rivers. Environment Agency R\&D project W2-020.
Crisp, D.T., Matthews, A.M., and Westlake, D.F. 1982. The temperatures of nine flowing waters in southern England. Hydrobiologia 89: 193-204.
Crook, D.A., Robertson, A.I., King, A.J., and Humphries, P. 2001. The influence of spatial scale and habitat arrangement on diel patterns of habitat use by two lowland river fishes. Oecologia 129: 525-533.
Crossman, E.J. 1977. Displacement, and home range movements of muskellunge determined by ultrasonic tracking. Environmental Biology of Fishes 1(2): 145158.

Cummins, K.W. 1974. Structure and function of stream ecysystems. BioScience 24: 631-641.

Dalke, P.D., and Sime, P.R. 1938. Home and seasonal ranges of eastern cottontail in Connecticut. Transcripts of the North American Wildlife Conference 3: 659669.

Dauba, F., and Biro, P.A. 1992. Growth of bream, Abramis brama L., in two outside basins of different state of Lake Balaton. Hydrobiologia 77: 225-235.
David, B.O., and Closs, G.P. 2002. Behavior of a stream-dwelling fish before, during, and after high-discharge events. Transactions of the American Fiseries Society 131: 762-771.

Davies, C. E., Shelley, J., and Harding, P. T., McLean I. F. G., Gardiner, R. and Peirson G. 2004 Freshwater fishes in Britain: the species and their distribution. Harley Books, England. pp 176.

Dawson, F.H., Clinton, E.M.F., and Ladle, M. 1991. Invertebrates on cut weed removed during weed-cutting operations along and English river, the River Frome, Dorset. Aquaculture and Fisheries Management 22: 113-121.

Diana, J.S. 1980. Diel activity pattern and swimming speeds of Northern pike (Esox lucius) in Lac Ste. Anne, Alberta. Canadian Journal of Fisheries and Aquatic Sciences 37: 1454-1458.

Dixon, K.R., and Chapman, J.A. 1980. Harmonic mean measure of animal activity areas. Ecology 61: 1040-1044.
Doak, D.F. 1995. Source-sink models and the problem of habitat degradation: general models and applications to the Yellowstone grizzly. Conservation Biology 9: 1370-1379.

Downhower, J.F., and Brown, L. 1980. Mate preference of female mottled sculpins Cottus bairdi. Animal Behaviour 28: 215-223.

Dunham, J.B., and Rieman, B.E. 1999. Metapopulation structure of bull trout: Influences of physical, biotic, and geometrical landscape characteristics. Ecological Applications 9: 642-655.
Eklov, P., and Diehl, S. 1994. Piscivore efficiency and refuging prey: the importance of predator search mode. Oecologia 98: 345-353.
Erman, D.C., Andrews, E.D., and Yoder-Williams, M. 1988. Effects of winter floods on fishes in the Sierra Nevada. Canadian Journal of Fisheries and Aquatic Sciences 45: 2195-2200.

European Commission (2000) Directive 2000/60/EC of the European Parliament and of the Council of 23 October 2000 establishing a framework for Community action in the field of water policy In Official Journal of the European Community, pp. 1-72.

Fedriani, J.M., Delibes, M., Ferreras, P., and Roman, J. 2002. Local and landscape habitat determinants of water vole distribution in a patchy Mediterranean environment. Ecoscience 9(1): 12-19.

Fisher, D.O. 2000. Effect of vegetation structure, food and shelter on the home range and habitat use of an endangered wallaby. Journal of Applied Ecology 37: 660-671.

Ford, E. 1933. An account of herring investigations conducted at Plymouth during the years from 1924-1933. Journal of the Marine Biological Association 19: 305384.

Fraser, C.M. 1916. Growth of the spring salmon. Transactions of the Pacific Society Seattle, Second Annual Meeting, 1915: 29-39.
Fraser, D.F., Gilliam, J.F., MacGowan, M.P., Arcaro, C.M., and Guillozet, P.H. 1999. Habitat quality in a hostile river corridor. Ecology 80(2): 597-607.
Freeman, M.C., Bowen, Z.H., Bovee, K.D., and Irwin, E.R. 2001. Flow and habitat effects on juvenile fish abundance in natural and altered flow regimes. Ecological Applications 11: 179-190.
Frost, W.E., and Kipling, C. 1967. A study of reproduction, early life, weight-length relationship and growth of pike Esox lucius L., in Windermere. Journal of Animal Ecology 36: 659-693.
Fuller, M.R., K.E. Church, J.J. Millspaugh, and R.E. Kenward. (2005). Wildlife Telemetry. In Manual of Wildlife Management Techniques. (ed C.L. Braun), pp. 377-417. The Wildlife Society, Maryland
Garner, P. 1996. Microhabitat use and diet of 0+ cyprinid fishes in a lentic, regulated reach of the River Great Ouse, England. Journal of Fish Biology 48: 367-382.
Getz, W.M. and C.C. Wilmers. (2004) A local nearest-neighbour convex-hull construction of home ranges and utilisation distributions. Ecography, 27, 489505.

