

Potential vectors of equine arboviruses in the UK

G. E. Chapman, D. Archer, S. Torr, T. Solomon, M. Baylis

There is growing concern about the increasing risk of disease outbreaks caused by arthropod-borne viruses (arboviruses) in both human beings and animals. There are several mosquito-borne viral diseases that cause varying levels of morbidity and mortality in horses and that can have substantial welfare and economic ramifications. While none has been recorded in the UK, vector species for some of these viruses are present, suggesting that UK equines may be at risk. The authors undertook, therefore, the first study of mosquito species on equine premises in the UK. Mosquito magnet traps and red-box traps were used to sample adults, and larvae were collected from water sources such as tyres, buckets, ditches and pools. Several species that are known to be capable of transmitting important equine infectious arboviruses were trapped. The most abundant, with a maximum catch of 173 in 72 hours, was *Ochlerotatus detritus*, a competent vector of some flaviviruses; the highest densities were found near saltmarsh habitats. The most widespread species, recorded at >75 per cent of sites, was *Culiseta annulata*. This study demonstrates that potential mosquito vectors of arboviruses, including those known to be capable of infecting horses, are present and may be abundant on equine premises in the UK.

Globally, there is increasing concern over emerging infectious diseases, particularly arthropod-borne viruses (arboviruses) affecting human beings and livestock (Kilpatrick and Randolph 2012, Durand and others 2013). Examples from the UK include bluetongue and Schmallenberg viruses in ruminants. The introduction of West Nile virus (WNV) into North America demonstrated the effects of mosquito-borne disease on a naive host population, both human and equine, and concerns have also been raised over the potential for spread of other mosquito-borne arboviruses affecting horses (Brown and others 2008, Pages and others 2009, Durand and others 2013). Mosquito-borne arboviruses affecting horses include WNV, Japanese encephalitis virus (JEV), Eastern equine encephalitis virus (EEEV), Western equine encephalitis virus (WEEV), Venezuelan equine encephalitis virus (VEEV), Ross River virus (RRV), Murray Valley encephalitis virus (MVEV) and Getah virus (Table 1).

Further knowledge about potential vector mosquitoes in the UK and their ability to spread arboviruses will play a key role in control and surveillance of disease in the event of an outbreak. Climate change may increase the risk of emergence of arboviral diseases by several mechanisms. Higher temperatures increase the ability of vectors to transmit viruses (Guis and others 2012,

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MacKenzie-Impoinvil and others 2015) and also have the potential to increase the geographical range of mosquitoes (Elbers and others 2015). Increased winter rainfall may also contribute to increase in mosquito populations due to creation of more temporary freshwater sites for breeding and greater abundance of emerging mosquitoes in spring (Vardoulakis and Heaviside 2012).

There are 34 species of mosquito in the UK (Medlock and Vaux 2011) and species that are implicated as vectors of arboviruses of horses elsewhere in the world include Aedes cinereus, Aedes vexans, Anopheles maculipennis s.l., Coquillettidia richiardii, Culex pipiens, Culex modestus, Culiseta morsitans, Ochlerotatus caspius, Ochlerotatus dorsalis and Ochlerotatus flavescens and a number of these are widely distributed and locally abundant across the UK (Table 2). In addition, some mosquito species present in the UK have been shown in the laboratory to be competent vectors of at least one of these viruses including Ochlerotatus punctor, Oc. detritus, Cx. modestus, Ae. vexans, Cx. pipiens s.l. and Anopheles plumbeus (Table 2).

In the UK, there have been recent and ongoing sampling and surveillance of mosquito species (Snow and Medlock 2008, Medlock and Vaux 2013, 2014, 2015, Vaux and Medlock 2015, Vaux and others 2015); however, there has been no sampling of mosquito species with specific focus on the equine host. Accordingly, the authors carried out a survey of the mosquitoes present at 32 premises across England (see Fig 1 for approximate locations) to obtain baseline data on the species composition and abundance of mosquitoes that may interact readily with equines. The authors' results identify which species may play an important role in outbreaks of mosquito-borne equine viruses in the UK and hence contribute to the development of national strategies to monitor and manage this risk.

