Exploration of the *Ixodes ricinus* virosphere unveils an extensive virus diversity including novel coltiviruses and other reoviruses

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Abstract

Recent metagenomics studies have revealed several tick species to host a variety of previously undiscovered RNA viruses. *Ixodes ricinus*, which is known to be a vector for many viral, bacterial, and protozoan pathogens, is the most prevalent tick species in Europe. For this study, we decided to investigate the virosphere of Belgian *I. ricinus* ticks. High-throughput sequencing of tick pools collected from six different sampling sites revealed the presence of viruses belonging to many different viral orders and families, including Mononegavirales, Bunyavirales, Partitiviridae, and Reoviridae. Of particular interest was the detection of several new reoviruses, two of which cluster together with members of the genus Coltivirus. This includes a new strain of Eyach virus, a known causative agent of tick-borne encephalitis. All genome segments of this new strain are highly similar to those of previously published Eyach virus genomes, except for the fourth segment, encoding VP4, which is markedly more dissimilar, potentially indicating the occurrence of a genetic reassortment. Further polymerase chain reaction–based screening of over 230 tick pools for 14 selected viruses showed that most viruses could be found in all six sampling sites, indicating the wide spread of these viruses throughout the Belgian tick population. Taken together, these results illustrate the role of ticks as important virus reservoirs, highlighting the need for adequate tick control measures.

Key words: *Ixodes ricinus*; tick; virus discovery; Eyach virus; Coltivirus; reovirus; Belgium

1. Introduction

Recent metagenomics studies have revealed the extensive virus diversity within arthropods (Li et al., 2015; Shi et al., 2016). Of particular interest amongst arthropods are the haematophagous vectors. These animals feed on the blood of vertebrate hosts, presenting the ideal conditions for the transmission of pathogens. In addition to mosquitoes, ticks are some of the most important vectors of infectious diseases, in humans, livestock, and wildlife animals (Mansfield et al., 2020). Ticks are known to carry a diverse range of pathogens, including protozoa, bacteria, helminths, and viruses, although in Europe they are predominantly known for their ability to transmit *Borelia burgdorferi* bacteria, the causative agent for Lyme disease (Bush and Vazquez-Pertejo 2018; Boulanger et al., 2019; Madison-Antenucci et al., 2020). However, ticks also act as vectors for multiple viral pathogens, including tick-borne encephalitis virus (TBEV), the most prevalent viral tick-borne zoonotic agent in Europe, Crimean-Congo haemorrhagic fever virus, and Colorado tick fever virus (Shi et al., 2018). In the last decade, many articles have been published looking at the virus diversity in different tick species in different regions, revealing the presence of many previously undiscovered viruses (Tokarz et al., 2014, 2018; Xia et al., 2015; Moutailler et al., 2016; Pettersson et al., 2017; Harvey et al., 2019; Meng et al., 2019). Although estimating the relative risk of these novel viruses for human, animal, and plant health is difficult, their phylogenetic clustering within specific virus families and genera that are known to harbour tick-borne human, animal, and plant pathogens, such as *Reoviridae*, *Nairoviridae*, *Flaviviridae*, or *Phenuiviridae*, hints at the pathogenic potential for at least some of them.

Ticks are divided into three families: the *Argasidae* or ‘soft ticks’, the *Ixodidae* or ‘hard ticks’, and the *Nuttalliellidae*. Unlike the latter, which contains only one species, the families *Argasidae* and *Ixodidae* harbour several hundred member species, split across multiple genera (Guglielmone et al., 2010). In Belgium, only a limited number of tick species have been described, and almost all reported tick bites can be attributed to the castor bean tick or *Ixodes ricinus*, the type species of the genus *Ixodes* in the family of ‘hard ticks’ (Obsomer et al., 2013; Lernout et al., 2019).
This prominently European tick species requires three blood meals during its life cycle: to molt from larva to nymph and from nymph to adult, and to lay eggs once it has reached adulthood (Walker 2019). This triple-host life cycle makes I. ricinus an important vector for the transmission of several tick-borne diseases between a variety of mammalian hosts, including humans. Several viruses that pose a threat to human and animal health are already known to be transmitted by I. ricinus ticks, although, with the exception of TBEV, studies concerning human diseases of viral origin spread by I. ricinus are limited (see Hubalek and Rudolf, 2012, for a detailed overview). Furthermore, the ongoing discovery of new (putative) viral pathogens, such as the newly described Aologne virus, which was recently reported to be present in European I. ricinus ticks, indicates that I. ricinus ticks might harbor more zoonotic viruses than hitherto assumed (Kuivamen et al., 2019).

