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5	Orthophosphate-P in the nutrient impacted River Taw and its catchment
6	(SW England) between 1990 and 2013
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8 9	Alan D. Tappin*, Sean Comber, Paul J. Worsfold
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12	School of Geography, Earth and Environmental Sciences, Plymouth University,
13	Drake Circus, Plymouth PL4 8AA, UK
14	
15	
16	*Corresponding author: <u>atappin@plymouth.ac.uk</u>
17	Tel +44(0)1752584572
18	Fax +44(0)1752584710

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Term	Abbreviation	Definition
Above ordnance datum	AOD	Height (in m) above mean sea level at Newlyn
Base flow index	BFI	Generally, the contribution of groundwater flow
		to river runoff as a ratio; the higher the
		contribution the higher the BFI value
Daily mean flow	DMF	Mean river flow (in $m^3 s^{-1}$) in a water-day (09.00
-		to 09.00 GMT). Typically, flows are calculated
		on the basis of measurements at 15-minute
		intervals
Extended end-member mixing	E-EMMA	The use of water quality monitoring data to
analysis		generate plots of flux of a chemical analyte
		against flow which may be used to infer and
		quantify analyte retention and/or release within a
		river catchment
Ecological quality ratio	EQR	This determines the P concentration at the lower
		class boundary for each ecological class under
		the WFD (i.e. high, good, moderate, poor)
Flux		mass area ⁻¹ time ⁻¹
Good Ecological Status	GES	The aspiration of the WFD (see below) that all
-		surface waters reach this status by 2015
Load		mass time ⁻¹
Method 3		Load estimation algorithm from Littlewood. ¹
		Defined in section 2.3
Method 5		Load estimation algorithm from Littlewood et al.
		² Defined in section 2.3
Orthophosphate-P		Generic term used in the current study to
		describe P concentrations measured for
		regulatory purposes
Phosphorus Limiting	PLC	River phosphorus concentration below which
Concentrations		algal growth is nutrient limited
Population equivalent	pe	3 1
Q ₉₅ flow		Flow (in m ³ s ⁻¹) which was equalled or exceeded
		for 95% of the flow record
Reactive phosphorus	RP	Fraction of phosphorus passing through a 0.45
		µm filter membrane and measured by the
~	~~~~	phosphomolybdenum blue colorimetric method
Sewage treatment works	STW	
Soluble reactive phosphorus	SRP	Fraction of phosphorus measured by the
		phosphomolybdenum blue colorimetric method
		after settling of suspended particles from the
		river water sample
water Framework Directive	WFD	
water Information Management	WIMS	A chemical dataset collected and compiled by the
System		Environment Agency of England

41 Abstract

Excess dissolved phosphorus (as orthophosphate-P) contributes to reduced river water quality 42 within Europe and elsewhere. This study reports results from analysis of a 23-year (1990-43 2013) water quality dataset for orthophosphate-P in the rural Taw catchment (SW England). 44 Orthophosphate-P and river flow relationships and temporal variations in orthophosphate-P 45 concentrations indicate the significant contribution of sewage (across the catchment) and 46 industrial effluent (upper R. Taw) to orthophosphate-P concentrations (up to 96 %), 47 particularly during the low flow summer months when maximum algal growth occurs. In 48 contrast, concentrations of orthophosphate-P from diffuse sources within the catchment were 49 more important (> 80 %) at highest river flows. The results from a 3 end-member mixing 50 51 model incorporating effluent, groundwater and diffuse orthophosphate-P source terms suggested that sewage and/or industrial effluent contributes ≥ 50 % of the orthophosphate-P 52 load for 27 – 48 % of the time across the catchment. The Water Framework Directive (WFD) 53 Phase 2 standards for reactive phosphorus, introduced in 2015, showed the R. Taw to be 54 generally classified as Poor to Medium Ecological Status, with a Good Status occurring more 55 frequently in the tributary rivers. Failure to achieve Good Ecological Status occurred even 56 though, since the early-2000s, riverine OP concentrations have decreased (although the 57 mechanism(s) responsible for this could not be identified). For the first time it has been 58 demonstrated that sewage and industrial effluent sources of alkalinity to the river can give 59 erroneous boundary concentrations of orthophosphate-P for WFD Ecological Status 60 classification, the extent of which is dependent on the proportion of effluent alkalinity 61 present. This is likely to be a European – wide issue which should be examined in more 62 detail. 63

Keywords: Taw, orthophosphate-P, time series, loads, phosphorus standards, water
quality.

66 **1 Introduction**

The deleterious impacts of increased anthropogenic loads of phosphorus, principally 67 as orthophosphate-P, on river water quality within Europe and elsewhere have been 68 recognised for several decades.³ Within the EU, the Urban Waste Water Treatment Directive 69 (UWWTD, 91/271/EEC) has been the main legislative driver in reducing phosphorus inputs 70 from urban centres to surface waters, largely via improved waste water collection and 71 reductions in phosphorus concentrations in sewage treatment work (STW) effluents.⁴ Across 72 the EU, decreases in riverine orthophosphate-P concentrations of 2.1 % per year on average 73 in the two decades to 2012 have been ascribed to the mitigation measures implemented under 74 the UWWTD.^{5, 6} The mean orthophosphate-P concentration in ca. 1000 EU rivers in 2012 75 was 0.06 mg P $L^{-1.5}$ In the UK, decreasing concentrations since the late 1990s – mid 2000s 76 have been reported for the R. Thames^{7, 8} and its tributaries⁸, and the Dorset R. Frome.^{9, 10} In 77 78 the R. Tamar (southwest England) decreasing concentrations with time were also evident, although reduced sampling frequency in the later stages of the time series reduced confidence 79 in this conclusion.¹¹ 80

The EU Water Framework Directive (WFD; 2000/60/EC) was promulgated with the 81 significant objective to protect, enhance and restore all bodies of surface water with the aim 82 of achieving Good Ecological Status (GES) of surface and ground waters by 2015. Despite 83 the apparent success in reducing riverine orthophosphate-P concentrations there are still 84 many rivers across the EU that are failing to achieve GES for this nutrient, including many in 85 the UK.⁴ Surface waters receiving effluents from major UK urban centres have been most 86 impacted, and it is here that most effort has been applied to reduce nutrient inputs from STWs 87 (e.g.⁷). In rural catchments with relatively low human populations, however, it has become 88 increasingly apparent that sewage effluents also make an important contribution to riverine 89 orthophosphate-P concentrations, particularly during the lower flow, spring and summer 90

months when in-river algal growth can flourish.^{8, 9, 12, 13} While tertiary nutrient stripping is
being installed in larger STWs, most STWs in rural catchments are too small for this to be
required. Furthermore, septic tanks remain integral to waste disposal for many dwellings in
rural areas, and their not insignificant contribution to nutrient loading in surface waters is
now recognised.¹⁴

The river phosphorus (termed reactive phosphorus) concentrations which define the 96 Ecological Status boundaries prescribed in the WFD Phase 2 standards introduced in 2015 97 are estimates of natural phosphorus concentrations, taking into account the alkalinity 98 99 (measured as CaCO₃) of the river water and the altitude, above sea level, that would be expected in the absence of anthropogenic pressures.¹⁵⁻¹⁶ The hypothesis is that the alkalinity 100 concentration observed is a reflection of natural rock weathering and hydrological processes. 101 In pristine areas, unperturbed by anthropogenic impacts, this is likely to be correct. However, 102 in rivers receiving sewage and industrial effluents, this hypothesis is incorrect, as these 103 effluents can contain substantial quantities of alkalinity (e.g.^{17,18}). In these rivers, inclusion of 104 sewage alkalinity will serve to increase the concentration of phosphorus defining each 105 Ecological Status boundary, effectively providing a more 'relaxed' standard. The extent of 106 this effect will depend on the relative loads of alkalinity from sub-soil sources and sewage 107 and industrial effluent in a sample; for the same effluent alkalinity load the effect will be less 108 for rivers fed from a chalk aquifer than for rivers fed by low alkalinity groundwater or rivers 109 110 of low base flow index (BFI). The effect may also be most pronounced in rivers which are dominated by effluent flows during the low river flow periods that typify the summer algal 111 growing season. 112