Methods

A total of 32 sites were sampled—8 equine premises in each of northwest, northeast, southeast and southwest regions in

TABLE 1: Mosquito	-borne viruses affect	ting horses and	known morbidity	and morta	ality information			
	Virus							Getah
	JEV	WNV	EEEV	WEEV	VEEV	MVEV	RRV	virus
Inapparent infections common	Yes ⁹	Yes ⁷	Yes ²	Yes	No ⁹	Yes ¹³	Yes ¹²	Yes
Morbidity	0.03–1.4% of horses in a region ³	1 in 11-12 infections ⁷	61% ¹ of horses on some farms	low	10% of regional population (estimated) ^{10,11}	Low	Low	Unknown
Case mortality Vaccination available	5-40% ^{4,5,6} Y	38–57% ⁷ UK licensed	Up to 73% ¹ Y	20-30% ⁸ Y	40-90% ^{10,11} Y	Low	Low	Not fatal Y

Y—available in affected countries

¹Silva and others (2011)

²Pauvolid-Corrêa and others (2010)

³Spickler (2010)

⁴Ellis and others (2000)

⁵Hale and Witherington (1953)

⁶Nakamura (1972)

⁷Sellon and Long (2013)

⁸Long and Gibbs (2007)

⁹Rico-Hesse (2000)

¹⁰Sudia and others (1975)

¹¹Zehmer and others (1974)

¹²Vale and others (1991) ¹³Holmes and others (2012)

EEEV, Eastern equine encephalitis virus; JEV, Japanese encephalitis virus; MVEV, Murray Valley encephalitis virus; RRV, Ross River virus; VEEV, Venezuelan equine encephalitis virus; WEEV, Western equine encephalitis virus; WNV, West Nile virus

England (Fig 1). A stratified sampling approach was used due to the fact that many mosquito species have a patchy population distribution. In this case, four types of mosquito breeding habitat were identified: land associated with drainage ditches (drained farmland) or fenland (site 29), woodland, urban and salt marsh (Hutchinson and others 2007). The authors aimed to recruit two equine premises in each of the four habitats in each region (32 premises in total).

An internet search was conducted using Google Maps and The Phone Book from British Telecom, using the search terms 'Riding Schools', 'Livery', 'Stables', 'Stud'. BHS Riding Schools and Livery Yard Lists and the British Equestrian Directory and Newmarket Trainers Association lists were also used. This produced a list of businesses with publicly available contact details.

For each premises, the local area was investigated for potential mosquito habitats using Magic (www.magic.gov.uk) and Google Earth. Sites were assigned a category based on habitat (some sites qualified for two categories) and were graded based on area of presumed habitat and proximity of habitat to the premises. The authors aimed to locate premises within suitable habitats or, if that was not possible, within 500 m (woodland and urban sites), 1 km (urban and drained farmland sites) or 3 km (salt marsh). A maximum distance of 500 m for woodland sites was selected reflecting the relative ease of finding sites close to woodland. For salt marsh or grazing marsh, it was not possible to find sites in close proximity in many cases, but species associated with floodwater, such as Ae. vexans and coastal salt marsh such as Oc. detritus, tend to have greater dispersal capacity and Oc. detritus is capable of flying at least 2.5 miles (Service 1969, 1971b, Snow and Medlock 2008, Becker and others 2010, Verdonschot and Besse-Lototskaya 2014). In order to try and include all four habitat types within reasonable travelling distance, the four areas within the regions were chosen as follows: Wirral peninsula and Chester (northwest); between Scunthorpe, Gainsborough, Doncaster and Goole (northeast); within 20 miles of Exeter (southwest); a transect from Newmarket to the Wash (southeast).

Premises were recruited by sending out either a letter or email to the business, and following up with a telephone call. For sites where there was no response or a negative response, correspondence was then sent to a number of alternative second-choice sites, for that category of habitat, until 32 sites (eight in each of four regions) were recruited.

Mosquito sampling

Host-seeking adults

Each of the 32 sites was visited three times throughout the summer of 2015, and mosquitoes were trapped continuously for three days. Timing of visits was based on what is presently known about peaks in adult mosquito numbers of different species in the UK, visiting each of four regions within each of three seasonal peaks of mosquito activity in the months of May, late June–early July and September (Service 1969, 1977, Medlock and others 2007, Snow and Medlock 2008, Becker and others 2010, Medlock and Vaux 2015).