Gore, J.A., and Shields, F.D. 1995. Can large rivers be restored? BioScience 48: 367382.

Gotelli, N.J., and Taylor, C.M. 1999. Testing metapopulation models with stream-fish assemblages. Evolutionary Ecology Research 1: 835-845.
Gozlan, R., Mastrorillo, S., Dauba, F., Tourenq, J.-N., and Copp, G.H. 1998. Multiscale analysis of habitat use during late summer for $0+$ fishes in the River Garonne (France). Aquatic Sciences 60: 99-117.

Gozlan, R.E. 1998. Environmental biology and morphodynamics of the sofie Chondrostoma toxostoma (Cyprindidae), with emphasis on early development, Department of Environmental Sciences, University of Hertfordshire, London.
Greenhalgh, M. 1999. Freshwater fish. Octopus Publishing Group Ltd.
Grenouillet, G., Pont, D., and Herisse, C. 2004. Withiin-basin fish assemblage structure: the relative influence of habitat versus spatial position on local species richness. Canadian Journal of Fisheries and Aquatic Sciences 61: 93102

Grift, R.E., Buijse, A.D., Klein Breteler, J.G.P., van Densen, W.L.T., Machiels, M.A.M., and Backx, J.J.G.M. 2001. Migration of bream between the main channel and floodplain lakes along the lower River Rhine during the connection phase. Journal of Fish Biology 59: 1033-1055.
Grossman, G.D., Ratajczak, R.E., Crawford, M., and Freeman, M.C. 1998. Assemblage organization in stream fishes: effects of environmental variation and interspecific interactions. Ecological Monographs 68(3): 395-420.
Hanski, I.K. 1994. Patch-occupancy dynamics in fragmented landscapes. Trends in Ecology and Evolution 9(4): 131-135.
Hanski, I.K. 1999. Metapopulation Ecology. Oxford University Press.
Harden Jones, F.R. 1968. Fish Migration. Edward Arnold Ltd., London.
Harmata, A.R., and Montopoli, G.R. 2001. Analysis of bald eagle spatial use of linear habitat. Journal of Raptor Research 35(3): 207-213.

Harris, S., Cresswell, W.J., Forde, P.G., Trewhella, W.J., Woollard, T., and Wray, S. 1990. Home-range analysis using radio-tracking data - a review of problems and techniques particularly as applied to the study of mammals. Mammal Review 20(2/3): 97-123.
Hawkins, L.A., Armstrong, J.D., and Magurran, A.E. 2005. Aggregation in juvenile pike (Esox lucius): interactions between habitat and density in winter. Functional Ecology 19: 794-799.

Heywood, M.J.T., and Walling, D.E. 2003. Suspended sediment fluxes in chalk streams in the Hampshire Avon catchment, UK. Hydrobiologia 494((1-3)): 111-117

Hodder, K.H., Kenward, R.E., Walls, S.S., and Clarke, R.T. 1998. Estimating core ranges: A comparison of techniques using the common buzzard (Buteo buteo). Journal of Raptor Research 32(2): 82-89.
Hodder, K.H., Masters, J.E.G., Beaumont, W.R.C., Gozlan, R.E., Pinder, A.C., Knight, C.M., and Kenward, R.E. in press. Techniques for evaluating the spatial behaviour of river-fish. Hydrobiologia.
Hoeinghaus, D.J., Layman, C.A., Arrington, D.A., and Winemiller, K.O. 2003. Spatiotemporal variation in fish assemblage structure in tropical floodplain creeks. Environmental Biology of Fishes 67(4): 379-387.
Hohausová, E., Copp, G.H., and Jankovsky, P. 2003. Movement of fish between a river and it's backwater: diel activity and relation to environmental gradients. Ecology of Freshwater Fish 12.
Holmgren, K. 2003. Omitted spawning in compensatory-growing perch. Journal of Fish Biology 62: 918-927.

