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First record of the West Nile virus bridge vector *Culex modestus* Ficalbi (Diptera: Culicidae) in Belgium, validated by DNA barcoding

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Abstract

A thorough knowledge of the presence and spatio-temporal distribution patterns of vector species are pivotal to assess the risk of mosquito-borne diseases in Europe. In 2018, a *Culex* larva was collected during routine monitoring activities to intercept exotic *Aedes* mosquito species in the port of Antwerp (Kallo, Belgium). The larva, collected from a pond in mid-September, was morphologically identified as *Culex modestus*, and this identification was subsequently confirmed by *COI* barcoding. It is the first confirmed record of this West Nile virus bridge vector in Belgium. The present study also demonstrates the value of DNA-based identification techniques to validate the presence of potential vector species.

Key words: biodiversity, DNA-based identification, larva, monitoring, mosquito, new record

Introduction

Mosquitoes are notorious for their ability to transmit several pathogens of human diseases, and high costs are usually associated with the implementation of prevention and control strategies, health care, loss of economic productivity or recreational activities (Roiz *et al.* 2018; Thompson *et al.* 2020). Main factors contributing to the (re-)emergence of diseases caused by mosquito-borne pathogens are increased globalisation (causing introduction of exotic vectors and/or pathogens), altered landscape management (e.g. wetland restoration, urbanisation), changing socioeconomic behaviour and climatic changes (Becker 2008; Randolph & Rogers 2010; Roiz *et al.* 2015; Rochlin *et al.* 2016). In this context, thorough knowledge of mosquito diversity is crucial to assess the current and future risks of the local transmission of pathogens (Versteirt *et al.* 2013; Medlock *et al.* 2015; Calzolari 2016). For example, local chikungunya and dengue fever outbreaks in southern Europe resulted from local transmission of travel-related arboviruses by the established exotic vector species *Aedes albopictus* (Skuse) (ECDC 2018). Also, several indigenous European mosquito species can transmit mosquito-borne pathogens, such as West Nile, Tahyna, Sindbis, Batai and Inkoo viruses (Lundström 1999; Hubálek 2008).

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In Europe, West Nile virus (WNV) has been showing a strong resurgence since 1996 (Hubálek & Halouzka 1999; Zeller & Schuffenecker 2004). From 2010 onwards, WNV has engendered a permanent public health problem in Europe (Papa *et al.* 2010; Zeller *et al.* 2010; Ziegler *et al.* 2019). Its suspected or proven main vectors in this region are *Culex pipiens* Linnaeus, *Cx. modestus* Ficalbi and *Cx. perexiguus* Theobald (Balenghien *et al.* 2006; Balenghien *et al.* 2007; Becker *et al.* 2010; Brugman *et al.* 2018). These three species regularly feed on birds, humans and other mammals (Becker *et al.* 2010; Radrova *et al.* 2013; Brugman *et al.* 2017). As such, *Cx. modestus* is the principal WNV bridge vector in the Camargue wetlands of France (Ponçon *et al.* 2007). While *Cx. pipiens* is a widespread and abundant species in Belgium, *Cx. modestus* has not yet been recorded in the country (Versteirt *et al.* 2013; Boukraa *et al.* 2015). This study presents the first morphological and molecular evidence of its occurrence in Belgium.

Material and methods

Larval sampling

Within the framework of the 'Monitoring of Exotic MOsquito species' (MEMO) project, mosquito larvae were collected from April until November 2017, 2018 and 2019 at 23 localities in Belgium (Deblauwe *et al.* 2020). Larval sampling targeted predominantly artificial container habitats (e.g. flower vases, rain barrels, tyres), as these are the preferred larval habitats of exotic *Aedes* species, but also some natural sites were sporadically sampled. Fine-meshed aquarium nets were used to collect larvae from the water. The larvae were killed by a heat shock at 70°C in the laboratory, and subsequently transferred in 80% ethanol for storage at room temperature before species identification.

