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1	Revealing potential bridge vectors for BTV and SBV: a study on Culicoides
2	blood feeding preferences in natural ecosystems in Spain
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6	Culicoides host-feeding preferences
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10	Sandra Talavera ^{1*} , Francesc Muñoz-Muñoz ² , Marta Verdún ¹ , Nuria Pujol ¹ , Nitu Pagès ^{1,3}
11	¹ IRTA, Centre de Recerca en Sanitat Animal (CReSA, IRTA-UAB), Campus de la
12	Universitat Autònoma de Barcelona, 08193 Bellaterra, Spain.
13	² Departament de Biologia Animal, de Biologia Vegetal i d'Ecologia, Universitat
14	Autònoma de Barcelona, 08193 Bellaterra, Spain
15	³ Present address: CIRAD, UMR ASTRE, Petit-Bourg, Guadeloupe, France
16	
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19	*Corresponding author
20	E-mail: sandra.talavera@irta.cat (ST)
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Several species of *Culicoides* (Diptera: Ceratopogonidae) are vectors of pathogens, such as the Bluetongue (BTV) and the Schmallenberg (SBV) viruses, that cause important diseases in domestic and wild ruminants. Since wild ruminants can contribute to overwintering and epizootics of both diseases, knowledge of Culicoides host-feeding behaviour in natural ecosystems is an important parameter to better understand their epidemiology. Blood engorged Culicoides females trapped in natural areas inhabited by different wild ruminant species were genetically analysed to identify host species. The origin of blood meals was identified in 114 females of 14 species of Culicoides. A total of 104 (91.1%) Culicoides fed on mammals and 10 (8.9%) on birds. The most abundant host identified was red deer (66.7%), followed by human (13%) and fallow deer (6.1%). Eleven of the 14 species of *Culicoides* fed exclusively on mammalian hosts. Among them, 5 are mammalophilic species considered important BTV and/or SBV vectors. The results of the present study confirm that C. imicola, C. obsoletus, C. scoticus, C. pulicaris and C. punctatus fed on wild ruminants, and therefore support the hypothesis that these species can act as bridge vectors by facilitating the circulation of pathogens between wild and domestic ruminant communities.

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KEY WORDS: *Culicoides*, BTV, host preference, SBV, Spain, vector, wild ruminants

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INTRODUCTION

2	The genus <i>Culicoides</i> has been extensively studied due to the fact it contains species
3	that act as vectors of several diseases. There are approximately 1400 Culicoides
4	species in the world (Borkent 2012, 2014). The females of 96% of these species are
5	expected to be obligate blood-feeders, and in consequence, can act as vectors of
6	protozoa, filarial worms and viruses affecting birds, humans, and other mammals
7	(Mellor et al., 2000). One of the most economically important pathogens transmitted
8	by Culicoides is Bluetongue virus (BTV), which produces an infectious non-
9	contagious disease of domestic and wild ruminants (Mellor & Wittmann, 2001).
10	Culicoides transmit Schmallenberg virus (SBV) to ruminants as well, which was
11	detected for the first time in Germany and the Netherlands in summer and autumn
12	2011 (De Regge et al., 2012; Elbers et al., 2013; Goffredo et al., 2013; Larska et al.,
13	2013; Balenghien et al., 2014). Since then, the disease has been detected in almost
14	the whole of Europe. In Spain, it has been reported to affect sheep and goats
15	(Fernández-Aguilar et al., 2014). To date, six species of Culicoides have been
16	described as vectors for BTV and SBV in Europe: C. obsoletus, C. scoticus
17	(Carpenter et al., 2006; Carpenter et al., 2008a; De Liberato et al., 2005; Savini et
18	al., 2005, De Regge et al., 2012; Elbers et al., 2013; Goffredo et al., 2013), C.
19	dewulfi (Meiswinkel et al., 2007; De Regge et al., 2012; Elbers et al., 2013;
20	Goffredo et al., 2013), C. chiopterus (Dijkstra et al., 2008; De Regge et al., 2012;
21	Elbers et al., 2013), C. pulicaris (Caracappa et al., 2003; Balenghien et al., 2014),
22	and C. imicola (Balenghien et al., 2014). Culicoides punctatus (Larska et al., 2013)
23	and C. nubeculosus (Balenghien et al., 2014) are considered SBV vectors, and a
24	recent study indicates that C. newsteadi and C. punctatus might act as BTV vectors
25	(Goffredo et al., 2015).