113 The Taw catchment in south west England is a predominantly rural environment with 114 a low population and little industry. Nevertheless, thirteen water bodies across the catchment 115 were failing GES for phosphorus under the WFD Phase 1 cycle.¹⁹ An important aim of the

116 Taw River Improvement Project, a recently completed 2.5 y £1.86 M programme designed to improve ecological status of rivers across the catchment (L. Couldrick, pers. comm.) was to 117 undertake an assessment of river water quality data for phosphorus in the catchment. This 118 study is a contribution to that aim. The specific objectives for the current study were to: (i) 119 quantify temporal trends in orthophosphate-P concentrations between 1990 and 2013, and to 120 apportion orthophosphate-P sources, and (ii) calculate Ecological Status in relation to 121 orthophosphate-P using the WFD Phase 2 standards, and then examine if the effluent 122 component of alkalinity compromises the standards for orthophosphate-P in this catchment, 123 and by implication in catchments across the EU. The second objective provides the first 124 analysis of its kind. 125

126 **2 Methods**

127 **2.1 Study area**

The Taw system is part of the North Devon catchment of the UK South West River 128 Basin District²⁰ and covers an area of 1211 km². The R. Taw rises at Taw Head on Dartmoor 129 (altitude 550 m AOD) and flows northward to join the estuary at the tidal limit at Newbridge. 130 The major tributaries are the Mole, the Lapford Yeo and the Little Dart River (Fig. 1). The R. 131 Taw (68 %) and the R. Mole (29 %) drain 97 % of the catchment (Table 1). The mean, 132 maximum and Q₉₅ flows reflect the relative size of the drainage basins, with these values 133 highest at Umberleigh, located closest to the tidal limit (Table 1). The river responds quickly 134 to rainfall, with rapid rises in river levels.²¹ The river base flow indices are mid-range (0.42 -135 0.47; Table 1), and alluvial deposits in the main river valley may contribute to river flows 136 during dry periods.²¹ Catchment geology is dominated by the Carboniferous Bude and 137 Crackington formations, while > 95% of the catchment is covered by clay and clay loam 138 soils.²² About 75% of the catchment area is used for agriculture (59% grassland, 14% arable), 139 12% woodland and forest and 9% rough grassland.²³ An estimated 77% of the grassland is 140 used for beef cattle and sheep, whilst the remainder is used for dairy cattle.²² The main areas 141 of population are Barnstaple (population 47858 in 2009), Braunton (11491), South Molton 142 (13576) and Witheridge (2262), as shown in Fig. 1. 143

144

2.2 Data resource and general approach

Daily mean river flows (DMFs; calculated from 15 min interval instantaneous flow
data over 24 h) were obtained from the Environment Agency (EA) gauging stations (GS) at
Umberleigh and Taw Bridge on the R. Taw and Woodleigh on the R. Mole (Table 1 and Fig.
1) from 1990 onwards. Chemical determinand data from 1990, collected for statutory
monitoring purposes, were also obtained from the EA WIMS for the sampling sites shown in
Fig. 1 and detailed in Supplementary Information (SI) 1. Site selection was based on EA

evidence, since 2010, of failure of river waters to meet WFD Phase 1 GES for 151 orthophosphate-P because of effluent and diffuse pollution.¹⁹ Sample record end dates 152 occurred between 2006 and 2013, depending on site (SI 1). The chemical determinands 153 considered herein are orthophosphate-P and total alkalinity. Orthophosphate-P was 154 determined in the sample supernatant following settlement of suspended particles originally 155 present in the collected water sample. As the supernatant may contain colloids and fine 156 particles, as well as non-orthophosphate 'dissolved' P, the resulting measured P 157 concentrations may represent orthophosphate-P plus P associated with these other 158 159 components that is measurable by the analytical technique used. As a result, the orthophosphate-P as it is coded in the EA database, may be variously referred to as soluble 160 reactive phosphorus (SRP) or reactive phosphorus (RP). In this paper the term 161 orthophosphate-P is used by default, although the other terms are used when appropriate. 162 For sites sampled between 1990 and 2013, water samples were collected from a 163 maximum of 4.3 % of the 8766 gauged flow days. In general, 4 % of the chemical samples 164 were collected within 7 days of each other, 15 % within 14 days and 57 % within 30 days. 165 Relatively high frequency sampling (25-35 samples y⁻¹) was undertaken between 1991 and 166 1996 at site 2; otherwise the frequency at this site and elsewhere was 10-13 samples y^{-1} . 167 Effluent data for sewage treatment works (STWs) in the catchment were provided by the EA; 168 the orthophosphate-P concentration was 5.3 mg P L^{-1} (in the centre of the range of 1 - 10 mg 169 PL^{-1} for UK sewage effluent²⁴) while dry weather flows for each STW were given as an 170 171 annual mean. The EA also provided effluent orthophosphate-P concentration and flow data for the Taw Valley Creamery, located in the upper catchment between sites 9 and 10 (see Fig. 172 1), for 2006 - 2014. There are also many septic tanks throughout the area, which can act as 173 either point or diffuse inputs for nutrients¹⁴, but there were no data on either flows or nutrient 174 concentrations available for these potential sources. 175

In section 3 the data are generally reported for the R. Taw and its tributaries 176 separately. However, at three locations (two on the R. Taw and one on the R. Mole), river 177 flow and chemical concentration data were integrated at each of these locations individually 178 to provide additional insights into orthophosphate-P behaviour; subsequently, the results from 179 these three locations are discussed as a group. For the first location, the flow data from site 3 180 were combined with the chemical data from site 2, and for the second location chemical data 181 182 for site 14 were combined with flow data for the Woodleigh GS on the R. Mole. Site 3 is ca. 4.5 km upstream of site 2, the most frequently sampled site on the R. Taw, while the 183 184 Woodleigh GS is ca. 4 km upstream of site 14. For the third location, the Taw Bridge GS is co-incident with the chemical sampling at site 7 (Fig. 1). These locations are therefore 185 referred to as site 2, site 14 and site 7. In addition to orthophosphate-P there is also a more 186 extended treatment of the alkalinity data because of the key role of this parameter in the 187 definition of river water quality standards for phosphorus within the WFD. 188

189 **2.3 Temporal trend statistics and load estimation algorithms**

Exploratory data analysis indicated that there appeared to be decreasing trends in 190 orthophosphate-P concentrations in the time series at sites 2, 7 and 14, particularly from ca. 191 2003 onwards. To quantify the potential significance of these patterns at each individual site, 192 temporal trend analyses using the Spearman's Rho, Mann-Kendall and Seasonal Kendall 193 194 tests were undertaken. These tests are non-parametric, rank-based statistics designed to reveal gradual monotonic trends in time-series data^{25,26} and are appropriate for relatively 195 coarsely-resolved data.²⁷ The theoretical basis of the trend statistics are described in Hipel 196 and McLeod and Helsel²⁵ and Hirsch.²⁸ The Seasonal Kendall test performs the Mann-197 Kendall test for individual seasons of the year, where season is defined by the user.²⁶ The 198 Mann-Kendall and Seasonal Kendall tests require the data to show constant variance through 199 time and to not show autocorrelation.^{26,28} Log₁₀ transformation of the orthophosphate-P 200

variable satisfied the first assumption, while autocorrelation function analysis showed that the 201 orthophosphate-P data were largely within the limits of acceptable autocorrelation. 202 Autocorrelation is less likely to be observed in monthly resolved data of up to 10 years 203 duration, as is the case here.^{26,27} Trend analyses were undertaken on both non flow-adjusted 204 orthophosphate-P concentrations (representing the influence of both hydrological and non-205 hydrological factors) and flow-adjusted orthophosphate-P concentrations (representing non-206 hydrological factors only), following Hirsch et al.²⁹ and Jaruskova and Liska.³⁰ Human 207 related activity may account for a proportion of the latter.³¹ For the flow-adjusted analyses. 208 209 temporal trend analyses were undertaken on residuals obtained from ordinary least squares linear regression of orthophosphate-P concentration vs flow. The data were log₁₀ transformed 210 prior to regression in order to ensure regression residuals were homoscedastic in each of the 211 three cases, although Hirsch et al.²⁹ had previously defended the use of the parametric 212 procedure by stating that it was used only to remove variance explained by the flow variable, 213 rather than using it for statistical modelling per se. The significance of all temporal trend 214 results were tested at $\alpha = 0.05$. The Spearman's Rho tests were undertaken using Sigmaplot[®] 215 11, while the Mann-Kendall and Seasonal Kendall tests were undertaken using an executable 216 file developed by, and downloaded from, the US Geological Survey.²⁶ Significance of the 217 correlation coefficients was tested at the p < 0.05 level. Autocorrelation function analyses 218 were undertaken using Minitab[®]17. 219