In recent years, a number of studies have delved deeper into the virus diversity in I. ricinus in Europe, revealing the presence of several previously undetected viruses (Mouttaker et al., 2016; Pereira et al., 2017; Petersson et al., 2017; Kuivamen et al., 2019; Ohlendorf et al., 2019). In 2017, we reported the discovery of a novel nairovirus, Guroenhout virus, in Belgian I. ricinus, which was identified through next-generation sequencing of a pool of adult ticks (Vanmechelen et al., 2017). Because we also detected traces of other viruses, we decided to sequence additional tick pools to further characterize the virus diversity in Belgian ticks. Here, we report the detection of several novel as well as previously described viruses, including a rabdovirus and multiple reoviruses. Additionally, we performed polymerase chain reaction (PCR) screening for 14 different viruses, including the novel viruses reported here as well as some other viruses known to be carried by I. ricinus, on more than 200 tick pools from six sampling locations in Belgium, providing a broad overview of virus prevalence in Belgian I. ricinus ticks.

2. Materials and methods

2.1 Sample collection

Ticks were collected between 2009 and 2017 from six sampling locations in Belgium (Torhout [51°4′0.105″N, 3°3′25.9″E], Moerbeke [51°11′39.5″N, 3°55′4.3″E], Zoersel [51°14′41.8″N, 4°40′49″E], Gierle [51°17′19.3″N, 5°53′8.2″E], Chimay [50°0′1″N 4°14′52.5″E], and Heverlee [50°51′27.5″N 4°39′56.4″E]), by flagging the low-growing vegetation using a 1-m-by-2-m white cotton flag. Ticks from Gierle, Chimay, and Heverlee were kept at −20°C for long-term storage, while ticks from Torhout, Moerbeke, and Zoersel were stored at room temperature in 100 per cent ethanol. A total of 767 ticks were collected, including 33 adults, 513 nymphs, and 221 larvae. All ticks were sorted based on their location of origin and their developmental stage, and all were identified as I. ricinus based on phenotypical characteristics. An overview of the different sampling locations and the number of ticks caught at each location can be found in Fig. 1.

2.2 RNA extraction

Each adult tick or pool (1–10) of nymphs/larvae (see Supplementary Table S1) was added to a tube containing 100 µl phosphate buffered saline and zirconium oxide beads and subsequently homogenized for 150 s at 4,000 g in a Bertin Minilys homogenizer. Total RNA was extracted from the homogenate using the RNeasy Mini Kit (QIAGEN, Leiden, Netherlands) according to the manufacturer’s instruction. DNA carry-over was further depleted by DNase treatment using 2U TurboDNase (Invitrogen, ThermoFisher Scientific, CA, USA), and the remaining RNA was re-purified using the RNeasy MinElute Cleanup Kit (QIAGEN, Leiden, Netherlands).