High seroprevalence for BTV and SBV antibodies in wild ruminants suggests	
wildlife might play an important role in the transmission of such pathogens in	
Europe (Falconi et al., 2011; Chiari et al., 2014). Moreover, it has been observed	
that some Culicoides species feeding on domestic ruminants are abundant and in	
close contact with wild ruminants in natural ecosystems, which suggests that these	
species may be acting as bridge vectors (Talavera et al., 2015). To be sure of the	
role of these species as bridge vectors and to better understand the epidemiology of	
the different diseases transmitted by Culicoides, it is necessary to know the standing	
host preferences of Culicoides species inhabiting natural ecosystems. However,	
most of the studies dealing with Culicoides host feeding preferences in Europe have	
been conducted around livestock areas (e.g. Rawlings et al., 2003; Bartsch et al.,	
2009; Garros et al., 2011; Viennet et al., 2011; Lassen et al., 2011; Ninio et al.,	
2011; Calvo et al., 2012; Pettersson et al., 2013). Only recently have some studies	
been performed in natural ecosystems (Santiago-Alarcón et al., 2013, Bobeva et al.,	
2015), and to the best of our knowledge none of them in natural areas with an	
important presence of wild ruminants. While some studies have inferred Culicoides	
host preferences based on the morphology of the sensory organs present in the	
antennae and the palps (Blackwell 2004), direct evidence is needed to be sure of the	
food source of a particular species. Early studies on Culicoides feeding patterns	
were based on serological (Braverman et al., 1971; Walker & Davie, 1971; Nevill &	
Anderson, 1972; Boakye et al., 1999) and immunological assays (Blackwell et al.,	
1994; Blackwell et al., 1995). However, these methods allowed the identification of	
a limited number of potential vertebrate host species. Recently, molecular	
techniques have been developed, which have increased the accuracy of host	

identification at species level (Votýpka *et al.*, 2009; Bobeva *et al.*, 2015; Martínez

de la Puente *et al.*, 2015; Slama *et al.*, 2015).

Given the sparse knowledge about *Culicoides* hosts in natural ecosystems, the main

goal of the present study is to identify by means of molecular techniques the

vertebrate species on which *Culicoides* feed in these ecosystems. Knowing which

vertebrate species on which *Cuticoides* reed in these ecosystems. Knowing which vertebrate species the different species of *Culicoides* found in natural ecosystems feed on, will shed light on which species of *Culicoides* can act as bridge and enzootic vectors for *Culicoides* borne pathogens, and therefore on the mechanisms behind the spillover of BT and SB from domestic to wild ruminants (and viceversa). In addition, this information could indirectly indicate which reservoirs are

MATERIALS AND METHODS

significant in associated vector-borne diseases.

Sampling

Culicoides were trapped between years 2009-2010 in seven private Spanish areas of land characterized by their distinctive bioclimatic features and wild ruminant communities (S1 Table). Bioclimatic variables and altitude were obtained from a climatic atlas of the Iberian Peninsula (Ninyerola et al., 2005). Landscape variables were obtained from the Global Environment Monitoring database (Global Land Cover 2000). The distribution of ruminant species in Spain was obtained from the atlas of land mammals in Spain (Palomo et al., 2007). Three CDC black light traps (John W. Hock Company, Gainesville, FL, USA) were placed close to water points used as troughs for wild ruminants. The traps were located at a distance of more than 1 km from domestic ruminants and the distance between traps was about 15 m. Traps

were switched on at dusk and off at dawn the following day for three consecutive nights once a month during summer and autumn for two consecutive years. In Spain, seven species of wild ruminants are present, red deer (*Cervus elaphus*) being the most abundant, fallow deer (*Dama dama*), roe deer (*Capreolus capreolus*), mouflon (*Ovisaries musimon*), Spanish ibex (*Capra pyrenaica hispanica*), Pyrenean chamois (*Rupicapra pyrenaica*) and aoudad (*Ammotragus lervia*).

Morphological and molecular identification

Trapped insects were killed by drowning in the capture containers with soapy water, and then stored in 70% ethanol. *Culicoides* midges were morphologically identified under a stereomicroscope (Nikon SMZ) to species-group level according to wing pattern. An accurate morphological identification was performed of at least one female of each species. Specimens were dissected using sterilized ultrafine tweezers, mounted on slides with Canada Balsam solution, and identified to species level according to their morphology (Campbell & Pelham-Clinton, 1960; Kremer 1965; Kremer *et al.*, 1973; Delécolle 1985; Glukhova 1989) using a Nikon Eclipse E200 light microscope. Once females of the Obsoletus complex had been grouped according to wing pattern, species identification was performed genetically. Briefly, DNA was isolated from wild caught individuals and used as a template in a PCR reaction targeting the gene marker cytochrome oxidase I (COI), as described in Pagès & Sarto i Monteys, 2005 and Nolan *et al.*, 2007.