Hoopes, M.F., and Harrison, S. 1998. Metapopulation, source-sink and disturbance dynamics. In Conservation science and action. Edited by W.J. Sutherland. Blackwell Science Ltd.

Howden, N.J.K. 2004. Hydrogeological controls on surface/groundwater interactions in a lowland permeable chalk catchment: implications for water quality and numerical modelling, Environmental and Water Resources Engineering Department of Civil and Environmental Engineering, Imperial College London, PhD Thesis.
Huet, M. 1954. Profiles and biology of Western European streams as related to fish management. Transactions of the American Fiseries Society 88: 153-163.
Humphries, P., King, A.J., and Koehn, J.D. 1999. Fish, flows and flood plains: links between freshwater fishes and their environment in the Murray-Darling River system, Australia. Environmental Biology of Fishes 56: 129-151.
Ibbotson, A.T., Beaumont, W.R.C., Collinson, D., Wilkinson, A., and Pinder, A.C. 2004. A cross-river antenna array for the detection of miniature passive integrated transponder tags in deep fast flowing rivers. Journal of Fish Biology 65: 1441-1443.

Jackson, D.A., Peres-Neto, P.R., and Olden, J.D. 2001. What controls who is where in freshwater fish communities - the roles of biotic, abiotic and spatial factors. Canadian Journal of Fisheries and Aquatic Sciences 58: 157-170.

Jacobsen, L., Berg, S., Jepsen, N., and Skov, C. 2004. Does roach behaviour differ between shallow lakes of different environmental state? Journal of Fish Biology 65: 135-147.

Jager, H.I., Chandler, J.A., Lepla, K.B., and Winkle, W.V. 2001. A theoretical study of river fragmentation by dams and its effects of white sturgeon populations. Environmental Biology of Fishes 60: 247-261.
Jennrich, R.J., and Turner, F.B. 1969. Measurement of non-circular home range. Journal of Theoretical Biology 22: 227-237.

Jepsen, N., Beck, S., Skov, C., and Koed, A. 2001. Behavior of pike (Esox lucius L.) $>50 \mathrm{~cm}$ in a turbid reservior and a clearwater lake. Ecology of Freshwater Fish 10: 26-34.

Jepsen, N., Koed, A., Thorstad, E. B and Baras, E. 2002. Surgical implantation of telemetry transmitters in fish: how much have we learned? Hydrobiologia 483(1-3): 239-248.

Johnson, C.J., Seip, D.R., and Boyce, M.S. 2004. A quantitative approach to conservation planning: using resource selection functions to map the distribution of mountain caribou at multiple spatial scales. Journal of Applied Ecology 41(2): 238-251.
Jungwirth, M. 1998. River continuum and fish migration - going beyond the longitudinal river corridor in understanding ecological integrity. In Fish migration and fish bypasses. Edited by M. Jungwirth, S. Schmutz and S. Weiss. Fishing News Books, Oxford. pp. 19-32.
Junk, W.J., Bayley, P.B., and Sparks, R.E. 1989. The flood pulse concept in riverfloodplain systems. Canadian Special Publication of Fisheries and Aquatic Sciences 106: 110-127.

Karas, P., and Lehtonen, H. 1993. Patterns of the movement and migration of pike (Esox lucius L.) in the Baltic Sea. Nordic Journal of Freshwater Research 68: 72-79.

Kenward, R.E. 1987. Wildlife radio tagging: Equipment, field techniques and data analysis. Academic Press, London.

Kenward, R.E. 1992. Quantity versus quality: programming for collection and analysis of radio tag data. In Wildlife telemetry - remote monitoring and tracking of animals. Edited by I.G. Preide and S.M. Swift. Ellis Horwood, Chichester, UK. pp. 231-246.
Kenward, R.E. 2001. A manual for wildlife radio tagging. Academic Press.
Kenward, R.E., Clarke, R.T., Hodder, K.H., and Walls, S.S. 2001. Density and linkage estimators of home range: nearest-neighbor clustering defines multinuclear cores. Ecology 82(7): 1905-1920.
Kenward, R.E., and Hodder, K.H. 1998. Red squirrels (sciurus vulgaris) released in conifer woodland: the effects of source habitat, predation and interactions with grey squirrels (Sciurus carolinensis). Journal of the Zoological Society of London 244: 23-32.
Kernohan, B.J., Gitzen, R.A., and Millspaugh, J.J. 2001. Analysis of animal space use and movements. In Radio Tracking and Animal Populations. Edited by J.J. Millspaugh and J.M. Marzluff. Academic Press.