220

221

The estimation of orthophosphate-P loads (mass time⁻¹) reported in the current work are based on the following algorithms:

(i) the quantitation of the low (or base) flow end member load, and its division into a
 groundwater and an effluent contribution follows the extended end-member mixing analysis
 (E-EMMA):³

225

 $L_{baseflow} = L_{effluent} + L_{groundwater}$

Eqn (1)

226
$$L_{effluent} = \Sigma(C_{stw}, Q_{dwf_stw})$$
Eqn (2)227 $L_{groundwater} = C_{groundwater} \times Q_{groundwater}$ Eqn (3)228 $Q_{groundwater} = Q_{river_min} - Q_{effluent}$ Eqn (4)229 $Q_{effluent} = \Sigma Q_{dwf_stw}$ Eqn (5)

where L is determinand load in g s⁻¹, C is determinand concentration in g m⁻³, Q is flow in m³ s⁻¹, dwf_stw is sewage treatment work dry weather flow and river_min is the minimum daily mean river flow for the period of interest.

(ii) Method 5 is the favoured OSPARCOM algorithm for estimating determinand loads from
 periodic concentration and flow data; the load, L, is flow-weighted.²

$$L = \left(\frac{K\sum_{i=1}^{n}(Q_{i}C_{i})}{\sum_{i=1}^{n}Q_{i}}\right)\overline{Q}$$
 Eqn (6)

C_i is the deteminand concentration (in g m⁻³) in each of i = 1, n samples, Q_i is the corresponding daily mean river flow for that sample. K is 86400 seconds per day and the load L is g d⁻¹. \bar{Q} is the flow-weighted term and is given by:

where Q_k represents daily mean river flows for each day of the year (k = 1, N) and N is the number of days in the year.

240 (iii) Method 3 defined the load as:

$$L = K \sum_{i=1}^{n} (C_i \overline{Q}_p)$$
 Eqn (8)

241 where $\overline{Q_p}$ is the mean flow for the period between samples.¹

242 **3. Results and Discussion**

243 **3.1 Spatial and temporal variability in orthophosphate-P concentrations**

244 **3.1.1 Spatial variability**

The annual spatial and temporal variability in the concentrations (mean \pm one 245 standard deviation) of orthophosphate-P for the R. Taw are summarised in SI 2. 246 Concentrations were relatively low in the upper R. Taw at site 11 (generally $< 0.04 \text{ mg P L}^{-1}$), 247 became markedly higher downstream, 0.089 - 0.707 mg P L⁻¹ at site 9, and then decreased 248 further downstream to 0.036 - 0.186 mg P L⁻¹ at site 2, located close to the tidal limit at site 249 250 1. The lower concentrations are typical of rural sites in southwest England showing some evidence of anthropogenic impact.^{6, 10, 11} The elevated concentrations observed at sites 7, 8 251 and 9 were more typical of urban areas impacted by effluents^{32,33}, and were likely to have 252 been due to effluent discharged from the Taw Valley Creamery and STWs located on the 253 upper R. Taw (North Tawton, Belstone/South Tawton; see Fig. 1). Incorporation of effluent 254 orthophosphate-P concentration and flow data from the Creamery and the STWs into a 255 conservative dilution model for the short stretch of river reach between Belstone/South 256 Tawton and Taw Bridge (distance ca. 12 km) indicates that the Creamery effluent may have 257 been a significant contributor (up to 100 %) to the orthophosphate-P concentration observed 258 at site 7 (Taw Bridge), as shown in Fig. 2. Nevertheless, the STW (population equivalent 259 (p.e.) 2706) also was important, particularly during the summer. 260

The annual mean concentration data for orthophosphate-P in tributary rivers are given in SI 3. As for the R. Taw, mean concentrations varied by an order of magnitude across the 16 sites, and variations in concentrations about the mean were large at many sites. The relatively high concentrations at site 12, on the R. Mole, were probably due to effluent discharges from the STWs located upstream at South Molton and North Molton; the former is the second largest STW in the catchment and together they have an estimated P load of 2.75 t

 y^{-1} , equivalent to 26 % of the STW effluent P loads to rivers across the entire catchment. 267 Downstream of site 12, orthophosphate -P concentrations decreased (sites 13 and 14), 268 presumably due to dilution and/or loss of nutrient from solution. In addition, the tributaries 269 draining into the Mole upstream of site 13, at sites 15 and 16, were low in orthophosphate-P. 270 Sites 19 - 25 showed wide variability in mean concentrations. In this region there are a 271 number of STWs, with an aggregate p.e. of over 3000 and an effluent P load of ca. 1.4 t y⁻¹, 272 and it is likely that these inputs would have contributed to this variability. Lowest mean 273 concentrations occurred at site 18, and at sites 26 and 27 located in the more northerly part of 274 275 the catchment.

A decreasing concentration of any dissolved determinand with increasing river flow 276 (the type 1 response¹⁰) can be largely explained by the dilution of determinand rich effluents 277 or groundwater, or both, with determinand poor diffuse run-off from the catchment. The 278 concentration – flow relationships for orthophosphate-P at sites 2, 7 and 14 exemplify this 279 behaviour (SI 4(a)). Recasting the orthophosphate-P data as a monthly time series, shown in 280 SI 4(b), similarly reveals that the highest concentrations (and largest standard deviations in 281 concentrations) consistently occurred during the low flow, summer months. Thus, the 282 maximum mean concentrations occurred in September at site 7 (0.94 \pm 1.09 mg P L⁻¹), in 283 August at site 2 (0.13 \pm 0.09 mg P L⁻¹) and in July at site 14 (0.07 \pm 0.05 mg P L⁻¹). 284

285 **3.1.2 Temporal trends**

The results of the temporal trend analyses are reported in Table 2. While the correlation coefficients are weak to moderate (ca. -0.3 to -0.5), all three tests show that for each site there was a significant downward trend in orthophosphate-P concentrations over the sampling periods, for both the flow adjusted and non-flow adjusted data. However, the similarity in the magnitude of the correlation coefficients from analyses on the flow adjusted and non-flow adjusted concentrations indicates that hydrological variability may have played

a role. There have been changes to river water abstractions and augmentation in the Taw 292 catchment (as noted in Table 1) although it is not possible to attribute any of the changes in 293 orthophosphate-P concentrations to these factors. It has been shown that median (Q_{50}) flows 294 during the summer and autumn (June – November) for the period 1969 – 2008 increased by 295 10 - 30 % in the southwest of England, probably due to climatic factors.³⁴ In principle, 296 increased river flows during summer and autumn would, inter alia, lead to increased dilution 297 of the point source orthophosphate-P loads that tend to dominate during these periods, and 298 hence give rise to lower river orthophosphate-P concentrations. This may explain, to some 299 300 extent, why the Seasonal Kendall test returns larger positive correlations in each case than the Mann Kendall test, which doesn't explicitly account for seasonal scale changes in the data 301 (Table 2). However, this hypothetical scenario is likely to be confounded by the complexity 302 of catchment nutrient cycling, sources and losses, and their changes with time. Recent 303 decreases in orthophosphate-P concentrations observed in English rivers in the last decade or 304 so have been due to nutrient stripping of STW effluents prior to final discharge.^{7, 10, 35, 36} In 305 the Taw catchment, orthophosphate-P stripping has not been implemented under the 306 UWWTD because of the relatively small sizes of the individual STWs within the catchment, 307 and so the downward trends in orthophosphate-P concentrations cannot be ascribed to this 308 driver. Reductions in P-loading of domestic detergents may have played a role³⁷, but as 309 temporal trend data for effluent P loads from STW in the catchment were not available, this 310 suggestion is tentative. The downward trend may also be explained, in part, by changes in 311 P₂O₅ fertiliser applications over the last decade; for grassland in southwest England, 312 applications have decreased by ca. 30 % since 2002, to 7 kg ha⁻¹ P in 2013, while for all crops 313 and grass the inputs have halved over the same period to 14 kg ha⁻¹ P in 2013.^{38, 39} An 314 additional factor that could generate an apparent trend in decreasing orthophosphate-P 315 concentrations is a change to lower resolution sampling and subsequent flattening of the 316

signal.⁴⁰⁻⁴² In the current study, the apparent downward trends in concentrations observed
from the early-2000s was not accompanied by a clear change in sampling frequency, and so
the trends, if real, were due to factors that are unidentified at present. The potential
consequences for improved ecological status of these apparent reductions in orthophosphateP concentrations, in relation to phytoplankton growth and community composition, are
examined in section 3.3.