Blood identification

Blood engorged females were used for DNA extraction. The abdomen of each blood engorged female was dissected with sterilized ultrafine tweezers. The abdomen served as a template for DNA extraction using DNeasy Blood and Tissue kit (Qiagen), in accordance with the manufacturer's recommendations. DNA was used for amplification of a fragment of COI gene using a nested PCR with degenerated primers, as described in Alcaide *et al.* 2009. DNA purified PCR products were sequenced on both strands using Big Dye Terminator version 3.1 cycle sequencing kit (Applied Biosystems) and analysed on an ABI PRISM 3730 Automated sequencer (Applied Biosystems).

DNA sequences were edited using Bioedit sequence alignment editor software (version 5.0.9. for Windows; Hall, 1999). Fasta files were used for nucleotide BLAST (Basic Local Alignment Search Tool) for host species identification.

RESULTS

Collection of biting midges

A total of 102,693 *Culicoides* specimens were collected during the study period. Less than 1% of collected *Culicoides* were blood engorged females (n=264). Engorged females were collected at all sampling sites except in Boumort where collections were scarce (0.1% total captures; S1 Table). Among the 264 blood engorged females, genetic characterization of blood-meal origin was successful in 114 (44%) of them, probably due to an enzymatic degradation of the blood within the digestive tract of the midge. Identification of hosts selected by *Culicoides* to feed on their blood was achieved for 14 *Culicoides* species. Among them, 47 were

1	obtained from C. obsoletus, 13 from C. scoticus, 10 from C. punctatus and 9 from
2	C. imicola (Table 1).
3	
4	Blood host identification
5	Culicoides fed on 12 different host species, 6 mammals and 6 birds. Eleven of the
6	14 Culicoides species exclusively had bloodmeals of mammalian origin, two had
7	blood only from avian hosts, and one species was found to be fed both on mammals
8	and birds (Table 1).
9	At the individual level, among the 114 females in which blood origin was
10	successfully identified, 95 (83.5%) fed on species of wild vertebrates, 15 (13%) on
11	humans and 4 (3.5%) on cattle. All <i>Culicoides</i> with cattle blood were collected in La
12	Almoraima, the closest sampling site to livestock facilities (one kilometre; S1
13	Table). At a taxonomic level, 104 (91%) of the 114 bloodmeals identified were of
14	mammals and 10 (9%) of birds (Table 1). In bloodmeals, the most frequently
15	identified host was red deer (Cervus elaphus, 66.7%), human (Homo sapiens, 13%)
16	and fallow deer (Dama dama, 6.1%). The most frequent avian hosts were blackbird
17	(Turdus merula, 3.5%) and hoopoe (Upupa epops, 1.8%) (Table 1).
18	In all areas the most abundant ruminant host was red deer (Cervus elaphus) except
19	for El Juanar where it was Spanish ibex (Capra pyrenaica hispanica) (S1 Table).
20	The most abundant wild ruminant of each zone according to local censuses was
21	detected as the preferred host for Culicoides, except in La Morera where only
22	human and hoopoe blood was detected (Table 1).
23	

DISCUSSION

Recent studies highlight that most *Culicoides* species are able to feed on several 1 2 vertebrate species, but usually present preferences for mammals or for birds (Martínez de la Punte et al., 2015), demonstrating that many species of Culicoides 3 are not generalist (Santiago-Alarcon et al., 2012). The results obtained here support 4 those from the aforementioned previous studies, since *Culicoides* specimens 5 6 collected at natural ecosystems were found to feed mainly on one or other of the two 7 groups, mammals or birds. Nevertheless, mammalophilic and ornithophilic 8 Culicoides species might feed, respectively, on different mammal or bird species opportunistically. Host selection generally seems to reflect host availability 9 10 (Pettersson *et al.*, 2013). Culicoides species previously described as strict mammal bloodfeeders based on 11 morphological traits (Talavera et al., 2015) contained exclusively blood of 12 mammalian origin. Similarly, species previously described as ornithophilic feeders 13 (Talavera et al., 2015) had avian blood except C. univitatus, which was found to 14 15 have only fed on humans. However, all these specimens were trapped in the same 16 locality and during the same period. Culicoides festivipennis mainly fed on birds; however one of the seven bloodmeals identified in this species was of human origin. 17 18 Thus, although according to our data C. festivipennis seems to prefer birds, they also partially support studies describing it as a generalist feeder (Santiago-Alarcon et al., 19 2012). Culicoides species with generalist preferences are of special interest because 20 they could promote the emergence of new diseases (Santiago-Alarcon et al., 2012). 21 22 The genetic identification of bloodmeals allowed us to report four new Culicoides hosts for the first time in Europe: two wild ruminants, fallow deer (Dama dama) and 23 24 Spanish ibex (Capra pyrenaica), and two birds, Sylvia cantillans and Upupa epops 25 (for purposes of comparison see Martínez de la Puente et al., 2015).