Koizumi, I., and Maekawa, K. 2004. Metapopulation structure of stream-dwelling Dolly Varden charr inferred from patterns of occurrence in the Sorachi River basin, Hokkaido, Japan. Freshwater Biology 49: 973-981.

Kraus, C., and Rodel, H.G. 2004. Where have all the cavies gone? Causes and consequences of predation by the minor grison on a wild cavy population. Oikos 105(3): 489-500.
Kruuk, H. (1978) Spatial organisation and territorial behaviour of the European badger Meles meles. Journal of Zoology, 184, 1-19.
Krebs, J.R., and Davies, N.B. 1997. Behavioural ecology: an evolutionary approach. Blackwell Science Ltd.
Ladle, M., and Westlake, D.F. 1995. River and stream ecosystems of Great Britain. In River and stream ecosystems. Edited by C.E. Cushing, K.W. Cummins and G.W. Minshall. Elsevier, Amsterdam. pp. 343-388.

Lam, T.J., Nagahama, Y., Chan, K., and Hoar, W.S. 1978. Overripe eggs and postovulatory copora lutea in the three spine stickleback, Gasterosteus aculeatus L., form trachurus. Canadian Journal of Zoology 56: 2029-2036.
Lance, A.N., and Watson, A. 1980. A comment on the use of radio tracking in ecological research. In A Handbook on Biotelemetry and Radio Tracking.

Edited by C.J. Amlaner and D.W. MacDonald. Pergamon Press, Oxford. pp. 355-359.

Langford, T.E. 1979. Observations on sonic-tagged coarse fish in rivers. Proceedings of the 1st British Freshwater Fisheries Conference, University of Liverpool, Liverpool, pp. 106-114.

Langler, G.J., and Smith, C. 2001. Effects of habitat enhancement on 0 -group fishes in a lowland river. Regulated Rivers: Research \& Management 17: 677-686.
Larson, M.A. (2001). A catalogue of software to analyse radiotelemetry data. In Radio Tracking and Animal Populations (eds J.J. Millspaugh \& J.M. Marzluff), pp. 398-421. Academic Press, San Diego, USA.

Leopold, L.B., and Maddock, T. 1953. The hydraulic geometry of stream channels and some physiographic implications. U.S Geological Survey Professional Paper 252: 57p.

Lindenmayer, D.B., Ball, I., Possingham, H.P., McCarthy, M.A., and Pope, M.L. 2001. A landscape-scale test of the predictive ability of a spatially explicit model for population viability analysis. Journal of Applied Ecology 38: 36-48.
Lindenmayer, D.B., Burgman, M.A., Akcakaya, H.R., Lacy, R.C., and Possingham, H.P. 1995. A review of the generic computer programs ALEX, RAMAS/Space and Vortex for modeling the viability of wildlife populations. Ecological Modelling 82: 161-174.
Locke, H. (1996) Yellowstone to Yukon. Wildlife Conservation, 99, 24-32.
Lucas, M.C. 1992. Spawning activity of male and female pike, Esox lucius L., determined by acoustic tracking. Canadian Journal of Zoology 70: 191-196.
Lucas, M.C., and Baras, E. 2001. Migration of freshwater fishes. Blackwell Science.
Lucas, M.C., Mercer, T., McGinty, S., and Armstrong, J.D. 2000. Development and evaluation of a flat-bed passive integrated transponder detection system for recording movement of lowland river fishes through a baffled pass. In Advances in Fish Telemetry. Edited by A. Moore and I. Russell. CEFAS, Lowestoft. pp. 117-127.
Lucas, M.C., Priede, I.G., Armstrong, J.D., Gindy, A.N.Z., and De Vera, L. 1991. Direct measurements of metabolism, activity and feeding behaviour of pike, Esox lucius L., in the wild, by the use of heart rate telemetry. Journal of Fish Biology 39: 325-345.