2.3 Attempt at virus isolation

Virus rescue was attempted using African green monkey kidney cells (Vero E6, American Type Culture Collection, C1008) and hamster kidney fibroblasts (BHK21). Vero E6 cells were maintained in Dulbecco’s Modified Eagle Medium (Thermo Fisher Scientific, MA, USA) supplemented with 10 per cent fetal bovine serum (FBS; Biowest, France), 1 per cent sodium bicarbonate, 1 per cent penicillin-streptomycin-1-glutamine, 1 per cent gentamicin, and 0.2 per cent fungizone (all Thermo Fisher Scientific). BHK21s were kept in Minimal Essential Medium Rega-3 (Thermo Fisher Scientific), supplemented with +10 per cent FBS and 200 mM 1-glutamine (Thermo Fisher Scientific). To attempt virus rescue, cells were seeded in 6-well plates and allowed to grow overnight. The medium was removed from the resulting monolayers, which were subsequently inoculated with 200 µl fresh tick homogenate for 1 h, after which 2 ml 2 per cent FBS-supplemented medium was added. Cells were observed daily for cytopathogenic effect for 2 weeks, and the cell medium was refreshed every 3–4 days. Collected supernatant was subjected to RNA extraction using the RNeasy Mini Kit (QIAGEN), followed by reverse transcription PCR (RT-PCR) screening as described below.

2.4 Nanopore sequencing

RNA extracted from a single female adult tick was converted to complementary DNA (cDNA) using the Complete Whole Transcriptome Amplification Kit (WTA2, Sigma-Aldrich, Saint Louis, MO, USA). Purification of cDNA was done using the MSB Spin PCRapace kit (Stratec Molecular, Berlin, Germany). The resulting cDNA was prepared for nanopore sequencing using the SQK-LSK108 kit (Oxford Nanopore Technologies, Oxford, UK) and loaded on an R9.4.1 flow cell. Reads corresponding to tick ribosomal RNA (rRNA) were removed, and following size selection, reads between 200 and 2,000 nucleotides were assembled using Canu v1.5 (Karen et al., 2017). Identification of the resulting contigs was done using BLAST (https://blast.ncbi.nlm.nih.gov/Blast.cgi).

2.5 Illumina sequencing

Ten pools were made for Illumina sequencing, containing three to five RNA extracts each. An overview of the ten pools can be found in Table 1. RNA was converted to cDNA using the WTA2 Kit (Sigma-Aldrich). cDNA was purified using the MSB Spin PCRapace kit (Stratec Molecular). Sequencing libraries were prepared using the Illumina NexteraXT DNA Library Prep Kit (Illumina, San Diego, USA). Libraries were double indexed using Set A-barcoded adapters (Illumina, San Diego, USA). In-house modifications on the standard library preparation were conducted following Conceicao-Neto et al. (2018). Sequencing was performed on a HiSeq4000 platform (Illumina, San Diego, USA) at VIB Nucleomics Core (Leuven, Belgium). CLC Genomics Workbench (v10.1.1) was used to trim the reads by adapter content, quality (Phred q-score >35), and length (>25 bp) and to assemble the resulting trimmed reads (default settings). Diamond (v0.9.24) and KronaTools (v2.7.1) were used to classify the obtained contigs, using the blastx algorithm against the complete NCBI non-redundant protein database (nr; GenBank release 239) (Ondov, Bergman, and Phillippy 2011; Buchfink, Xie, and Huson 2015). The HMMER v3.2.2 web server (www.hmmer.org) was used to identify divergent virus segments that could not be classified by Diamond.
Figure 1. Tick sampling sites. Ticks were collected from six sampling sites in Belgium, marked in red and scaled according to the number of ticks caught at each location. Dark grey, white, and light grey lines represent province, arrondissement, and municipality borders, respectively.

Table 1. Illumina sequencing data overview.

| Pool number | RNA pools used | #Ticks | Adult/nymph/larva | Location          | #Reads    | #Viral reads | #Contigs | #Viral contigs |
|-------------|----------------|--------|-------------------|-------------------|-----------|--------------|----------|---------------|
| S1          | 142 144        | 147    | 4 Adults          | Moerbeke          | 62,886,446| 977          | 4,874    | 13            |
| S2          | 4 6           | 87     | 4 Adults          | Chimay Gierle     | 65,898,424| 1,576        | 6,918    | 16            |
| S3          | 140 162       | 168    | 3 Adults          | Heverlee Torhout  | 48,976,538| 6,621        | 25,558   | 24            |
| S4          | 143 146       | 149    | 3 Adults          | Moerbeke          | 68,520,456| 12,952       | 65,273   | 65            |
| S5          | 89 90         | 95     | 3 Adults          | Gierle            | 68,966,000| 750,760      | 6,771    | 94            |
| S6          | 129 159       | 161    | 3 Nymphs 4 Larvae | Heverlee Moerbeke| 63,439,638| 149,654      | 74,219   | 693           |
| S7          | 47 48 49      | 51     | 12 Nymphs         | Chimay            | 76,725,768| 7,301        | 15,224   | 66            |
| S8          | 107 110       | 114    | 12 Nymphs         | Chimay Gierle     | 58,129,700| 366,933      | 53,572   | 132           |
| S9          | 106 110       | 112    | 15 Nymphs         | Gierle            | 83,201,118| 293,361      | 35,840   | 842           |
| S10         | 39 44 45      | 135    | 13 Nymphs         | Chimay Heverlee   | 70,097,012| 47,741       | 15,718   | 96            |