323 **3.2 Orthophosphate-P sources and loads**

324 **3.2.1 Catchment integrated diffuse sources**

325 The approximately asymptotic decrease in orthophosphate-P concentrations as flows increased, described in section 3.1.1, can provide information on the integrated diffuse (run-326 off from the surface and unsaturated zone that could include agricultural and septic tank 327 sources^{35, 43}) orthophosphate-P contribution from the catchment to the river. A quantitative 328 estimate of this contribution can be derived from the gradient of the regression of 329 orthophosphate-P load vs flow.^{43,44} The underlying assumption of this approach is that an 330 increase in orthophosphate-P load is due only to this diffuse term, while loads from other 331 important sources (groundwater and STW/industrial effluent) remain uniform regardless of 332 overall river flows.^{9, 44} We do not have the data to test this assumption however, as is outlined 333 in section 3.2.2. In the current work, concentration and flow data for all years at each site 334 have been used for the quantitation because annual sampling frequency was generally low 335 $(10-13 \text{ samples y}^{-1})$. Thus, the results reflect an integration of all the spatial and temporal 336 variation in inputs across the catchment plus any in-water orthophosphate-P gain/loss that 337 may have occurred (e.g.⁴³). In an earlier study⁴⁴ ordinary least squares regression (OLS) was 338 used to quantify the relationship between load and flow. However, in the current work a 339 robust linear regression (bisquare weight method) has been used to quantify the relationship 340 as this technique is less sensitive than OLS to heteroscedasticity in the data and the presence 341

342	of outliers. ⁴⁵ The results of the regression analyses are reported in SI 4(c) for sites 2 and 14
343	which represent the riverine outlets from the Taw and Mole sub-catchments. The
344	relationships between load and flow at both sites are significantly $(p < 0.001)$ and positively
345	linearly correlated, with R^2 values of 0.93 and 0.95, respectively. The slopes of the lines
346	indicate that the Taw and Mole sub-catchment contributions to orthophosphate-P
347	concentrations were of the order 0.035 mg P L^{-1} (0.034 – 0.036 mg P L^{-1} , 95 % CI) and 0.019
348	mg P L^{-1} (0.018 – 0.021 mg P L^{-1} , 95 % CI), respectively. In order to reveal variability around
349	these 1990-2013 integrated values, separate regressions between load and flow were
350	undertaken for site 2 for 1991 to 1996 when sampling was more frequent (25-35 samples y ⁻¹).
351	In all cases the R^2 values were strongly positive (range 0.75 - 0.94) and slope values (mean
352	0.054 mg P L ⁻¹ and range 0.029 – 0.076 mg P L ⁻¹) similar to or higher than the concentration
353	shown in SI 4(c). It is noteworthy that the reported year on year reductions in fertiliser P
354	loads have not obviously influenced the catchment integrated diffuse loads of
355	orthophosphate-P. This de-coupling, if real, may simply be a reflection of complexity in the
356	mobilisation and transfer of phosphorus at large areal scales. ⁴⁶ The diffuse catchment
357	concentrations estimated here are similar to those observed for the rural R. Tamar catchment
358	(southwest England) (range $0.025 - 0.118 \text{ mg P L}^{-1}$) but lower than those calculated for the
359	urbanised R. Thames and R. Thame catchments (southeast England) (range 0.097 – 0.298 mg
360	$P L^{-1}$). ^{11, 36}

The approximately conservative behaviour of orthophosphate-P evidenced by the positive and significant coefficients of determination in the load vs flow plots in SI 4(c) is perhaps counter-intuitive given the multitude of orthophosphate-P sources across the catchment, the downward temporal trend in river orthophosphate-P concentrations and known in situ orthophosphate-P reactivity in rivers.^{3, 43} With respect to in situ reactivity, exchange of phosphorus between river bed sediments, pore waters and overlying waters in

the upper Taw river and tributaries has been reported to be limited.⁴⁷ In addition, within-river
P cycling was unimportant relative to the source loadings of orthophosphate-P in the Dorset
R. Frome, located in southern England.⁹ A more rigorous assessment of orthophosphate-P
behaviour (i.e. conservative, non-conservative) is desirable, but this would require higher
temporal sampling resolutions (daily, weekly) than the monthly sampling adopted by the
EA.³

373 3.2.2 Sources and loads at river base flow

Equations (1) - (5), given in section 2.3, were used to quantify the low (or base) flow 374 end member loads of orthophosphate-P from groundwater and effluent. There are no direct 375 measurements of groundwater orthophosphate-P concentrations in the Taw catchment, but 376 typical concentrations of total dissolved phosphorus (orthophosphate-P + refractory inorganic 377 P + organic P) in a wide range of English groundwaters, including from the nearby Dartmoor 378 granite, are $< 0.1 \text{ mg P L}^{-1}$. ^{48, 49} Incorporation of the groundwater (0.1 mg P L⁻¹ used for 379 concentration) and STW flow and concentration data into equations (1) to (5) revealed that 380 STW effluent could theoretically contribute 96 %, 85 % and 72 % of the orthophosphate-P 381 load at site 7 (the Creamery effluent was included for this site; median P concentration is 5.5 382 mg L⁻¹), site 2 and site 14, respectively, during lowest river flows. The estimated dominance 383 of the effluent signal at site 7 is in good agreement with the results from the dilution model 384 assessment shown in Fig. 2. From this perspective, i.e. the dominance of effluent 385 orthophosphate-P at low river flows, the Taw is typical of rivers across the UK.^{10, 13, 35, 36, 43, 44} 386

387

3.2.3 A source apportionment model for orthophosphate-P

The catchment integrated diffuse source concentrations of orthophosphate-P reported in section 3.2.1 were combined with the orthophosphate-P loads from effluents and groundwater calculated in section 3.2.2 to give a source apportionment model (SAM) that is

analogous to the Type 1 E-EMMA described in ³. This three end-member load mixing model 391 can calculate the contribution of each source (effluent, groundwater, catchment) to the overall 392 orthophosphate-P load as river flow increases. As such, the SAM can be viewed as an initial, 393 pragmatic, attempt to examine the relative importance of orthophosphate-P loads under 394 contrasting flow regimes, and provides a basis for more detailed studies of orthophosphate-P 395 loads if or when higher temporal resolution orthophosphate-P concentration data become 396 available. In the SAM the same assumption was made regarding changes in load as given in 397 section 3.2.1, that only the diffuse loads increased with flow (thus for each 1 m³ s⁻¹ increase 398 in flow the diffuse load would increase by 0.035 g P s⁻¹ and 0.019 g P s⁻¹ for the Taw and 399 Mole sub-catchments, respectively). The results from the SAM are given in Fig. 3. At the 400 lowest flows, effluent loads dominate (72 - 85 %) at the outlets of the Taw and Mole sub-401 catchments (consistent with the loads estimated using equations 1 - 5), while groundwaters 402 contribute 15 - 28 %. At the highest river flows, in contrast, diffuse loads dominate (91 -403 93 %) and effluent loads are only ca. 6 % of the total. Nevertheless, effluent orthophosphate-404 P contributes \geq 50 % of load for approximately a half and a guarter of the time in the Taw 405 and Mole, respectively; which again emphasises the importance of effluent orthophosphate-P 406 during the critical low-flow, algal growing season. 407