Lusk, S., Halacka, K., Luskova, V., and Horak, V. 2001. Annual dynamics of the fish stock in a backwater of the River Dyje. Regulated Rivers: Research \& Management 17: 571-581.
MacDonald, D.W., Tew, T.E., and Todd, I.A. 2004. The ecology of weasels (Mustela nivalis) on mixed farmland in southern England. Biologia 59(2): 235-241.
Madsen, T., and Shine, R. 1996. Seasonal migration of predators and prey - a study of pythons and rats in tropical Australia. Ecology 77(1): 149-156.
Maekawa, K., Iguchi, K., and Katano, O. 1996. Reproductive success in male Japanese minnows, Pseudorasbora parva: observations under experimental conditions. Ichthyological Research 43: 257-266.
Major, R.E., and Gowing, G. 2001. Survival of red-capped robins (Petroica goodenovii) in woodland remnants of central western New South Wales, Australia. Wildlife Research 28(6): 565-571.
Mann, R.H.K. 1973. Observations on the age, growth, reproduction and food of the roach Rutilus rutilus (L.) in two rivers in southern England. Journal of Fish Biology 5: 707-736.

Mann, R.H.K. 1974. Observation on the age, growth, reproduction and food of the dace Leuciscus leuciscus (L.), in two rivers in Southern England. Journal of Fish Biology 6: 237-253.
Mann, R.H.K. 1976a. Observations on the age, growth, reproduction and food of the chub Squalius cephalus (L.) in the River Stour, Dorset. Journal of Fish Biology 8: 265-288.
Mann, R.H.K. 1976b. Observations on the age, growth, reproduction and food of the pike Esox lucius (L.) in two rivers in southern England. Journal of Fish Biology 8: 179-197.
Mann, R. H. K. 1982 The annual food consumption and prey preferences of pike (Esox lucius) in the River Frome, Dorset. Journal of Animal Ecology. 51. 8195
Mann, R.H.K., and Mills, C.A. 1986. Biological and climatic influences on the dace Leuciscus leuciscus in a Southern chalk-stream. Report of the Freshwater Biological Association. pp. 123-136.
Marchetti, M.P., Moyle, P.B. and R. Levine, (2004) Alien fishes in California watersheds: characteristics of successful and failed invaders. Ecological Applications, 14, 587-596.

Markus, N., and Hall, L. 2004. Foraging behaviour of the black flying-fox (Pteropus alecto) in the urban landscape of Brisbane, Queensland. Wildlife Research 31(3): 345-355.

Masters, J.E.G., Welton, J.S., Beaumont, W.R.C., Hodder, K.H., Pinder, A.C., Gozlan, R.E., and Ladle, M. 2002. Habitat utilisation by pike Esox lucius L. during winter floods in a southern English chalk river. Hydrobiologia 483: 185-191.

Matthiopoulos, J. (2003) Model-supervised kernel smoothing for the estimation of spatial usage. Oikos, 102, 367-377
Miller, L.M., Kallemeyn, L., and Senanan, W. 2001. Spawning-site and natal-site fidelity by northern pike in a large lake: mark-recapture and genetic evidence. Transactions of the American Fisheries Society 130: 307-316.
Mills, C. A. 1991 Reproduction and life history. In Winfield, I. J. and Nelson, J. S. (eds) Cyprinid Fishes, Systematics, Biology and Exploitation. Chapman and Hall. London. 483-504.

Minns, C.K. 1995. Allometry of home-range size in lake and river fishes Canadian Journal of Fisheries and Aquatic Sciences 52(7): 1499-1508
Molls, F. 1999. New insights into the migration and habitat use by bream and white bream in the floodplain of the River Rhine. Journal of Fish Biology 55: 11871200.

Morbey, Y.E., and Ydenberg, R.C. 2001. Protandrous arrival timing to breeding areas: a review. Ecology Letters 4: 663-673.
Morita, K., and Yamamoto, S. 2002. Effects of habitat fragmentation by damming on the persistence of stream-dwelling charr populations. Conservation Biology 16: 1318-1323.
Muller, U.K., Stamhuis, E.J., and Videler, J.J. 2000. Hydrodynamics of unsteady fish swimming and the effects of body size: Comparing the flow fields of fish larvae and adults. Journal of Experimental Biology 203(2): 193-206.
Mysterud, A., Perex-Barberia, F.J., and Gordon, I.J. 2001. The effect of season, sex and feeding style on home range area versus body mass scaling in temperate ruminants. Oecologia 127(1): 30-39.
Neumann, D., Staas, S., Molls, F., Seidenberg-Busse, C., Petermeier, A., and Rutschke, J. 1996. The significance of man-made lentic waters for the ecology
of the Lower River Rhine, especially for the recruitment of potamal fish. Archiv für Hydrobiologie Supplementband 113: 267-278.
Niaman, R.J., and Decamps, H. 1990. The ecology and management of aquaticterrestrial ecotones. UNESCO and The Parthenon Publishing Group.
Nicholls, A. O. and Margules, C. R. 1993 An upgraded reserve selection algorithm. Biological Conservation 64: 165-169.
Nilsson, P.A., Nilsson, K., and Nystrom, P. 2000. Does risk of intraspecific interactions induce shifts in prey-size preference in aquatic predators. Behavioural Ecology and Sociobiology 48: 268-275.
Northcote, T.G. 1978. Migratory strategies and production in freshwater fishes. In Ecology of Freshwater Fish Production. Edited by S.D. Gerking. Blackwell Science. pp. 326-259.