See Supplementary Table S1.
2.6 PCR screening and genome completion

The Qiagen OneStep RT-PCR kit was used to screen for the different viruses and to complete gaps in the obtained genome sequences. The following cycling conditions were used: 50°C for 30 min, 95°C for 15 min, 40 cycles of 94°C for 30 s, a primer-specific annealing temperature for 30 s and 72°C for 60 s, and a final extension step at 72°C for 10 min. Completion of genome ends was done using the Roche 5'/3' RACE kit, 2nd generation. An overview of all used screening primer sets and annealing temperatures can be found in Supplementary Table S2. PCR products were visualized by 2 per cent agarose gel electrophoresis and Sanger sequenced to confirm their identity. Sequencing was done by Macrogen Europe (Macrogen Europe B.V., Amsterdam, The Netherlands) after the products were purified in-house with PureIT ExoZAP (Ampliqon, Odense, Denmark).

2.7 Phylogenetic analysis

For the newly discovered rhabdovirus and reoviruses, multiple sequence alignments were made based on deduced amino acid sequences of the RNA-dependent RNA polymerase (RdRp) sequence of members of the families Rhabdoviridae and Reoviridae. For the rhabdovirus phylogenetic tree, amino acid sequences for all currently recognized International Committee of the Taxonomy of Viruses species, supplemented with all unclassified rhabdoviruses >5,500 bp (NCBI: txid35303), were downloaded from NCBI-Nucleotide (https://www.ncbi.nlm.nih.gov/nucleotide). Segkit was used to keep only protein sequences >1,800 amino acids, which were aligned using MAFFT (v7.310; localpair algorithm) (Katoh and Standley 2013; Shen et al., 2016). Based on this alignment, the closest related sequences to Chimay rhabdovirus (and Rabies virus as an outgroup) were realigned using MAFFT. The resulting alignment was trimmed using TrimAl (v1.4.rev15; gappyout setting), and model selection and phylogenetic tree calculation were done using IQ-TREE (v1.6.12), employing 1,000 bootstrap replicates (Capella-Gutierrez, Silla-Martinez, and Gabaldon 2009; Nguyen et al., 2015). The resulting tree was visualized using Figtree v1.4.3. For the reovirus tree, the RdRp protein sequence was deduced from all (putative) members of the subfamily Spinareovirinae for whom an RdRp-encoding segment could be identified. Alignment generation, trimming and phylogenetic tree calculation, and visualization were performed as described above. An overview of the accession numbers of all used sequences can be found in Supplementary Table S4.