408

3.2.4 Loads derived from EA WIMS data

Orthophosphate-P loads at each of the gauged sites were also calculated directly from river flow and EA WIMS monitoring data for the days for which orthophosphate-P data were available. Concentrations given as less than (these were << 1 % of the total dataset) were divided by 2 for the calculation.⁵⁰ The favoured OSPARCOM approach for load estimation was used² (the flow weighted Method 5, equation (6) in section 2.3). Mean and median daily load values were then calculated for each year, and the results are shown in Fig. 4(a). In each case the mean loads were similar to or larger than the median loads; the largest differences

were observed at site 7, presumably because of the occurrence of enhanced orthophosphate-P 416 concentrations at this site (SI 2). Highest mean loads were observed at site 2 and were in the 417 range $34 - 191 \text{ kg P d}^{-1}$, reflecting the relatively large size of its drainage area and hence river 418 flows. The next highest loads occurred at site 7 ($12 - 116 \text{ kg P d}^{-1}$). Loads from site 14 on the 419 R. Mole were in the range $20 - 60 \text{ kg P d}^{-1}$ and showed much less annual variation than the R. 420 Taw sites. The magnitude of the loads decreased in the latter half of the time series, a trend 421 that is consistent with the reduction in concentrations described in section 3.1.2. The trend of 422 decreasing loads is particularly marked at site 7 in the upper R. Taw and site 14 on the R. 423 424 Mole.

Bowes et al.⁴⁰ calculated loads of SRP in the high BFI (0.84) R. Frome (southern 425 England) using an algorithm assessed to be the most accurate (lowest bias) and least 426 imprecise (Method 3, equation 8 in section 2.3). Based on monthly sampling, bias in annual 427 load estimates for SRP in the R. Frome were in the range -10.6 to + 12.2 % (with one at 428 +27.9 %) relative to the 'true' load calculated from more frequently collected samples 429 (average 3.7 samples per day for one year). Load bias may be greater in the Taw because it is 430 a lower BFI (0.43) catchment, although because it also has a low population density, this may 431 not necessarily be the case.⁴¹ Load estimates for total reactive phosphorus (equivalent to 432 orthophosphate-P in the current work) using EA monitoring data (6 samples y⁻¹; May 2011 – 433 September 2012), using the Method 5 algorithm, gave a bias of +7.1 % compared with 434 higher resolution (hourly) data, in the low BFI (< 0.50) R. Leith (northwest England) 435 catchment.⁵¹ However, loads calculated during periods dominated by either low or high river 436 flows showed much poorer agreement with the high sample resolution based loads, implying, 437 inter alia, that the data record timespan used for inter-comparisons is important. Indeed, it has 438 been shown that high resolution sampling is necessary in order to quantify short term 439 variability in orthophosphate-P concentrations and hence loads.⁴⁰ From these studies it can be 440

441 concluded that the load estimates given in the current work may have a bias of up to ± 20 %. 442 Maier et al.⁵² reported the average orthophosphate-P load at Umberleigh (site 3 in the current 443 study) to be 37 t y⁻¹ for the period 1990 – 2004. Extrapolation of our estimates at site 2 to an 444 annual basis gives an average of 44 t y⁻¹ over the same period and 38 t y⁻¹ over the longer 445 period of 1990 – 2013 examined herein, both of which are within the bias error given above.

A comparison of the loads derived from the EA WIMS monitoring data and the SAM 446 was undertaken for sites 2 and 14. The EA derived mean and median loads for each year were 447 plotted against the mean river flow for the year (the latter calculated from the flow weighted 448 Method 5) and compared against the SAM calculated loads for the same flow range; the 449 results are presented in Fig. 4(b). While the SAM calculated loads increase monotonically 450 with flow (because only river flow changes in the model) they fall centrally within the ranges 451 calculated using Method 5 for both sites. This agreement indicates that the inclusion of 452 temporally variable orthophosphate-P loads into the SAM would improve the correlation 453 between these two load estimation approaches. Diffuse load data for orthophosphate-P at site 454 2 (cf. SI 4(c)) were calculated separately for the years 1991 to 1996 when sampling frequency 455 was relatively high $(25 - 35 \text{ samples y}^{-1})$, and the modelled loads recalculated for each of 456 those years (< 8 % of the data used to calculate the diffuse values were excluded as outliers). 457 Subsequent bisquare weight linear regression of EA derived median loads against SAM 458 calculated loads gave an R^2 of 0.91 and a slope of 1.23 (n = 5, 1995 excluded). Although n is 459 small, this good fit indicates that the approach used for estimating the diffuse concentration, 460 exemplified in SI 4(c), is robust and that, on an annual basis, the estimated loads calculated 461 by the two methods are not likely to be significantly different given that the bias (23 %) is of 462 a similar order to the bias in orthophosphate-P load estimations reported previously.^{40,51} 463 While this agreement is encouraging, the advantage of the SAM approach, relative to the 464

Method 5 load estimation, is the ability to separately identify the diffuse and point loadcomponents.

The load estimation approaches described in sections 3.2.1 to 3.2.4 can be 467 summarised and linked in the following way. Base flow orthophosphate-P loads from 468 groundwater and effluents revealed that at lowest river flows, effluent contributed 72 - 96 % 469 of the load at the three sites examined. Data from the base flow loads and integrated 470 catchment inputs of orthophosphate-P were combined to give a source apportionment model 471 which showed that effluent orthophosphate-P contributed at least half of the orthophosphate-472 P load for 27 - 48 % of the time. This outcome is consistent with recent reports of the 473 importance of effluent orthophosphate-P at low river flows in other English catchments 474 (e.g.¹⁰). Orthophosphate-P loads were also calculated from EA WIMS data using the 475 OSPARCOM Method 5 and compared with the loads derived from the SAM. The proximity 476 between the load estimates derived from the two approaches appeared to be of the same 477 magnitude as the bias in orthophosphate-P loads reported for other lower BFI English 478 rivers.⁵¹ It is anticipated that inclusion of more highly temporally resolved data would 479 improve the agreement between these two load estimation approaches, although there appear 480 to be a dearth of studies on the relationship between sampling frequency and orthophosphate-481 P load uncertainties in effluent impacted low BFI UK rivers. 482

483 **3.3 Water quality**

484 3.3.1 Water Framework Directive Phase 2 standards for reactive phosphorus

Site specific standards for reactive phosphorus (RP) under Phase 2 of the WFD
standard setting process were implemented in 2015.^{16, 53} In Phase 2, RP is defined as
phosphorus that will pass through a 0.45 µm pore size filter membrane and can be measured
by the phosphomolybdenum blue colorimetric method.

489 The RP standard, which is a calculated annual mean concentration, is defined by490 equation (9):

491	RP standard ($\mu g P L^{-1}$) = 10 ^{((1.0497 log_{10}(EQR) + 1.066) * (log_{10}(reference))}									
492	condition RP) – $\log_{10}(3500)$) + $\log_{10}(3500)$) Eqn (9)									
493	The RP standard is the concentration estimated for the lower class boundary of the High,									
494	Good, Moderate and Poor Ecological Status. The Ecological Status depends on the value of									
495	EQR used, where EQR is the site independent ecological quality ratio at the class									
496	boundary. ^{15, 16} The 'reference condition RP' is the RP concentration expected at near natural									
497	conditions, subject to local geology and geography. It can be estimated as:									
498	Reference condition RP = $10^{(0.454(\log_{10} alkalinity) - 0.0018(altitude) + 0.476)}$ Eqn (10)									
499	Alkalinity is the mean annual total alkalinity (mg CaCO ₃ L^{-1}) of the water (a proxy for									
500	geology and location) at a given site and the altitude is height (in m, AOD). The annual									
501	spatial and temporal variability in the concentrations (mean \pm one standard deviation) of									
502	alkalinity for the R. Taw and the tributary rivers are summarised in SI 5 and SI 6,									
503	respectively. There were no clear temporal trends in mean annual concentrations of alkalinity									
504	across the catchment, but there were marked spatial differences. On the R. Taw there was a									
505	notable increase in alkalinity concentrations in the upper catchment between site 10 and the									
506	downstream site 9. This pattern, and indeed the trends in the remainder of the R. Taw,									
507	mirrored those of orthophosphate – P. In the tributary rivers, the highest mean concentrations									
508	occurred at sites $22 - 25$ and to a lesser extent at sites $19 - 21$. It is also noteworthy that the									
509	sites with the highest mean concentrations also had the largest standard deviations about the									
510	mean concentration.									