Northcote, T.G. 1998. Migratory behaviour of fish and its significnace to movement through riverine fish passage facilities. In Fish migration and fish bypasses. Edited by M. Jungwirth, S. Schmutz and S. Weiss. Fishing News Books, Oxford. pp. 3-18.
Oberdoff, T., Hugueny, B., and Vigneron, T. 2001. Is assemblage variability related to environmental variability? An answer for riverine fish. Oikos 93: 419-428.
Olson, D.M., and Dinerstein, E. 1998. The globe 200: A representation approach to conserving the earth's most biologically valuable ecoregions. Conservation Biology 12(3): 502-515.
Ortlepp, J., and Murle, U. 2003. Effects of experimental flooding on brown trout (Salmo trutta fario L.): The River Spol, Swiss National Park. Aquatic Sciences 65: 232-238.
Paolillo, S.A.G. 1969. Hydroecology of the River Frome catchment (Southern England). Memorie E Note Dell'instituto Di Geologia Applicata Napoli 11: 169.

Peach, W.J., Denny, M., Cotton, P.A., Hill, I.F., Gruar, D., Barritt, D., Impey, A., and Mallord, J. 2004. Habitat selection by song thrushes in stable and declining farmland populations. Journal of Applied Ecology 41: 275-293.
Penczak, T., Zeieba, G., Koszalinski, H., and Kruk, A. 2003. The importance of oxbow lakes for fish recruitment in a river system. Archiv für Hydrobiologie 158(2): 267-281.

Perry, G., and Garland, T. 2002. Lizard home ranges revisited: Effects of sex, body size, diet, habitat and phylogeny. Ecology 83(7): 1870-1885.
Pessanha, A.L.M., Araujo, F.G., C., D.A.M.C., and Gomes, I.D. 2003. Diel and seasonal changes in the distribution of fish on a South East Brazil sandy beach. Marine Biology 143(6): 1047-1055.

Poff, N.L. 1997. Landscape filters and species traits: towards mechanistic understanding and prediction in stream ecology. Journal of the North American Benthological Society 16(2): 391-409.
Powell, R.A. 2000. Animal home ranges and territories and home range estimators. In Research Techniques in Animal Ecology: Controversies and Consequences. Edited by T.K. Fuller. Columbia University Press, New York. pp. 65-110.
Pretty, J.L., Harrison, S.S.C., Shepherd, D.J., Smith, C., Hildrew, A.G., and Hey, R.D. 2003. River rehabiliitation and fish populations: assessing the benefit of instream structures. Journal of Applied Ecology 40: 251-265.
Pulliam, H.R., Dunning Jr, J.B., and Liu, J. 1992. Population dynamics in a complex landscape: a case study. Ecological Applications 2: 165-177.
Raat, A.J.P. 1988. Synopsis of biological data on the northern pike Esox lucius Linaeus 1758. FAO Fisheries Synopsis No. 30(Rev 2): 178p.
Redpath, S.M. (1995) Habitat fragmentation and the individual: tawny owls Strix aluco in woodland patches. Journal of Animal Ecology, 64, 652-661.

Reyes-Gavilan, F.G., Garrido, R., Nicieza, A.G., Toledo, M.M., and Brana, F. 1996. Fish community variation along physical gradient in short streams of northern Spain and the disruptive effect of dams. Hydrobiologia 321: 155-163.
Rideout, R.M., Rose, G.A., and Burton, M.P.M. 2005. Skipped spawning in female iteroparous fishes. Fish and Fisheries 6: 50-72.
Robertson, P.A., Aebisher, N.J., Kenward, R.E., Hanski, I.K., and Williams, N.P. 1998. Simulation and jack-knifing assessment of home-range indices based on underlying trajectories. Journal of Applied Ecology 35(6): 928-940.
Rosell, R.S., and MacOscar, K.C. 2002. Movements of pike, Esox lucius, in Lower Lough Erne, determined by mark-recapture between 1994 and 2000. Fisheries Management and Ecology 9(4): 189-196.
Ross, S.T., and Baker, J.A. 1983. The response to fishes to periodic spring floods in a Southeastern stream. The American Midland Naturalist 109(1): 1-14.