3. Results

3.1 Detection of multiple viruses in a single tick

We reported previously the discovery of a novel nairovirus, Grotenhout virus, in a pooled library of ten adult tick RNA extracts (Vanmechelen et al., 2017). Even though we recovered a complete L and S segment from the Illumina data, we found no traces of an M segment. However, we did find one 267-bp contig that resembled part of a phlebovirus segment. In an attempt to obtain more sequence information for this phlebovirus and to simultaneously look for the missing Grotenhout virus M segment, we screened all ten RNA extracts used to make the illumina pool described in Vanmechelen et al. (2017) by RT-PCR for Grotenhout virus and the novel phlebovirus and selected one dual-positive extract (see Supplementary Table S1, extract 7) for RNA sequencing using the Oxford Nanopore MinION. This run yielded ~4.6 million reads, covering ~1.81 Gb. Following rRNA removal and length trimming (keeping reads between 200 and 2,000 bp), assembly using Canu (v1.5) yielded 31 contigs. Together, these contigs cover the full length of both the Grotenhout virus L and S segments as well as the L and S segments of the aforementioned phlebovirus, which we provisionally named Leuven phlebovirus (GenBank ID: MG407659-MG407660). Genome sequences similar to this virus (98.5–99.1 per cent nucleotide identity) have recently also been discovered in northern European ticks, under the name Norway phlebovirus 1 (Pettersson et al., 2017). All four segments had excellent coverage (see Table 2), but for neither virus, a contig matching a putative M segment was detected. The remaining contigs could all be classified as being of host or bacterial origin, except for three contigs that displayed limited similarity with members of the family Rhabdoviridae and appeared to be part of the same novel rhabdovirus. Correction by Sanger sequencing and RACE was used to obtain the complete 13,706-bp genome sequence for this virus, which was given the name Chimay rhabdovirus (GenBank ID: MF975531). The genome has the typical rhabdovirus genome architecture, encoding five open reading frames (ORFs): 3'-N-P-M-G-L-5' and clusters alongside Blacklegged tick rhabdovirus 1 and IRE CTVM19-associated rhabdovirus, both tick-derived viruses (Fig. 2). Chimay rhabdovirus was recently classified as a separate species (Chimay betaricinrhavirus) in a novel genus (Betaricinrhavirus) in the family Rhabdoviridae.

3.2 Tick sample set

The finding of three separate viruses in the RNA isolated from a single tick prompted us to further investigate the viral diversity in Belgian ticks. Therefore, we decided to collect additional ticks of different developmental stages and from different locations in Belgium and used next-generation sequencing for the detection of known and novel viruses in these ticks. Ten tick pools, made from pooling three to five RNA extracts, were sequenced on an Illumina 4000 HiSeq, yielding 48,976,538–83,201,118 reads per pool. Variable numbers of ticks from mixed locations of origin were used in the different pools to maximize the overall observed diversity. However, this approach also means that the pools should not be compared quantitatively with each other, as such comparisons would be biased due to the differences in sample handling and pooling. Following adapter trimming, between 4,874 and 74,219 contigs could be assembled for each pool (Table 1). Although the fraction of viral reads compared to host material was low for most pools, we did detect reads and contigs of many different viruses, with strong heterogeneity between pools (Fig. 3). While some pools were largely dominated by the

Table 2. Tick extract 7 nanopore data.

| Virus ID                        | Segment length (bp) | #Contigs | Merged contig length (bp) | #Reads | Average coverage |
|--------------------------------|---------------------|----------|---------------------------|--------|------------------|
| Grotenhout virus L segment     | 14,854              | 6        | 14,806                    | 38,615 | 1.433X           |
| Grotenhout virus S segment     | 3,728               | 3        | 3,716                     | 27,061 | 3.564X           |
| Leuven phlebovirus L segment   | 6,730               | 1        | 6,699                     | 6,769  | 580X             |
| Leuven phlebovirus S segment   | 2,501               | 1        | 2,467                     | 9,816  | 1,806            |
| Chimay rhabdovirus            | 13,706              | 3        | 13,562                    | 3,210  | 125X             |
Figure 2. Phylogenetic clustering of Chimay rhabdovirus. Maximum-likelihood tree based on the protein sequence of the RdRp of Chimay rhabdovirus and the closest-related (putative) members of the family Rhabdoviridae, including Rabies virus as an outgroup. Chimay rhabdovirus clusters near Blacklegged tick rhabdovirus-1 and IRE CTVM19-associated rhabdovirus, both tick-derived viruses. The scale bar indicates the number of amino acid substitutions per site. The numbers at the node indicate the bootstrap support for each node, based on 1,000 replicates. Only values <100 are shown. All used accession numbers are provided in Supplementary Table S4.