Knowledge about vector host preferences is critical to better understand the epidemiology of vector borne diseases (Mukabana et al., 2002; Lyimo & Ferguson, 2009). Thus, the opportunistic or selective host tendencies of a vector may affect the spread of a disease (Burkot 1988). The *Culicoides* host preferences are dependent on the intrinsic features of each *Culicoides* species, as well as on environmental factors such as host availability (Lyimo & Ferguson, 2009; Burkot 1988). However, most studies of *Culicoides* host preferences have usually been performed in farms or rural areas with livestock because of the economic importance of Culicoides transmitted pathogens for livestock production (Rawlings et al., 2003; Bartsch et al., 2009; Garros et al., 2011; Viennet et al., 2011; Lassen et al., 2011; Ninio et al., 2011; Pettersson et al., 2013). The main BTV and SBV vectors (C. imicola, C. obsoletus, C. scoticus, C. pulicaris and C. punctatus) collected in this study were found to feed exclusively on mammals, mainly wild ruminants (70/82; Table 1). The present study reports for the first time the origin of C. imicola bloodmeals molecularly identified in Europe. In Africa C. imicola was described feeding mainly on horses, cattle and sheep (Logan et al., 2010), and also on humans, birds and dogs (Slama et al., 2015). The results of the present study supports the hypothesis that C. imicola, C. obsoletus, C. scoticus, C. pulicaris and C. punctatus have the potential to act as bridge vectors and circulate pathogens at the interface between wild and domestic ruminant communities. Interestingly, two thirds of the bloodmeals identified in the study were of red deer, whose population density is similar to that of domestic livestock in some areas of Europe. Red deer is susceptible to BTV and might carry the virus asymptomatically for long periods, thus acting as a potential reservoir candidate (Rodríguez-Sánchez et al., 2010). In fact, previous studies found that

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- there were more BT cases in domestic ungulates in areas where red deer was present
- 2 (Falconi *et al.*, 2011).
- There are some species described as ornithophilic or with unclear feeding habits as
- 4 C. parroti, C. saevus and C. yemenensis (Talavera et al., 2015), with low catches,
- but feeding exclusively on wild ruminants in natural ecosystems. These species have
- been poorly studied to date because of sparse collection in domestic ruminant farms.
- 7 Different authors have pointed out that UV light-traps do not accurately reflect
- 8 proportions of *Culicoides* biting in the field (Logan *et al.*, 2010; Venter *et al.*, 2012),
- 9 suggesting that these traps could underestimate the assessment of BTV risk
- 10 (Carpenter et al., 2008b; Gerry et al., 2009). C. parroti belongs to the subgenus
- 11 *Monoculicoides*, which includes the main BTV vectors in the US (*C. sonorensis* C.
- variipennis and C. occidentalis), and has a wide distribution through Europe to
- Russia (Augot *et al.*, 2013; Grogan & Lysyk, 2015). Thus, its potential role in the
- epidemiology of BTV and SBV should be further investigated.
- A total of 13% of engorged *Culicoides*, belonging to seven different species, were
- found to bite on humans, which concurs with previous studies which have reported
- that common mammalophilic *Culicoides* species in Europe, including arbovirus
- vectors, occasionally feed on humans (Slama et al., 2015; Garros et al., 2011;
- 19 Santiago-Alarcon et al., 2012; Carpenter et al., 2013). Attacks on humans mainly
- 20 occur when preferred hosts such as livestock or wildlife are either scarce or absent.
- 21 Worldwide, the role of *Culicoides* in the transmission of zoonotic arboviruses is
- currently unclear, but thought to be limited (Carpenter *et al.*, 2013).
- Field seroprevalence and experimental infections suggest that wild ruminants are
- susceptible to BTV and SBV infection (Falconi et al., 2011; Chiari et al., 2014;
- Ruiz-Fons et al., 2008; Ruiz-Fons et al., 2014; García et al., 2009; García et al.,