Morphological and DNA-based species identification

Larvae were morphologically identified using the keys of Becker *et al.* (2010) and Gunay *et al.* (2018). For every species, one or multiple larvae were selected as voucher specimens. After morphological identification, the anterior part of each larval abdomen was used for DNA-based species identification, the head and posterior part of the abdomen of these voucher larvae were slide-mounted using dimethylhydantoin formaldehyde resin, microscope cavity slides and square cover slips. The voucher specimens and their respective DNA were submitted to the Belgian Culicidae collection housed in the Royal Belgian Institute of Natural Sciences.

DNA-based species identification was performed to confirm the morphological species identifications. Therefore, DNA was extracted from the anterior part of the larval abdomen using the NucleoSpin® Tissue extraction kit (MACHEREY-NAGEL GmbH & Co. KG, Düren, Germany) following the manufacturer's protocol, but with the elution volume set to 70 μ l. A fragment of the mitochondrial cytochrome *c* oxidase subunit I (*COI*) gene was amplified using the universal primers LCO1490 and HCO2198 (Folmer *et al.* 1994). PCR reactions and conditions, purification and sequencing were carried out as done by Ibáñez-Justicia *et al.* (2020). Finally, the *COI* consensus sequence was compared against the BLAST web application of GenBank (https://blast.ncbi.nlm.nih.gov/Blast.cgi) and compared against the Identification System of BOLD, with the Species Level Barcode Records option (www. boldsystems.org).

To proceed with the species validation of the suspected *Cx. modestus* larva, all publicly available *COI* sequences of *Culex* species known to occur in Belgium (Boukraa *et al.* 2015), and all the *COI* sequences of *Cx. modestus*, were downloaded from the BOLD online repository (Table 1; http://www.boldsystems.org/index.php/databases; November 2019). Sequences with a minimum size of 300 bp were retained, aligned using ClustalW in Geneious® v10.0.4 (Biomatters Ltd, Auckland, New Zealand) and checked for stop codons. *COI* sequences of *Coquillettidia richiardii* (Ficalbi) (KM25803, KU876993) and *Cq. fuscopennata* (Theobald) (GQ165802) were included as outgroup, and the alignment was then trimmed to only retain the 658-bp *COI* region (Folmer *et al.* 1994). Duplicate sequences were discarded per species using BioEdit v.7.2.6 (Hall 1999). Subsequently, a rooted Neighbor-Joining (NJ) tree based on Tamura-Nei distances was constructed using Geneious® v10.0.4. Nodal support was assessed by bootstrapping over 1000 replicates with a cut-off threshold of 70%.

TABLE 1. List of *Culex* species found to occur in Belgium (Boukraa *et al.* 2015), including the newly recorded *Cx. modestus*, their general occurrence in Belgium and the number of *COI* sequences (N_{COI}) downloaded from the BOLD repository (November 2019). The number of unique sequences (as defined by BioEdit) are enclosed in parentheses. Sequences were downloaded for neighbouring countries with confirmed occurrences of *Cx. modestus*. The alignment was delimited to the barcode region (Folmer *et al.* 1994).

Species	Occurrence in Belgium	Total N _{COI}	N _{COI} Belgium
Cx. territans	Rare	260 (228)	4 (4)
Cx. pipiens	Common	945 (213)	34 (10)
Cx. hortensis	Rare	18 (17)	0
Cx. torrentium	Common	340 (108)	5 (5)
Cx. modestus	First record	247 (121)	0

Species	N _{COI} neighbouring countries					
	France	Luxembourg	Germany	Netherlands	United Kingdom	
Cx. territans	0	0	1	0	0	
Cx. pipiens	0	0	424 (13)	0	146 (16)	
Cx. hortensis	1	0	15 (14)	0	0	
Cx. torrentium	0	0	282 (26)	0	36 (22)	
Cx. modestus	30 (18)	0	51 (26)	0	98 (27)	

TABLE 1. (Continued)

Results

Over the entire study period (2017–2019), one larva was morphologically identified as Cx. modestus. The larva was distinguished from other *Culex* species by the absence of lateral siphonal setae, the absence of a distinct median spine on the comb scales, the siphonal index (total length / diameter at base) between 3 and 4.5 (here 4.5) and the straight sides of the siphon. Most typical are the ventral siphonal setae, which appear in disarray (Fig. 1A), in contrast to the paired siphonal setae of the commonly collected and closely related species Cx. pipiens (Fig. 1B) and Cx. torrentium Martini (Becker et al. 2010; Harbach 2012).