presence of one specific virus (Pool 5), others displayed strong diversity, containing contigs of viruses belonging to many different families. Interestingly, while most of the virus families identified in our data are known to contain tick-borne viruses, we also observed fungus/plant-specific viral families (most notably Botourmiaviridae, Chrysoviridae, Endornaviridae, Hypoviridae, Polymoviridae, and Quadriviridae) especially in Pool 6. This pool also displayed the largest diversity, but this is most likely attributable to the presence of fungi in the sample used for RNA extraction. Whether these fungi were infecting the ticks in this sample or
Figure 3. Abundance of viral families in the different Illumina pools. Fraction of mapped viral reads per pool, colour-coded based on the taxonomical classification (according to KronaTools) of the closest available reference, as determined by DIAMOND. Virus families representing <1 per cent of the virus reads in a sample are grouped as ‘Other’. Virus families known to be families of bacteriophages are grouped together as ‘Phage’, as they are unlikely to represent true tick-associated viruses.

should be considered environmental contaminants remains to be determined.

### 3.3 Reconstruction of viral genomes

While for most detected viruses only fragments of the genome were retrieved, some were present more abundantly, allowing the reconstruction of near-complete genome sequences. These include Grotenhout virus and the aforementioned Leuven phlebovirus, which have also recently been observed in *I. ricinus* in Norway (Pettersson et al., 2017). With the exception of Norway luteo-like virus 2, all viruses that were discovered in this Norwegian study are also present in our data, although only Bronnoya virus, Norway partiti-like virus 1 and Norway luteo-like virus 1 had sufficient read support to recover contigs covering most of the genome. These contigs showed high nucleotide identity (>90 per cent) to the genomes of the Norwegian strains and are not further discussed here. Lastly, we also managed to retrieve near-complete genome sequences of three reoviruses. These include a novel strain of Eyach virus, named strain Heverlee (GenBank ID: MW874081-MW874092), as well as two previously undescribed viruses, named Gierle tick virus (GenBank ID: MW874093-MW874104) and Zoersel tick virus (GenBank ID: MW874105-MW874114), respectively. For Eyach virus strain Heverlee, a combination of Illumina and Sanger sequencing was used to obtain the complete sequence of all 12 genome segments. While almost all segments are highly similar (>94 per cent nucleotide identity) to previously identified strains of Eyach virus (Table 3), the fourth segment, encoding VP4, is markedly dissimilar, displaying <78 per cent nucleotide similarity. No other reovirus segments were detected in samples positive for this strain of Eyach virus. Akin to the Eyach virus, we managed to identify all 12 genome segments of Gierle tick virus, a putative new viral species in the genus *Coltivirus* most similar to Tarumizu tick virus and Kundal virus (Fig. 4). Lastly, we also managed to retrieve the near-complete genome of a third reovirus, Zoersel tick virus. Although markedly dissimilar from all currently classified reoviruses, this virus shares some similarity with the unclassified reoviruses High Island virus, Eccles virus, and ‘Reoviridae sp. BF02/7/10’, and the unclassified Riboviria Shelly beach virus and Hubei diptera virus 21. Together, these five viruses form a
The notably lower identity values for segment 4 are marked in bold.
Figure 4. Reovirus phylogeny and genome organization. Left: genome organization of Eyach virus strain Heverlee, Gierle tick virus, and Zoersel tick virus, with segment lengths drawn according to scale. Ribbed lines indicate incomplete genome ends. The deduced amino acid length of all proteins is shown next to the corresponding segments. *Incomplete ORFs. Right: maximum-likelihood tree based on the protein sequence of the RdRp of all the (putative) members of the subfamily Spinareovirinae for whom such a sequence is available, shows Eyach virus strain Heverlee and Gierle tick virus clustering within the genus Coltivirus, with Gierle tick virus representing a novel species. Zoersel tick virus forms a separate branch within the subfamily alongside several yet unclassified viruses. The scale bar indicates the number of amino acids substitutions per site. The numbers at the node indicate the bootstrap support for each node, based on 1,000 replicates. Only values <100 are shown. All used accession numbers are provided in Supplementary Table S4.
Perhaps, the most interesting discovery is the finding of two members of the genus Coltivirus. This genus currently contains five member species, although putative additional species have been reported (Attoui et al., 2002; Gao et al., 2020; Harvey et al., 2019; Walker et al., 2020). Two of its members, Colorado tick fever virus and Eyach virus, are known human pathogens capable of causing severe disease (Attoui et al., 2005). While the other members, Tarumizu tick virus, Tai Forest reovirus, and Kundal virus, have not been implicated in human disease, they have been shown to be capable of infecting mammalian and/or human cell lines (Fujita et al., 2017; Weiss et al., 2017; Yadav et al., 2019). One of the two coltiviruses reported here appears to be a strain of Eyach virus, sharing >94 per cent nucleotide similarity for all but one segments. To our knowledge, this marks the first detection of Eyach virus or any known viral human pathogen in the Belgian tick population, further expanding the known geographical range in which this virus is found (see Chastel 1998). Interestingly, the fourth segment is markedly dissimilar from known Eyach virus sequences, indicating that some form of reassortment has occurred, hinting at the existence of other, yet to be discovered coltiviruses.