Trapping at each site was carried out using a mosquito magnet, Independence model (Woodstream Europe). The mosquito magnet is designed to catch host-seeking mosquitoes by using propane as a fuel source to produce heat, moisture and carbon dioxide. The trap was also baited with 1-octen-3-ol (as supplied by the trap manufacturer). The mosquito magnet trap was run continuously for ~72 hours starting in the morning and a data logger was placed underneath the body of the trap to record the environmental temperature and relative humidity for this time period. The traps were emptied as time allowed, so that the samples were not in the trap for more than 48 hours, to reduce damage, and the subsamples were combined and identified as the full 72-hour catch.

Attempts were made to catch mosquitoes landing on hosts in order to confirm horse biting. These attempts were made in the afternoon and wherever possible at dusk (Service 1969, 1971a). To fit in with the trapping schedule, four sites in each area were sampled in June/July and September in the mid-late afternoon, and four sites around dusk. Therefore, one site per day was sampled in the afternoon and one in the evening for Monday to Thursday of the trapping week in each sampling area. For each sampling effort, a group of horses was observed for 15 minutes to see whether any mosquitoes could be identified landing on them. If no mosquitoes were observed, then another group of horses was observed for 15 minutes. Group sizes ranged from 1 to 10 as horses were in their normal grazing environment (with the exception of site 25 where sampling was attempted in the stable as there was no grazing). If no mosquitoes were observed on two groups, the attempt was abandoned. If mosquitoes were observed, landings were counted for 2 minutes and then mosquitoes sampled from the head and neck of the horse (for reasons of safety) for 30 minutes to allow for species identification. Some premises could not be sampled at dusk due to access restrictions, so were only sampled in the

Species	Host biting ^{5,7}	Evidence of equine biting	Vector status
Aedes cinereus/Aedes geminus	M ^{31, 32} B ^{31, 32}	Morocco ¹ , Switzerland ³²	EEEV (I) ¹⁸
Aedes vexans	$M^{31, 32}, B^{31, 32}$	France ² , Switzerland ³²	WNV (I) ⁵ , EEV (I, L*) ^{18,19,20,21,22,}
Anopheles algeriensis	M		· · · · · · · ·
Anopheles claviger	M ³²	Switzerland ³²	
Anopheles maculipennis s.l.	M ^{31,32} , B ³¹	UK ^{4,8} , Switzerland ³²	WNV (I) ⁵
Anopheles plumbeus	M ³¹	France ²	WNV (L) ¹⁴
Coquillettidia richiardii	$M^{31,32}$, B^{32}	France ² , Switzerland ³²	WNV (I) ⁵
Culiseta alaskaensis	M		
Culiseta annulata	M ^{14,32} , B ³²	UK ³ , France ² , Switzerland ³²	WNV (L) ³³
Culiseta fumipennis	В		
Culiseta litorea	M, B		
Culiseta longiareolata	В		
Culiseta morsitans	M ³¹ , B ³¹		EEV (Z) ^{17, 19}
Culiseta subochrea	M ²	France ²	
Culex europaeus	A, R, B		
Culex modestus	M^2 , B^2	France ²	WNV (V, L) ^{2.5}
Culex pipiens s.l.	$M^{31} B^{31,32}$	France ²	WNV $(V, L)^{23, 27}$, JEV $(L^{+})^{23}$, EEV $(N)^{26}$, WEEV $(N)^{24,25}$, VEEV $(N)^{27}$
Culex torrentium	B ³¹ , M ³¹		· · · · · · · · · · · · · · · · · · ·
Ochlerotatus annulipes	M ^{9,31} , B	France ²	
Ochlerotatus cantans	M ³¹ , B ³²	UK ⁹ , Switzerland ³²	
Ochlerotatus caspius	M^2 , B^2	UK ³ , France ²	WNV (I, L†) ^{2,5}
Ochlerotatus communis	M ³¹		
Ochlerotatus detritus	M², B	UK³, France²	WNV (L) ¹⁶ , JEV (L) ¹³
Ochlerotatus dorsalis	M ⁶	UK ⁶	WEEV (I, L) ^{28,30}
Ochlerotatus flavescens	M ^{11,12}	Denmark, Canada ^{11, 12}	
Ochlerotatus geniculatus	M ^{2,31}	France ²	
Ochlerotatus leucomelas			
Ochlerotatus punctor	М ¹⁰ , В	UK ¹⁰	WNV (L) ¹⁴
Ochlerotatus rusticus	M ^{31, 32} , B	Switzerland ³²	
Ochlerotatus sticticus	M ^{31, 32} , B ³¹	Switzerland ³²	
Orthopodomyia pulcripalpis	В		