Equations (9) and (10) have been used to retrospectively examine the compliance of
the R. Taw and its tributaries with the Phase 2 RP standards and the results are reported in

513	Fig. 5 and SI 7. For the R. Taw itself, most sites (4, 5, 6, 7, 8, 9) would be classified as
514	'Poor', while sites 2, 3 and 10 would be classified as 'Moderate' and site 10 occasionally as
515	'Good'. The ratio of the measured RP to reference condition RP can be calculated for each
516	year for each site and is a quantitation of the human impact on ambient phosphate-P
517	concentrations. ¹⁵ The ratio values reflect the range found for Ecological Status, with
518	measured orthophosphate-P concentrations at 'Poor' sites factors of 10 or more above the
519	reference condition, implying marked anthropogenic nutrient pressures at these locations. For
520	the tributary rivers the Ecological Status would be better, with a number of sites achieving
521	'Good' in some years (sites 14, 15, 16, 18, 26, 27). Nevertheless, sites 19 - 25 appear
522	particularly impacted, with a 'Poor' status dominant and ratios generally > 10.
523	While there are uncertainties in correlating RP standards with actual biological
524	impacts, ¹⁶ it is likely that concentrations of orthophosphate-P in the Taw catchment rivers,
525	particularly during summer, were generally above concentrations considered detrimental to
526	periphyton and benthic diatom communities. Figure 6 shows orthophosphate-P
527	concentrations for the summer growing season (April to September) for two sites on the Taw
528	with the lowest orthophosphate-P values; site 10 in the upper catchment and site 2 at the
529	catchment outlet (for the tributaries, only sites 18, 26 and 27 exhibited similarly low
530	concentrations). Also shown on Fig. 6 are reported Phosphorus Limiting Concentrations
531	(PLC) for some UK and US rivers below which periphyton biomass accrual rates were
532	observed to decline (summarised in ⁵⁴). It can be seen that most orthophosphate-P
533	concentrations were in the range of these PLC values, particularly in the later part of the time
534	series. Targeted experiments on the Taw itself would be required to see if a decrease in algal
535	biomass had indeed occurred. As these sites probably represent the best case scenarios it is
536	likely that the remaining sites in the Taw catchment would require more comprehensive
537	mitigation measures in order to achieve GES with respect to orthophosphate-P.

538

3.3.2 Does the use of alkalinity compromise the phosphorus standards?

Within the WFD Phase 2, the phosphorus-P standards for rivers are dependent on 539 alkalinity and altitude, as described above. In the unperturbed, or 'reference' sites of Phase 2, 540 alkalinity and altitude explain most of the variation in river RP concentrations because these 541 parameters "take into account the main sources and controls of the natural variation in soluble 542 phosphate concentrations (i.e. rock weathering)".¹⁵ Thus the reference conditions are defined 543 by an absence of anthropogenic pressures and indicate that river flows are driven by natural 544 catchment hydrology. The resulting linear regression of alkalinity/altitude with RP is used to 545 calculate the 'expected' concentration of RP at any site in the UK. This value is also used to 546 determine the RP concentrations that delineate the boundaries between the five different 547 Ecological Status classifications, as described in section 3.3.1. 548

While the Phase 2 RP standards are framed as annual mean concentrations, many 549 studies have highlighted the importance of orthophosphate-P to periphyton and benthic 550 diatom growth during the low flow, summer months.^{13, 55} From equation (1) it has been 551 established that low river flows can have contributions from both groundwater and effluents 552 from sewage treatment and industrial sources, and the relative importance of these sources to 553 orthophosphate-P at three sites in the Taw was estimated from equations (1) to (5). This 554 approach can be extended to alkalinity because sewage (and some industrial) effluent 555 contains alkalinity (principally carbonate/bicarbonate, but also borate and organic acids, and 556 orthophosphate-P itself¹⁸), and so, at low flows, riverine alkalinity will have both a 557 groundwater and an effluent component. Summary plots of alkalinity concentration vs river 558 flow and monthly variations in alkalinity concentrations for the sites 2, 7 and 14 in the Taw 559 catchment (Figs. 7(a) and (b), respectively) are consistent with this contention, with the 560 highest concentrations occurring at low flows and during the summer months. Notably, these 561 trends mirror those for orthophosphate-P shown in SI 4(a) and (b). 562

The presence of effluent alkalinity implies that the calculated "reference condition" 563 and "Ecological Status boundary" RP concentrations may be incorrect, the size of the 564 discrepancy being dependent on the proportion of effluent alkalinity. In addition, the 565 calculated RP concentration at a given boundary may be higher in effluent influenced rivers, 566 relative to groundwater dominated rivers, thereby giving a misleading impression of the 567 ecological status of the water body. Separation of the measured alkalinity at low river flows 568 into these two components, using equations (1) to (5), has been done for sites 2, 7 and 14. 569 Box 1 summarises the calculations, including the data used and underlying assumption made. 570 571 Thus, it has been estimated that ca. 43, 76 and 26 %, respectively, of the measured river alkalinity at these sites was from effluent. Reference condition RP and High/Good RP 572 boundaries were then calculated with and without this effluent component for the 573 ecologically important summer period of April to September (Fig. 8). The measured : 574 predicted RP ratio increased by 29 %, 89 % and 15 % at sites 2, 7 and 14, respectively, while 575 the reference condition RP concentrations decreased by the same margins. The associated RP 576 concentrations delineating the High/Good boundary fell by 26 %, 78 % and 13 %, 577 respectively, at these sites. 578

The significance of these results with respect to nutrient reduction strategies, at this 579 time, should be placed in the context that the Phase 2 standards are for annual mean 580 concentrations of both alkalinity and RP. For example, using the data in Box 1, it is possible 581 to apportion river loads of alkalinity from the different sources (effluent, groundwater, 582 diffuse) against changes in river flow (cf. orthophosphate-P in Fig. 3). The results, shown in 583 Fig. 9, suggest that both effluent and groundwater alkalinity are important at low flows, 584 consistent with the calculations above. Thus, for flows $\leq Q_{95}$ value, effluents may account for 585 25 - 50 % of the alkalinity load for the sites representing the outlets of the Taw and Mole 586 catchments. Nevertheless, because of the paucity of data on flows and concentrations used in 587

- this analysis, as noted in footnotes d and e to Box 1, these results probably provide only a
- 589 first order assessment of the importance of alkalinity sources under contrasting river flows,
- and in particular at low flows. Additional work using more constrained datasets would serve
- 591 to reduce the uncertainties and arguably contribute to a more refined set of phosphorus
- standards for inclusion in a Phase 3 cycle of standards revisions under the WFD.