Rothley, K. D. 1999. Designing bioreserve networks to satisfy multiple, conflicting demands. Ecological Applications 9: 741-750.

Rowland, W.J. 1989. The effects of body size, aggression and nuptial coloration on competition for territories in male threespine sticklebacks Gasterosteus aculeatus. Animal Behaviour 37: 282-289.

Rueda, M., and Defeo, O. 2003. Spatial structure of fish assemblages in a tropical estuarine lagoon: combining multivariate and geostatistical techniques. Journal of Experimental Marine Biology and Ecology 296(1): 93-112.
Sauer, T.M., Ben-David, M., and Bowyer, D.T. 1999. A new application of the adaptive-kernal method: Estimating linear home ranges of river otters, Lutra canadensis. Canadian Field-Naturalist 113(3): 419-424.

Schad, S., Revilla, E., Wiegand, T., Knauer, F., Kaczensky, P., Breitenmoser, U., Bufka, L., Cerveny, J., Koubek, P., Huber, T., Stanisa, C., and Terepl, L. 2002. Assessing the suitability of central European landscapes for the reintroduction of Eurasian lynx. Journal of Applied Ecology 39: 189-203.

Schaffer, W.M., and Elson, P.F. 1975. The adaptive significance of variations in the life history among local populations of Atlantic salmon in North America. Ecology 56: 577-590.
Schiemer, F. 1999. Conservation of biodiversity in floodplain rivers. Archiv für Hydrobiologie Supplementband 115(3): 423-438.
Schmutz, S., and Jungwirth, M. 1999. Fish as indicators of large river connectivity: the Danube and its tributaries. Archiv für Hydrobiologie Supplementband 115(3): 329-348.
Schulze, T., Kahl, U., Radke, R.J., and Benndorf, J. 2004. Consumption, abundance and habitat use of Anguilla anguilla in a mesotrophic reservoir. Journal of Fish Biology 65: 1543-1562.
Scott, M.T., and Nielsen, L.A. 1989. Young fish distribution in backwaters and mainchannel norders of the Kanawha River, West Virginia. Journal of Fish Biology 35(Suppl. A): 21-27.
Skov, C., Brodersen, J., Bronmark, C., Hansson, L. A., Hertonsson, P. and Nilsson, P. A. 2005. PIT tagging in cyprinids. Journal of Fish Biology. 67(5): 1195-1201.

Slipke, J.W., Sammons, S.M., and Maceina, M.J. 2005. Importance of the connectivity of backwater areas for fish productions in Demopolis Reservoir, Alabama. Journal of Freshwater Ecology 20(3): 479-485.

Sliwa, A. 2004. Home range size and social organisation of black-footed cats (Felis nigripes). Mammalian Biology 69(2): 96-107.

Sokal, R.R., and Rohlf, F.J. 1995. Biometry: the principles and practice of statistics in biological research. W. H. Freeman and Company, New York.
Sommer, T.R., Nobriga, M.L., Harrel, W.C., Batham, W., and Kimmerer, W.J. 2001. Floodplain rearing of juvenile chinook salmon: evidence of enhanced growth and survival. Canadian Journal of Fisheries and Aquatic Sciences 58: 325-333.
Starrett, W.C. 1972. Man and the Illinois River. In River ecology and man. Edited by R.T. Oglesby, C.A. Carlson and J.A. McCann. Academic Press, New York. pp. 131-167.
StatSoft.Inc. 2006. Electronic Statistics Textbook WEB: http://www.statsoft.com/textbook/stathome.html. Tulsa, OK.
Sunde, P., and Bolstad, M.S. 2004. A telemetry study of the social organization of a tawny owl (Strix aluco) population. Journal of Zoology 263(1): 65-76.
Sutherland, W.J. 1996. From individual behaviour to population ecology. Oxford University Press, Oxford.

Swihart, T.K. 1986. Home range - body-mass allometry in rabbits and hares (Leporidae). Acta Theriologica 31(1-14): 139-148.
Szacki, J. 1999. Spatially structured populations: how much do they match the classic metapopulation concept? Landscape Ecology 14: 369-379.
Taylor, W.A., and Skinner, J.D. 2003. Activity patterns, home ranges and burrow use of aardvarks (Orycteropus afer) in the Karoo. Journal of the Zoological Society of London 261: 291-297.
Ter Braak, C.J.F., and Prentice, I.C. 1988. A theory of gradient analysis. Advances in Ecological Research 18: 271-317.