A, amphibians; B, birds; EEEV, Eastern equine encephalitis virus; I, implicated in disease transmission worldwide; JEV, Japanese encephalitis virus; L, laboratory-competent vector; M, mammals; N, non-competent as laboratory vector; R, reptiles; V, ecologically significant bridge vector worldwide; VEEV, Venezuelan equine encephalitis virus; WEEV, Western equine encephalitis virus; WNV, West Nile virus; Z, ecologically significant enzootic vector worldwide

afternoon. In order to trap mosquitoes feeding on horses, a mechanical pooter (Watkins and Doncaster) was modified with an elongated inlet tube and was muffled so as to avoid startling the horse. Individual horse behaviour was discussed with the yard owner in advance, and permission to attempt landing catches with each horse or group of horses was obtained.

Resting adults

The resting box trap was a $40\times30\times20\,\mathrm{cm}$ black box (Morris 1981), painted red inside (red-box trap) and was designed to aid in the capture of blood-fed mosquitoes (Fig 2). It was set in an open area facing west and was emptied on two mornings (either at 24 and 72 hours after deployment or 48 and 72 hours) by

Species in bold were sampled during the present study *Variable laboratory competence in a number of studies †Relatively inefficient laboratory vector

¹Faraj and others (2009) ²Balenghien and others (2006)

³Pilot work for this study—site 8, 2014

⁴Danabalan (2010)

⁵Medlock and others (2005)

⁶Service (1971a)

⁷Becker and others (2010)

⁸Hutchinson (2004) ⁹Medlock and Vaux (2011)

¹⁰Service and others (1986)

¹¹Service and Smith (1972)

¹²Rempel and others (1946)

¹³MacKenzie-Impoinvil and others (2015)

¹⁴Vermeil and others (1960)

¹⁵Balenghien and others (2008)

¹⁶Blagrove and others, (2016)

¹⁷Andreadis and others (1998)

¹⁸Armstrong and Andreadis (2010)

¹⁹Centers for Disease Control and Prevention (CDC) (2006)

²⁰Vaidyanathan and others (1997)

²¹Davis (1940)

²²Chamberlain and others (1954)

²³Turell and others (2006)

²⁴Aviles and others (1990)

²⁵Hammon and Reeves (1943)

²⁶Merrill and others (1934)

²⁷Turell (2012)

²⁸Kramer and others (1998)

²⁹Vaux and others (2015)

³⁰Zacks and Paessler (2010)

³¹Börstler and others (2016)

³²Schönenberger and others (2016)

³³M.S.C. Blagrove, personal communication

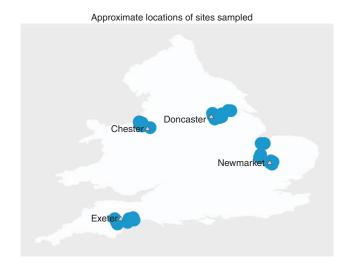


FIG 1: Map of study areas

placing a perspex cover on the open front of the box and aspirating resting mosquitoes.

Immature mosquitoes

Larval sampling was undertaken on the equine premises themselves and, where there was access, on neighbouring land within 500 m of the mosquito magnet or of grazing horses. The aim was to sample all water sources within the boundary of the premises, including all collections of artificial containers. This was not always possible due to access constraints or on larger premises. Larvae and pupae were sampled using a dipper. This is a 500 ml container with a long handle that is used to sample water sources. Each dip was then emptied into a white tray and searched for larvae. For larger water bodies, 5×500 ml dips were used in different parts of the water body, whereas for small containers only one dip sample or partial dip samples could be obtained.

Sample handling and identification

Mosquitoes were removed from the traps with a mechanical aspirator and 'Fly-nap' (Carolina Biological Supply Company) was used to produce knockdown. Adult mosquitoes were stored dry and identified within four days. Blood-fed mosquitoes were stored in 90 per cent ethanol immediately.

Larvae were pipetted into universal containers for storage. Fourth instar larvae were killed by gradually adding 90 per cent ethanol. Pupae were allowed to emerge for ease of identification.



FIG 2: Red-box trap

Live second and third instar larvae were allowed to continue to develop until the end of the fieldwork week for ease of identification. Containers were inspected daily and any dead larvae or pupae were preserved using 90 per cent ethanol for identification (Snow 1991).

