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) (CDC 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).
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 μl. 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%.

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### Table 1

| Species           | Occurrence in Belgium | Total N<sub>COI</sub> | N<sub>COI</sub> 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                       |

### Table 1 (Continued)

| Species         | N<sub>COI</sub> 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)        |

### 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...
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.](image-url)
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.

Acknowledgements

We would like to thank all cooperating companies for giving access to their private property during mosquito sampling surveys. We want to thank the laboratory staff and technicians at the Institute of Tropical Medicine Antwerp for conducting the field work, especially Jacobus De Witte, Anna Schneider and Ingrid Verlé. We thank Slimane Boukraa (Unit of Entomology, Gembloux Agro-Bio Tech, ULiège), two anonymous reviewers and the editor for their input and improvements to the manuscript.

Funding

This work is part of the MEMO project (CES-2016-02 Belgium), funded by the Flemish, Wallonian and Brussels authorities and the FPS Health, Food Chain Safety and Environment within the framework of the Belgian national collaboration agreement in the domains of environment and health (NEHAP). The Outbreak Research Team of the Institute of Tropical Medicine is funded by the Department of Economy, Science and Innovation of the Flemish government. The Barcoding Facility for Organisms and Tissues of Policy Concern (BopCo) is part of the Belgian federal contribution of the Belgian Science Policy Office (Belspo) to the European Research Infrastructure Consortium LifeWatch.

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