593

4. Summary and conclusions

The results from a source apportionment model incorporating effluent, groundwater 594 and diffuse loads of orthophosphate-P suggested that effluent discharged to the rivers in the 595 Taw catchment contributes approximately half of the orthophosphate-P load for up to half of 596 the time across the catchment. However, during the more biologically important summer 597 months, significant contributions of sewage (across the catchment) and industrial effluent 598 (upper R. Taw) to orthophosphate-P concentrations (up to 96 %) occur. These sources 599 probably contribute to the generally Poor to Moderate Ecological Status of the rivers with 600 respect to the recently introduced WFD Phase 2 standards for phosphorus. Since the early-601 602 2000s, orthophosphate-P concentrations appear to have decreased, for reasons not identified 603 herein, with no clear improvement in overall ecological status. To compound the failure to meet Good Ecological Status at most sites sampled, it has been demonstrated that sewage and 604 industrial effluent sources of alkalinity to the river can give erroneous boundary 605 concentrations of reactive phosphorus for WFD Ecological Status classification, and in effect 606 relax the standards. 607

While the sampling resolution of the EA chemical monitoring programme, 608 particularly in the temporal dimension, frequently attracts criticism, the resulting data 609 archived in the WIMS database, at a national level, are an important resource that can be 610 611 used, with due diligence, to assess and address problems in catchment management. In the current work, the high orthophosphate-P concentrations observed in the upper R. Taw were 612 largely due to industrial effluent from a single plant, coupled with more minor contributions 613 from STWs, while at the catchment scale, the enhanced summer concentrations of 614 orthophosphate-P were due to STW effluents. These point sources appeared to be the major 615 cause of failure to meet GES under the WFD. Reductions in orthophosphate-P loads from 616 effluents could be achieved via chemical precipitation using metal chloride, although the 617

costs may be prohibitive. In the future, the development of an EU-wide Phosphorus Circular 618 Economy to severely restrict P losses to the environment, as advocated by the European 619 Sustainable Phosphorus Platform, may drive through these mitigation measures. The 620 estimated diffuse catchment runoff concentrations of ca. 0.03 mg L⁻¹ orthophosphate-P, if 621 maintained at this level, would alone cause many sites to fail GES under the WFD Phase 2 622 standards. There is no requirement under the current UWWTD to further reduce 623 orthophosphate-P loads from STW and industrial effluents in the Taw catchment and so 624 unless driven by the WFD, it is unlikely that most rivers in the Taw catchment will achieve 625 626 GES in the foreseeable future.

For catchments like the Taw, with an extensive sampling history, both spatially and 627 temporally, it is arguable that the regulatory agencies should investigate alternative sampling 628 strategies that would provide more useful data for management / policy purposes while at the 629 same time be resource neutral and still fulfil statutory monitoring obligations. For example, 630 the number of sites regularly sampled could be reduced to those with known issues plus sites 631 located at the catchment outlets, including sub-catchments, where sampling frequency could 632 be markedly increased. The latter approach would provide improved estimates of catchment 633 nutrient loads, as defined in this study, and at the same time allow more accurate and precise 634 land to sea flux estimates of contaminants required by OSPAR. Reduced routine sample 635 throughput could also allow a more comprehensive set of analyses per sample, of particular 636 importance for phosphorus because it occurs in a variety of inorganic and organic fractions 637 with varying bioavailabilities. With anticipated changes in rainfall patterns, in particular 638 increases in the frequency of short sharp summer rain events, it may be prudent to undertake 639 "smart sampling" during these events (using a combination of meteorological predictions for 640 summer rainfall with in situ measurement technologies) in order to improve understanding of 641 nutrient transfers during intermittent wetting up of catchments. The aspiration for in situ 642

measurements, inter alia, is that they should measure the bioavailable P as defined by the 643 Environmental Quality Standard that is current at the time; while in situ measurements of 644 (dissolved) reactive phosphorus in rivers are now being more frequently reported, it is evident 645 that the instrumentation requires further development, particularly in relation to the more 646 specific definition of filtered reactive phosphorus given in the WFD Phase 2 standard. 647 The contribution of sewage effluent to riverine concentrations of alkalinity appeared 648 significant in the Taw, particularly during the key summer months. Given the extent of 649 650 urbanisation within Europe it would appear prudent to extend this analysis in order to properly assess the contributions of effluent alkalinity to river alkalinity across the EU. This 651

would no doubt provide a more nuanced derivation of river phosphorus standards in a future

653 WFD river basin management cycle.

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Gauging	NRFA station	% of total catchment	Comments	Baseflow		Daily flo (m ³	ow data s ⁻¹)	
station	#	drained		index	Mean	Min	Max	Q95
Umberleigh (Taw)	50001	68	Significant modification to flows owing to public water supply abstraction. Augmentation from the Exe catchment at low flows stopped end 2002.	0.43	18.0	0.20	364	1.23
Taw Bridge (Taw)	50007	6	Water abstractions at Taw Marsh ceased ~ 1999. Cheese factory at North Taw bridge abstracts from borehole but also compensates into river at low flow.	0.47	1.80	0.02	51.1	0.17
Woodleigh (Mole)	50006	29	Low flows moderately affected by public water supply abstraction and augmentation from Exe- Taw transfers.	0.47	8.79	0.20	143	0.87

Table 1. River flow gauging stations included in this study.

773 Data from the National River Flow Archive (NRFA; http://www.ceh.ac.uk/data/nrfa/)

	Orthophosphate-P (flow adjusted concentrations)								
Site (n)	r _s	Probrs	trend	τ	Prob _{MK}	trend	τ	Prob _{SK}	trend
2	0.22	< 0.001		0.22	< 0.001	I	0.25	< 0.001	I
(126)	-0.32	< 0.001	\downarrow	-0.22	< 0.001	\downarrow	-0.35	< 0.001	\downarrow
7	0.42	< 0.001		0.20	< 0.001	I	0.22	< 0.001	I
(129)	-0.42	< 0.001	\downarrow	-0.28	< 0.001	\checkmark	-0.32	< 0.001	\checkmark
14	0.27	.0.001		0.22	.0.001	I	-0.30	0.007	I
(83)	-0.37	< 0.001	\downarrow	-0.22	< 0.001	\downarrow			\downarrow
			Orthopho	sphate-P (r	ion-flow adji	isted conce	entrations)		
	r _s	Prob r _s	trend	τ	Prob _{MK}	trend	τ	Prob _{SK}	trend
2	0.00	0.001		0.00	0.001	I	0.26	0.001	1
(126)	-0.32	< 0.001	\downarrow	-0.22	2 < 0.001	0.001 ↓	-0.36	< 0.001	\downarrow
7	0.05	0.001		0.00	0.001	I	0.05	0.001	1
(129)	-0.35	< 0.001	\downarrow	-0.23	< 0.001	\downarrow	↓ -0.37	< 0.001	\downarrow
14	0.40	0.001		0.04	0.001	I	0.45	0.001	1
(83)	-0.40	< 0.001	\downarrow	-0.26	< 0.001	\downarrow	-0.47	< 0.001	\downarrow

Table 2. Temporal trend analysis of orthophosphate-P concentrations for sites 2, 7 (2003 – 2013) and 14 (2003 – 2009).

n, number of data points; r_s , Spearman's Rho correlation coefficient; τ , Kendall's tau correlation coefficient; prob, p value of the significance of the trend; MK, Mann Kendall; SK, Seasonal Kendall

Box 1. Estimation of the contribution of effluents, groundwater and diffuse catchment sources to river alkalinity loads at sites 2, 7 and 14 in the Taw catchment.

Effluent and groundwater sources										
Calculations: the approach uses equations $(1) - (5)$ given in the main text										
ComponentUnitSite 2Site 7Site 14										
Qdwf_stw ^a	$m^{3} s^{-1}$	0.067	0.031	0.018						
Qriver_min ^b	$m^{3} s^{-1}$	0.692	0.103	0.399						
Q _{groundwater} ^c	$m^{3} s^{-1}$	0.625	0.072	0.381						
Alkalinity _{stw} ^d	g CaCO ₃ m ⁻³	51.4	51.4	51.4						
Alkalinitygroundwater ^e	g CaCO ₃ m ⁻³	7.1	7.1	7.1						
Alkalinity_load _{effluent}	$g s^{-1}$	3.42	1.59	0.95						
Alkalinity_load _{groundwater}	$g s^{-1}$	4.46	0.51	2.71						
% Alkalinity from effluent		43.4	75.6	25.8						

^a Site 2 = Taw creamery + 13 STW; Site 7 = Taw creamery + 2 STW; Site 14 = 3 STW. Dry Weather Flow (dwf) data from the EA.

^b Flow values are from the period coincident with the chemical data.

^c $Q_{groundwater} = Q_{river_min} - \Sigma Q_{dwf_stw}$. ^d Value from Beaford STW on the nearby R. Torridge. There are no alkalinity data for Taw STWs, but the range for South West River Basin District STWs is 14 - 444 mg CaCO₃ L⁻¹. Data from the EA. ^e Median value for nearby Dartmoor groundwater (range $4.1 - 34.4 \text{ mg CaCO}_3 \text{ L}^{-1}$; from ⁴⁸). No data for

Taw catchment.