Mosquitoes of all stages were identified morphologically as far as possible to species or species complex using keys of British and European mosquitoes (Marshall 1938, Cranston and others 1987, Snow 1991, Schaffner and others 2001, Becker and others 2010). *Cx. pipiens* was differentiated from *Culex torrentium* by molecular methods as described by Hesson and others (2010).

Due to the skewed distribution of the catch data, geometric mean is used for comparison in the text and Fig 3. Fig 4, showing total catch counts, is displayed with a log(10) scale.

Results Host-seeking adults

It was not possible to find drained farmland in the southwest area sampled, so two more exposed hillside sites were chosen as a comparison (sites 18 and 19, at altitudes of 120 and 114 m, respectively). At one of these hillside locations (location 19, Table 3), trapping was not carried out in September 2015 due to loss of the propane canister. A number of specimens could not be identified positively to species level due to trap damage and are recorded as unidentified *Aedes* species.

A total of 917 adult mosquitoes of 14 species were caught over a total of 285 trapping days over the 32 locations (Table 3). The geometric mean catch for each mosquito magnet trapping period (approximately 72 hours) was 3.7 (sd 3.4), across all locations and seasons. Totals caught were 487, 217, 160 and 53 in the areas sampled in the northwest, southeast, northeast and southwest, respectively.

For locations given one habitat classification, the geometric mean catch (nine days across three sampling periods) from a mosquito magnet was 6.9 (sd 5.90), 3.8 (2.5), 6.1 (3.3) and 36.5 (5.2) for premises associated with woodland, urban, drained farmland and saltmarsh habitats, respectively (Fig 3).

The most abundantly trapped species was *Oc. detritus* with a total of 499 adults caught. All three sites with total catches >100 were associated with the saltmarsh habitat of this species.

The second most abundantly trapped species was *Cs. annulata*, with 154 adults caught. *Cs. annulata* had the highest presence and was trapped on 75 per cent (24/32) of sites.

Total catch was highest in September (Fig 4), and the difference in catch was significantly higher (P<0.005) than that in May and that in June/July in a general linear model with a

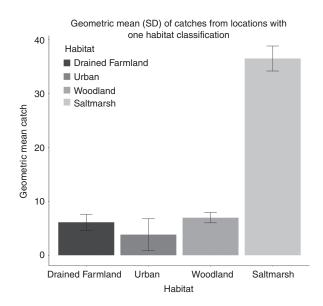


FIG 3: Geometric mean of total catch per location for each habitat type (locations only included if given one habitat)

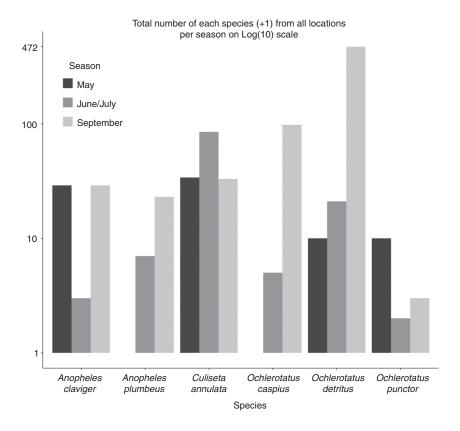


FIG 4: Total adult catches by season for each of six most abundant species

Location number and region Habitat		Anopheles ats claviger	Most abundantly caught mosquito species						
	Habitats		Anopheles plumbeus	Culiseta annulata	Ochlerotatus caspius	Ochlerotatus detritus	Ochlerotatus punctor	0ther	Total
NW 1	D	0	0	12	0	0	0		12
NW 2	U	6	0	7	24	1	0	<i>UA-</i> -5	43
NW 3	D	5	0	0	0	0	0		5
NW 4	U, S	0	1	12	3	53	0	<i>UA-</i> -5	74
NW 5	W	0	1	5	0	2	0	OCA-3	1
NW 6	W	0	0	8	3	17	0		28
NW 7	S	0	0	14	1	176	0	UA—4	19
NW 8	W, S	3	11	12	4	85	0	UA—4	119
NW 9	U	0	0	1	0	0	0		
NW 10	W, D	16	0	15	2	0	10	<i>OCA</i> —3, <i>CR</i> —2, <i>UA</i> -9 <i>AV</i> —3	60
NE 11	W	0	0	6	0	0	0		
NE 12	W, U	1	0	20	0	0	0	CuS—1, CR—1	2
NE 13	S	8	0	2	19	0	0		2
NE 14	S	5	0	0	3	1	0	CR-3	1
NE 15	D	6	0	15	0	0	0	AnM—1, CR—1 UA—1	2
NE 16	U	3	0	2	0	0	0		
SW 17	W	1	0	2	0	0	0	AnM—1	
SW 18	Н	0	0	0	0	0	0		(
SW 19	Н	0	0	0	0	0	0	CxP—1	
SW 20	W, S	3	0	2	0	4	0	CxP—1	1
SW 21	s, u	0	1	0	8	0	0	CxP—1, UA—8	18
SW 22	w, u	0	0	0	0	1	2	ŕ	
SW 23	W	0	13	1	0	0	0		1.
SW 24	W	0	1	1	0	0	0	CxP—1	
SE 25	U	0	0	0	0	0	0	CxP—1	
SE 26	W	0	0	1	0	0	0	<i>0D</i> —1	
SE 27	W, U	0	0	0	0	0	0	CR—1	
SE 28	W	0	0	6	0	0	0	OR—3	
SE 29	D	0	0	3	1	0	0	OR—2	
SE 30	S	0	0	4	33	155	0	_	19
SE 31	S	1	0	2	0	2	0		
SE 32	D	0	0	1	0	0	0		