The larva was collected on 18 September 2018 in a small pond close to the port of Antwerp (Kallo) in the province of East Flanders (51° 15' 11.14" N, 4° 12' 48.53" E). The vernal pond was mainly vegetated with common cattail (*Typha latifolia*), and was almost completely dried out at the time of the sampling (Fig. 2). In addition, larvae of *Culiseta annulata* Schrank (n = 1), *Anopheles maculipennis* Meigen *s.l.* (n = 1) and *Cx. pipiens* (n = 4) were collected at the same place on the same day. On 13 November 2018, the same pond was sampled again, but no mosquito larvae were found.

After sequence blasting in BOLD and GenBank, a 100% similarity was found with *COI* sequences of *Cx. modestus* from the United Kingdom (BOLD record CXOMD049-17, GenBank accession MK971827). The next 100 best matches all involved *Cx. modestus COI* sequences. The species identification was validated by the NJ tree, in which the generated consensus *COI* sequence of the Belgian specimen clustered with high confidence (81.6% bootstrap value) with other *Cx. modestus COI* sequences (Fig. 3). The *COI* barcode of the specimen was deposited in GenBank (accession MN978924). Thereby, DNA-based species identification confirmed the morphological identification of the larva.

Discussion

This is the first report of *Cx. modestus* in Belgium, based on the single larva collected at Kallo. The identification of the larva was confirmed by both morphology and *COI* barcoding. However, the species was expected to occur in Belgium based on its presence in neighbouring countries, and some suspected larvae collected at two cattle farms in Brecht and Somme-Leuze (Belgium) during 2008 and 2009 (Table 1) (Boukraa *et al.* 2011; Ries 2019; ECDC

2020). Nevertheless, in the absence of well-confirmed records it was not included in the latest checklist of the mosquito species known to occur in Belgium (Boukraa *et al.* 2015). Morphological identification of immature life stages, damaged specimens and sibling species of species complexes can quickly become challenging (Hebert *et al.* 2003; Versteirt *et al.* 2015). In these situations, DNA barcoding can support monitoring campaigns and help at providing correct species identifications.

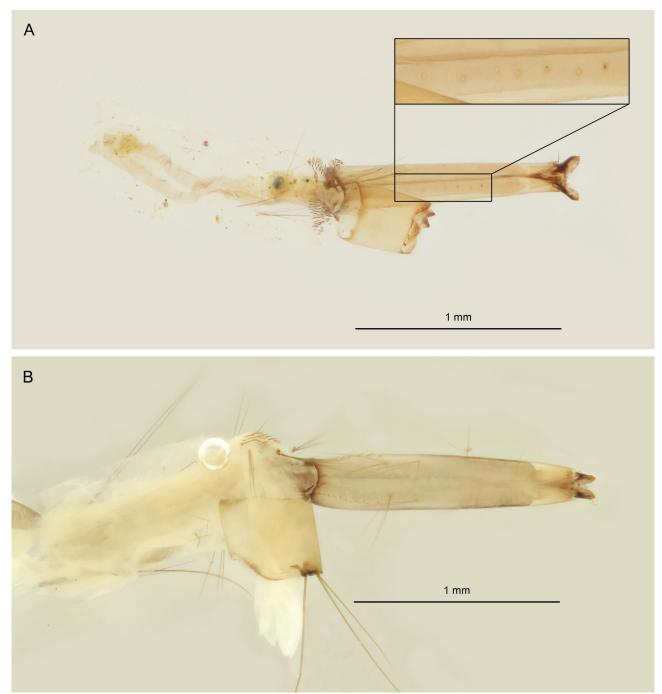


FIGURE 1. (A) Posterior part of the mounted *Cx. modestus* larva. Zoom on the diagnostic characteristic of the siphon, showing disarrayed insertion points of the ventral siphonal setae. (B) Posterior part of a mounted *Cx. pipiens* larva.