Ter Braak, C.J.F., and Verdonschot, P.F.M. 1995. Canonical correspondence analysis and related multivariate methods in aquatic ecology. Aquatic Sciences 57: 1015-1621.

Terry, E.L., McLellan, B.N., and Watts, G.S. 2000. Winter habitat ecology of mountain caribou in relation to forest management. Journal of Applied Ecology 37: 589-602.
Tesch, F.W. 1977. The eel; biology and management of anguillid eels. Chapman and Hall, London.

Thorstad, E.B., Hay, C.J., Naesje, T.F., and Okland, F. 2001. Movements and habitat utilization of three cichlid species in the Zambezi River, Namibia. Ecology of Freshwater Fish 10: 238-246.

Valdez, R.A., Hoffnagle, T.L., McIvor, C.C., Ted, M., and Leibfried, W.C. 2001. Effects of a test flood on fishes of the Colorado River in Grand Canyon, Arizona. Ecological Applications 11(3): 686-700.
Vannote, R.L., Minshall, W., Cummins, C., Sedell, J., and Cushing, C. 1980. The river continuum concept. Canadian Journal of Fisheries and Aquatic Sciences 37(1): 130-137.
Van Teeffelen, A. J. A., Cabeza, A. and Moilanen, A. 2006. Connectivity, probabilities and persistence: Comparing reserve selection strategies. Biodiversity and Conservation. 15: 899-919.

Vokoun, J.C. 2003. Kernel density estimates of linear home ranges for stream fishes: Advantages and data requirements. North American Journal of Fisheries Management 23: 1020-1029.
Von Bertalanffy, L. 1938. A quantitative theory of organic growth. Human Biology 10: 181-243.

Walford, L.A. 1946. A new graphic method for describing the growth of animals. Biological Bulletin of the Marine Biological Laboratory 90: 141-147.

Walls, S.S., Manosa, S., Fuller, R.M., Hodder, K.H., and Kenward, R.E. 1999. Is early dispersal enterprise or exile? Evidence from radio-tagged buzzards. Journal of Avian Biology 30: 407-415.
Ward, J.V. 1989. The four-dimensional nature of lotic ecosystems. Journal of the North American Benthological Society 8(1): 2-8.
Ward, J.V., and Stanford, J.A. 1995. The serial discontinuity concept: extending the model to floodplain rivers. Regulated Rivers: Research \& Management 10: 159-168.
Welcomme, R.L. 1985. River Fisheries. In Technical paper 262. FAO Fisheries. WFD CIS Guidance Document No. 10. 2003. River and lakes - Typology, reference conditions and classification systems, Published by the Directorate General Environment of the European Commission, Brussels.
Wheeler, A. 1969 Fishes of the British Isles and North-West Europe. Macmillan. London. pp 613.

White, G.C., and Garrott, R.A. 1990. Analysis of wildlife radio-tracking data. Academic Press, New York, USA.

Wielgus, R. B. (2002) Minimum viable population and reserve sizes for naturally regulated grizzly bears in British Columbia. Biological Conservation. 106, 381-388.

Woodroffe, R., Donnelly, C., Cox, D.R., Bourne, F.J., Cheeseman, C.L., Delahay, R.J., Gettinby, G., McInerney, J.P., and Morrison, W.I. 2006. Effects of culling on badger Metes metes spatial organization: implications for the control of bovine tuberculosis. Journal of Applied Ecology 43: 1-10.

Wootton, R.J. 1990. Ecology of teleost fishes. Chapman and Hall, London.
Worton, B.J. 1989. Kernel methods for estimating the utilization distribution in homerange studies. Ecology 70: 164-168.

Yoder, J.M., Marschall, E.A., and Swanson, D.A. 2004. The cost of dispersal: predation as a function of movement and site familiarity in ruffed grouse. Behavioural Ecology 15(3): 469-476.
Zydlewski, G.B., Haro, A., Whalen, K.G., and McCormick, S.D. 2001. Performance of stationary and portable passive transponder detection systems for monitoring of fish movements. Journal of Fish Biology 58: 1471-1475.



[^0]:    * Spawning location is classified either as spawning observed (in all cases of river spawning location and some cases of side-channel spawning location) or as side-channel visited and spawning suspected.