AnM, An. maculipennis; AV, Ae. vexans; CR, Cq. richiardii; CuS, Cs. subochrea; CxP, Cx. pipiens s.l.; D, drained farmland; NE, northeast; NW, northwest; OCA, Oc. cantans; OD, Oc. dorsalis; OR, Oc. rusticus; S, salt marsh; SE, southeast; SW, southwest; U, urban; UA, unidentified Aedes species; W, woodland

negative binomial distribution using the R statistical programme and the MASS package (The R Foundation (2016) R: The R Project for Statistical Computing. The R Foundation. https://www.r-project.org/. Accessed June 15, 2016; Ripley and others 2016). Thirty-one sites were sampled (one site not sampled), and total mosquito number from all locations was 679 with a geometric mean of 5.6 (sd 5.1) per location.

No mosquitoes were trapped while feeding on horses. Only three blood-fed mosquitoes were trapped, all were part-fed individuals caught in the mosquito magnet, of which two were *Oc. detritus* and one was *Cs. annulata*. One mosquito (*Oc. caspius*) was sampled landing on a human host. Pilot host-landing catches using horses carried out in September 2014 yielded 20 *Oc. detritus*, 3 *Oc. caspius* and 2 *Cs. annulata*, in two 15-minute daytime sampling efforts at site 8.

Resting adults

Sampling of resting mosquitoes was unsuccessful. No mosquitoes were found in the red-box traps.

Immature mosquitoes

Immature mosquitoes were recovered by dipping of water sources on 23 of 32 premises. A total of 61 samples containing mosquito larvae or pupae were collected from a variety of water sources including ditches, buckets and water butts, tyres, ruts, muck heaps, pools and ponds.

Cx. pipiens s.l., Cx. torrentium, Cs. annulata/alaskaensis/subochrea, Culiseta fumipennis, Cs. morsitans, Oc. caspius, Anopheles claviger and An. maculipennis s.l. were captured using dipping techniques.

The majority of samples were from artificial containers with small amounts of water, such as tyres. Therefore on most occasions, samples from each container were less than 500 ml, so it was not considered appropriate to state the numbers sampled, nor possible to compare larval numbers across sites. Larval samples were used to identify the presence of a species rather than its relative abundance.

A selection of larvae identified morphologically as *Cx. pipiens/torrentium* was further identified by molecular methods for each location. Of the 23 sites from which samples were obtained, *Cx. pipiens* larvae were identified from 65.2 per cent of locations, *Cx. torrentium* from 47.8 per cent. Both species were found on 21.7 per cent of these 23 locations. Both *Cx. pipiens* and *Cx. torrentium* larvae were obtained from at least two sites in all four regions.

Cs. annulata/alaskaensis/subochrea larvae cannot be differentiated morphologically and were obtained at 28.1 per cent (9/32) of sites. Due to the rarity of Cs. alaskaensis and the relative abundance of Cs. annulata, it is likely that the vast majority of these were Cs. annulata. The total of sites with presence of Cs. annulata/alaskaensis/subochrea including adult samples was 84.4 per cent (27/32).