1	2011; Rodríguez-Sánchez et al., 2010; Linden et al., 2012; Laloy et al., 2014;
2	Fernández Aguilar et al., 2014). Molecular analyses of bloodmeals of Culicoides
3	species collected in natural ecosystems confirm that the major putative BTV and
4	SBV vectors in Europe (C. imicola, C. obsoletus, C. scoticus, C. pulicaris and C.
5	punctatus) are mostly mammalophilic and bite more frequently on wild ruminants.
6	Therefore, these species might act as bridge vectors leading the spillover of both
7	diseases between domestic and wild ruminants in epizootic areas.
8	
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15	conflict of interest regarding the publication of this article.
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Table 1. Blood meal sources (vertebrates hosts) of engorged *Culicoides.* **Pro.,** Proaza; **P.P.,** Puig la Penya; **Q.M.,** Quintos de Mora; **L.M.,** La Morera; **E.J.,** El Juanar; **L.A.,** La Almoraima

	1		Mamm	nals					Bi	rds			
Species	Capra pyrenaica	Cervus elaphus	Dama dama	Homo sapiens	Sus scrofa	Bos taurus	Cyanistes caeruleus	Delichon urbica	Erithacus rubecula	Sylvia cantillans	Turdus merula	Upupa epops	% Blood engorged
C. cataneii	pyronaida	olapitao	dama	Gaprone	Corola	taarao	L.A. 1	unbiou	rabooala	caritilario	moraid	L.M.1	1.8
C. circumscriptus											Q.M.1	L.A.1	1,8
C. festivipennis				L.M.1				Pro. 1	Q.M.1	Q.M.1	Q.M. 2 L.A. 1		6,1
C. imicola		L.A. 6				L.A. 3							7,9
C. newsteadi		L.A. 1	L.A. 2	L.A. 1		L.A. 1							4,4
C. obsoletus	E.J. 1	Pro. 26 P.P. 10 Q.M. 2 L.A. 4		Pro. 1 Q.M. 2	Pro. 1								41,1
C. parroti		Q.M. 3 L.A. 1	L.A.1										4,4
C. pulicaris		P.P. 2	P.P. 1	Pro.1									3,5
C. punctatus		P.P. 2 Q.M. 6	P.P. 2										8,8
C. saevus		Q.M. 2											1,8
C. scoticus		Pro. 4 P.P. 1 L.A. 4	P.P.1	L.A. 3									11,3
C. subfagineus		L.A. 1									The state of the s		0,9
C. univitattus				L.A. 5									4,4
C. yemenensis		L.A. 1		L.A. 1									1,8
% Blood engorged	0,9	66,7	6,1	13	0,9	3,5	0,9	0,9	0,9	0,9	3,5	1,8	

S1 Table. Data summary of ecological variables and characterization of the sampling sites (Palomo *et al.*, 2007; Global Land Cober 2000). **A,** altitude; **AP,** annual precipitation; **LT,** mean low temperature of the coldest month; **HT,** mean high temperature of warmest month (Ninyerola *et al.*, 2005)

Sampling site	Geographical variables	Bioclimatic	Bic	Bioclimatic variables	variabl	es	Environment near	Environment near the sampling site	Ruminants in sampling place	Blood engorged Culicoides
	Coordinates	Classification	A(m)	HT(°C)	.T(°C)	AP(mm)	LT(°C) AP(mm) Domestic ruminants (distance in km)	Water (distance in meters)	(more abundant - less abundant)	% total
Proaza	43°12' N 06°03' W	Temperate oceanic submediterranean 349	349	25		1000	1000 Farm (4,5)	wet soil, not water on surface	red deer, roe deer	29,8
R.N.C.Boumort	N.C.Boumort 42°12' N 01°06' E	Temperate hyperoceanic	1276	. 25	7,5	006	Free domestic livestock (1)	Pond (<5)	red deer, fallow deer, roe deer, chamois	0
Puig la Penya	uig la Penya 42°31' N 02°32' E	Mediterranean pluviseasonal oceanic 228	228	27,5	2,5	1000	Farm (2)	Pond (25)	red deer, fallow deer, mouflon	16,7
Quintos de Mora	uintos de Mora 39°23' N 04º06' W	Mediterranean pluviseasonal oceanic 718	718	35		482	Farm (10)	Pond (<5)	red deer, fallow deer, roe deer	17,6
La Morera	38°54' N 04°16' W	Mediterranean pluviseasonal oceanic 707	707	35	2,5	200	Farm (4)	Pond (5)	red deer, mouflon, aoudad	1,8
El Juanar	36°34' N 04°53,'W	Mediterranean pluviseasonal oceanic 870	870	27,5	2.5-5	850	Farm (5)	Cement trough (<5)	spanish ibex	1 88'0
La Almoraima	-a Almoraima 36°27' N 05°43' W	Mediterranean pluviseasonal oceanic 45	45	30,5	2,5	922	Farm (4)	wet soil, not water on surface	red deer, fallow deer, roe deer, mouflon 33,3	33,3