The larva was collected in a small pond in the port area of Antwerp, which is part of a protected ecological corridor network constructed in 2016. This network consists of several aquatic sites preserved for the reproduction of endangered freshwater species, e.g. the Natterjack toad *Epidalea calamita* Laurenti. The port area is rich in freshwater to slightly saline wetlands, ponds and marshes, which are the preferred habitats of *Cx. modestus*, and harbour migratory and resident bird populations (Becker *et al.* 2010; Jacobs *et al.* 2020). The larvae of *Cx. modestus* show

a preference for sunlit habitats, generally characterised by rich vegetation such as reed beds, e.g. the Camargue wetlands of France (Ponçon *et al.* 2007; Radrova *et al.* 2013).

Culex modestus was not recorded during the MODIRISK project (2007–2010), a large nationwide survey conducted to inventory and assess the distribution of mosquito species (adults) in Belgium (Versteirt *et al.* 2013). Nor were larvae of this species collected during a study conducted from 1997 until 2009 to map the distribution of mosquito larvae in Flanders (Lock *et al.* 2012). However, the latter survey focused on monitoring the water quality of streams. Therefore, as stagnant waterbodies, including marshes, ponds and wetlands were underrepresented, *Cx. modestus* may have been missed. Still, while a high mosquito nuisance was reported in the port area of Kallo in 2013, no adults of *Cx. modestus* were trapped in 2014 (Sohier & Grootaert 2015).

During the MEMO monitoring project, vegetated ponds and other bodies of ground water were not among the habitats targeted for larvae of exotic *Aedes* mosquito species. So, more targeted surveys are needed to determine the actual occurrence and distribution of *Cx. modestus* in the country. For example, the species was only discovered by focused sampling of marsh areas in England (Golding *et al.* 2012). This observation is important in light of the northward spread of WNV in Europe, which led to the detection of human cases in the Netherlands in 2020 (Vlaskamp *et al.* 2020). The increasing restoration and protection of wetlands near densely populated urban areas could create suitable habitats for potential vector species, including *Cx. modestus*, or support changes in the spatio-temporal distribution of species (Reusken *et al.* 2010; Medlock & Vaux 2015). The confirmed presence of *Cx. modestus* in Belgium confirms the trend of increased detections throughout Europe. In countries like England and the Czech Republic, studies have provided evidence of increased local abundance and spread of this previously seemingly rather rare or non-indigenous species (Radrova *et al.* 2013; Hernández-Triana *et al.* 2020).



FIGURE 2. The small vernal pond vegetated with common cattail (*Typha latifolia*) where the *Cx. modestus* larva was collected.

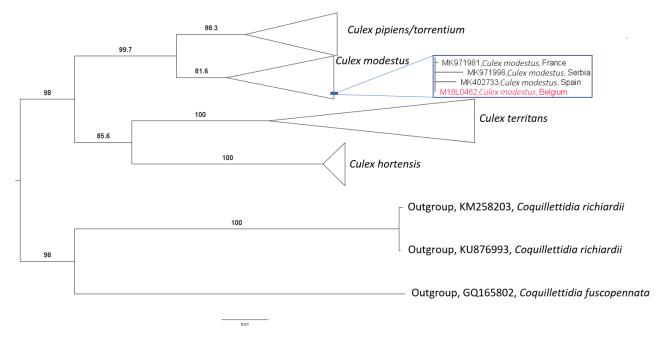


FIGURE 3. NJ tree based on *COI* sequences downloaded from BOLD, including *Culex* species recorded from Belgium (Boukraa *et al.* 2015) and sequences of *Cx. modestus* (Table 1). Bootstrap values are indicated above the branches. The blue square gives a zoom view of the un-collapsed tree.

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