Discussion

This study is, to the authors' knowledge, the first survey of mosquito species on equine premises in the UK. This work has demonstrated the presence of several mosquito species that are candidate vectors of pathogens affecting horses. Commonly found mosquito species on equine premises during this study included Oc. detritus, Oc. caspius, Cs. annulata, Cx. pipiens s.l., Cx. torrentium, An. claviger, An. plumbeus and Oc. punctor. Although mosquito density could be considered low at most of the sites sampled, this could be partly explained by the fact that the months of March, April and May in 2015 were relatively dry for all of the regions except the northwest (Met Office 2016). Climate change predictions suggest increased temperature and potential for flooding events in the UK (Met Office 2010, Caminade and others 2012, Medlock and Leach 2015), which are likely to increase the abundance of native mosquito species. It therefore seems likely that in the future there may be significantly increased horse-vector interaction, particularly with mosquito species that thrive in warmer regions of Europe, such as

Cx. annulata, Oc. caspius, Cx. pipiens s.l. Oc. detritus, An. plumbeus, Cq. richiardii, An. maculipennis and Ae. vexans (Balenghien and others 2008). The species trapped in the current study are all considered mammalophilic or bite both birds and mammals, with the exception of *Cx. torrentium* that is strongly ornithophilic (bird biting). Three European studies provide evidence that Cx. pipiens s.l. found in rural areas will bite mammals, including horses (Balenghien and others 2008, Börstler and others 2016, Schönenberger and others 2016). Although not all of these studies differentiated Cx. pipiens form pipiens from Cx. pipiens form molestus, the study of Börstler and others (2016) records a significant number of Cx. pipiens form pipiens with mammalian blood meals. Oc. detritus, Oc. caspius, Cx. pipiens s.l., Oc. punctor and An. plumbeus have all been shown to transmit arboviruses affecting horses and therefore are important when considering the risk of mosquito-borne equine disease. Also, 11 of the 16 species found on equine premises during this study are laboratory-competent vectors of, or are implicated in, naturally occurring disease cycles for at least one arbovirus-affecting horses (Table 2). An important aspect of this study is that the authors trapped very few blood-fed mosquitoes: just three in the mosquito magnet and none by other methods. This begs the question of whether the mosquitoes present at equine premises in the UK only rarely feed on equines or whether they feed but were not caught. A number of factors suggest that the latter is the most likely explanation: (i) the mosquito magnet is designed to trap host-seeking rather than blood-fed adults; (ii) many of the premises had other potential hosts present (human beings, cattle, small mammals), indicating that the low number of trapped blood-fed mosquitoes cannot be attributed to the specific avoidance of equids; (iii) in pilot work in September 2014, mosquitoes Cs. annulata, Oc. caspius and Oc. detritus were directly observed by the author feeding on horses; and (iv) most of the species caught in this study have been reported, in other studies, to feed on horses and/or transmit arboviruses to horses. Nevertheless, and probably due to the inherent difficulties in trapping blood-fed mosquitoes in the UK (Brugman and others 2015), blood-feeding on horses has not been confirmed in this study. A large sampling effort and high mosquito densities are required to maximise trapping of blood-fed mosquitoes. The number of sites included in this study dictated that sampling effort on each site was necessarily lower than that of other recent studies (Brugman 2016), and seasonal variation in abundance due to climatic conditions, for example, a dry early summer period (Met Office 2016), may have supressed mosquito density. However, all of the species sampled in this study, with the exception of Cx. torrentium and Cs. morsitans, have been shown to bite equines (Table 2), and four of the six most abundant species in adult catches have been shown to bite horses in the UK either in previous studies or in pilot work for this study. Further work would be required to investigate the feeding rate of UK populations of these mosquitoes on horses, and host bait catches (Schönenberger and others 2016) would seem most likely to provide useful information.

The comparatively high numbers of Oc. detritus and Oc. caspius caught on some saltmarsh-associated sites are consistent with previous studies and reports of significant nuisance biting (Clarkson and Setzkorn 2011, Medlock and others 2012, Medlock and Vaux 2013) and confirm that there is significant potential for host-vector interaction between these species and horses. These two species are competent vectors of WNV (Vermeil and others 1960, Blagrove and others, 2016). Detailed, high resolution information regarding horse and mosquito species distribution is lacking (Lo Iacono and others 2013). However, using previously published horse distribution data at postcode scale (Boden and others 2012, Lo Iacono and others 2013) and saltmarsh distribution (Adnitt and others 2007), in combination with mosquito species records, several coastal areas of England appear worthy of further investigation for hostvector interaction potential. These areas have high horse density, saltmarsh presence and records of Oc. detritus and Oc. caspius (The Walter Reed Biosystematics Unit 2014, National Biodiversity Network 2016a, b) and include the Severn estuary, South Devon coast, the south coast of England from Swanage to Chichester and the Dee and Mersey estuaries. Two of these areas were sampled during this study: Wirral (Dee estuary) and the South Devon coast.