Diffuse catchment source

Requirement: the key requirement for this approach is that alkalinity should be conservative in the river. Using bivariate plots analogous to those given in SI 4(c) for orthophosphate-P it can be seen that alkalinity is approximately conservative, as shown in the three figures below.





Figure 1. Left panel: place and river names; right panel: sampling sites (number) and gauging stations (U = Umberleigh, TB = Taw Bridge, W = Woodleigh). The baseline Ecological Status classification of 2009²⁰ is also shown in the right panel. **REQUIRES MINOR EDITS**

Figure 2. Estimated concentrations of orthophosphate-P in the river at site 7 (Taw Bridge), between October 2006 and March 2014, due to effluent inputs from the Taw Valley Creamery and the sewage treatment works located on the upper R. Taw. Also shown are the measured concentrations of orthophosphate-P from the EA WIMS dataset for the same period. For the estimated concentrations, orthophosphate-P loads (g P d⁻¹, derived from effluent flow and concentration data) in the effluents were diluted in to the river flow (m³ d⁻¹) gauged for that day and conservative behaviour of orthophosphate-P assumed.

Figure 3. Calculated orthophosphate-P load from effluent, groundwater and diffuse run-off (%) vs river flow $(m^3 s^{-1})$ at (a) site 2 on the R. Taw and (b) site 14 on the R. Mole. Note changes in scales.

Figure 4. (a) Orthophosphate-P load (kg P d⁻¹) for each year and (b) orthophosphate-P load (kg P d⁻¹) vs annual mean flow (m³ s⁻¹) for each year at three sites on the R. Taw and R. Mole. Note changes in scales. Thin solid line, Method 5 mean value; dashed line, Method 5 median value; thick solid line, source apportionment model value.

Figure 5. Water Framework Directive (phase 2) reactive phosphorus standards for the R. Taw. Solid line, poor/moderate boundary; dashed line, moderate/good boundary; dotted line, good/high boundary (all μ g P L⁻¹). Observed mean annual concentrations of orthophosphate-P (•; μ g P L⁻¹). Vertical bars show the ratio of the observed and predicted reactive phosphorus concentrations.

Figure 6. Orthophosphate-P (mg P L⁻¹) concentrations at site 10 and site 2 on the R. Taw during the April to September algal growing period. The horizontal lines show the Phosphorus Limiting Concentrations (PLC), reported for different UK and US rivers, below which P was the limiting nutrient and periphyton growth declined. The dashed line shows the concentration below which a positive change in diatom community composition was observed. See Bowes et al.⁵⁴ for a more detailed account of the PLC data.

Figure 7. (a) Concentrations of total alkalinity (mg CaCO₃ L⁻¹) vs river flow (m³ s⁻¹) for all years at gauged sites on the R. Taw and R. Mole. (b) Monthly mean (± 1 standard deviation) concentrations of total alkalinity (mg CaCO₃ L⁻¹) for the same sites. Note changes in concentration scales.

Figure 8. Right panel: Bar chart shows the % contribution of effluent to the measured river alkalinity at each site and the % increase in the measured : predicted reactive phosphorus (RP) ratio after subtraction of the effluent component of the measured alkalinity. Left panel: Solid lines show the decrease in the High/Good boundary concentration of RP, while the circle/dashed lines show the decrease in the predicted RP concentration, after subtraction of the effluent component of the summer period (April – September) only.

Figure 9. Estimated contribution to river alkalinity load from effluent, groundwater and diffuse run-off (%) *vs* river flow ($m^3 s^{-1}$) at three sites on the R. Taw and R. Mole. The Q₉₅ flow values are from Table 1.

River (tributary)	Sampling site	Data years	NGR	Altitude (m) ^a
Taw	(1) New Bridge	1990-1996	SS5699028280	6
Taw	(2) Chapelton F'bridge	1990-2013	SS5822026100	11
Taw	(3) Umberleigh	1990-2013	SS6079923682	19
Taw	(4) Newnham Bridge	1990-2013	SS6603017320	47
Taw	(5) Kersham Bridge	1990-2006	SS6620713554	52
Taw	(6) Chenson	1990-2013	SS7021509533	68
Taw	(7) Taw Bridge	1990-2013	SS6729406589	89
Taw	(8) Bondleigh	1990-2006	SS6578104549	100
Taw	(9) Yeo Farm	1990-2006	SS6512602871	121
Taw	(10) Rowden Moor	1990-2006	SX6550299476	128
Taw	(11) Sticklepath	1990-2006	SX6435794030	195
Mole	(12) S Molton d/s STW	1990-2006	SS7227425547	102
Mole	(13) New Bridge	1990-2012	SS7248022570	77
Mole	(14) Head Barton	1990-2009	SS6674018270	43
Mole (Crooked Oak)	(15) Alswear	1990-2010	SS7247022280	137
Mole (Little Silver Stream)	(16) Alswear	1990-2012	SS7234822104	81
Little Dart	(17) Dart Bridge	2005-2009	SS6690613720	59
L. Dart (Huntacott Water)	(18) u/s Chulmleigh	1990-2010	SS6968113830	72
Lapford Yeo (Dalch)	(19) Canns Mill Bridge	1990-2012	SS7851010490	121
Lapford Yeo (Dalch)	(20) u/s Lapford STW	1991-2006	SS7362907623	80
Lapford Yeo (Dalch)	(21) u/s Lapford Yeo	1991-2006	SS7356407425	92
Lapford Yeo	(22) Bow Bridge	1990-2006	SS7173001740	102
Lapford Yeo	(23) Bury Bridge	1990-2010	SS7377806793	81
Lapford Yeo	(24) Nymet Bridge	1990-2006	SS7141709269	80
Lapford Yeo (Ash Brook)	(25) A377 bridge	1990-2006	SS7369006670	84
Barnstaple Yeo	(26) Collard Bridge	1990-2012	SS5955235665	39
Barnstaple Yeo	(27) Brockham Bridge	1900-2012	SS6034740817	137

Supplementary Information 1. Sampling site, data availability, location and altitude.

^a Altitudes from gridreferencefinder.com. d/s, downstream; u/s, upstream

Supplementary Information 2. Annual mean (± 1 standard deviation) concentrations of orthophosphate-P (mg P L⁻¹) in the R. Taw (site locations are shown in Fig. 1). Note changes in concentration scales. Concentrations at site 11 were generally less than the limit of detection (0.04 mg P L⁻¹) and are not included.

Supplementary Information 3. Annual mean (± 1 standard deviation) concentrations of orthophosphate-P (mg P L⁻¹) in the Taw tributary rivers (site locations are shown in Fig. 1). Note changes in concentration scales.

Supplementary Information 4. (a) Concentrations of orthophosphate-P (mg P L⁻¹) vs river flow (m³ s⁻¹) for all years (1990 – 2013) at gauged sites on the R. Taw and R. Mole (site locations are shown in Fig. 1). (b) Monthly mean (\pm 1 standard deviation) concentrations of orthophosphate-P (mg P L⁻¹) for the same sites. (c) Orthophosphate-P load (g P s⁻¹) vs river flow (m³ s⁻¹) for all years for the sites representing the outlets of the two rivers. Solid and dashed lines are the bisquare weight linear regression best fit and 95 % CI, respectively. Note changes in scales.

Supplementary Information 5. Annual mean (± 1 standard deviation) concentrations of total alkalinity (mg CaCO₃ L⁻¹) in the R. Taw (site locations are shown in Fig. 1)

Supplementary Information 6. Annual mean (± 1 standard deviation) concentrations of total alkalinity (mg CaCO₃ L⁻¹) in Taw tributary rivers (site locations are shown in Fig. 1).

Supplementary Information 7. Water Framework Directive (phase 2) reactive phosphorus standards for Taw tributary rivers. Solid line, poor/moderate boundary; dashed line, moderate/good boundary; dotted line, good/high boundary (all μ g P L⁻¹). Mean annual concentrations of orthophosphate-P (•; μ g P L⁻¹). Vertical bars show the ratio of the observed and predicted reactive phosphorus concentrations. Site locations are shown in Fig. 1.