The finding that the WNV vector *Cx. pipiens* was common on equine premises with suitable water sources is expected as this species has a widespread distribution (Medlock and others 2005, Medlock and Vaux 2011), but this study confirms that suitable container habitats are commonplace on equine premises. *Cx. torrentium* is a major enzootic (wildlife) vector of Sindbis virus in Scandinavia (Hesson and others 2015) and may therefore be capable of a similar role in transmission of other arboviruses. *Cx. pipiens* and *Cx. torrentium* were found on a number of occasions in all four regions, suggesting that *Cx. torrentium* may be more prevalent in the north of England than previously recognised (Medlock and others 2005).

One of the most interesting results to emerge from the current study was the presence of *Cs. annulata* on the majority of sites (27/32). It was also the second must abundant species in mosquito magnet samples, after *Oc. detritus*. While *Cs. annulata* is known to have a widespread distribution in the UK (Medlock and others 2005), this study provides evidence of the potential for host–vector interaction with UK equines. *Cs. annulata* has recently been demonstrated to be vector competent for WNV (M. S. C. Blagrove, personal communication), and as the species bites both birds and mammals including horses (Schönenberger and others 2016), it therefore has potential to transmit arboviruses from avian reservoirs and hence serve as a 'bridge vector'. Combined with its ability to breed in a variety of water sources and presence on most sites sampled, this makes it an important species for further study.

The authors' results suggest that mosquito species presence is determined mainly by local mosquito breeding habitat rather than equine host availability or management factors. However, biting of horses may be affected by practices such as use of repellents, rugs and masks, building design, and duration and timing of grazing.

Mosquito magnets are a commonly used trap in Europe for surveillance. They catch almost all mammalophagic species of mosquito, and catch more species than other systems and in greater numbers. BG sentinel and Centers for Disease Control and Prevention traps were not used in this study as they were considered less suitable due to the risk of unpredictable precipitation damaging samples, and because for wide-scale trapping in the UK, it may prove more practicable to use propane vendor's delivery services than to transport large amounts of dry ice or carbon dioxide. Red-box traps were used in the current study to attempt to trap blood-fed mosquitoes; however, no mosquitoes were captured. Similar but larger red-box traps have been successful in capturing An. maculipennis s.l., Cs. annulata and Culex species in England (Brugman 2016). At similar future levels of mosquito abundance and without the presence of invasive species, surveillance on equine premises in the UK should be based around the use of mosquito magnets and larval sampling.

Mosquito populations often have a patchy distribution (Medlock and others 2005, Snow and Medlock 2008, Golding 2013) and many are considered uncommon or rare. Simple random sampling of equine premises may have resulted in very low catches. It is also almost impossible to prove species absence, so using random sampling risked obtaining poor quality data. Stratified sampling is an alternative method, commonly used by ecologists studying rare species (Thompson 2012). Using the data obtained under this sampling regimen, it is not possible to estimate the risk of equine—mosquito interaction across the UK, but more accurate assessment of risk at individual sites based on local habitat is achievable.

There are a number of introduction pathways that could conceivably be involved in importation of arboviral disease to the UK. Perhaps the most often discussed is introduction of WNV

by migratory birds, but trade and transport of exotic birds and pets, and inadvertent vector transportation are also relevant risks. There is some recent evidence that human populations may continue epidemic transmission of VEEV in urban environments (Bowen and Calisher 1976, Watts and others 1998, Morrison and others 2008). Therefore in the event of an outbreak in the Americas, human movements as well as horse movements may constitute a risk (Adams and others 2012). Livestock transport, human transport and possibly mosquito eggs may present risk of RRV introduction (Harley and others 2001). Due to the complexity of the transmission cycles, virus introduction may not result in autochthonous (in-country) transmission.

In conclusion, the current study has highlighted a number of mosquito species that should be investigated with regard to vector competence and effectiveness of protection measures for equines. The authors' work has shown that horses in the UK are at risk of attack from a wide variety of mosquito species, several of which are known to be vectors of equine arboviruses in affected countries.

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