In addition to two coltiviruses, we also identified two reoviruses not belonging to any of the currently recognized genera in the family Reoviridae. While we only found three partial segments for one of them, Torhout tick virus appears to be distantly related to Fennes virus, a virus recently identified in ticks found on Antarctic penguins (Wille et al., 2020). Conversely, a near-complete genome was reconstructed for Zoersel tick virus. Phylogenetically, this virus is most closely related to a group of viruses for which only six genome segments are known, and, to a lesser extent, the ten-segment containing Operophtera brumata reovirus. Interestingly, several of the ten segments we identified for Zoersel tick virus share (almost) no sequence similarity with any known sequences. Without the minimal sequence similarity between segments 5, 7, and 10 and the corresponding segments of Operophtera brumata reovirus (NC_007559-NC_007568), identifying all ten segments would have been significantly more difficult. Thanks
to the availability of this reference, the erroneous conclusion that Zoersel tick virus has only six genome segments could be avoided. The high degree of sequence divergence of the additional segments of Zoersel tick virus illustrates the likely possibility that its closest known relatives might also have additional segments that are highly divergent. Such divergence could leave certain segments undiscovered by typical BLAST-based approaches, a concern previously raised by some of the groups that discovered these putative six-segmented viruses (Sadeghi et al., 2017; Medd et al., 2018; Temmam et al., 2019). Re-analysing these viruses’ sequencing sets with the help of additional related sequences, such as the ones presented here, might help identifying potentially missed segments.

For several of the viruses we detected in our sequencing data, highly similar or related viruses have previously been reported in different tick species from various locations, strengthening the hypothesis that these viruses are truly tick-associated. As noted by Pettersson and colleagues, this association can sometimes be difficult to claim based only on sequencing data, especially if the viruses in question are only found infrequently or in low abundance (Pettersson et al., 2017). Intriguingly, two of the viruses for which they note such low abundance, Norway mononegavirus 1 and Norway luteo-virus 4, are also some of the least-detected viruses in our sample set, although we did find traces of both. Their detection in multiple unrelated datasets might suggest that they are simply low abundant tick-associated viruses, rather than environmental contaminants. In addition to previously identified viruses, we also detected a variety of novel viruses in our data. For some of these viruses, (near)-complete genomes could be reconstructed (see above), albeit most were only present in low abundance, resulting in only small fragments of their genomes being recovered. The low abundance and associated lack of available sequence data make it difficult to confidently classify these viruses as tick-associated, although the similarity of some of them with members of virus families and orders known to contain tick-borne viruses (Rhabdoviridae, Phenuiviridae, Chuviridae, Orthomyxoviridae, and Mononegavirales) do seem to support this association.

For many other viruses of which we find traces, which only share similarities with unclassified viruses or viruses belonging to virus lineages with mixed hosts, the origin remains largely unknown. Nonetheless, they do provide further evidence that virus diversity in ticks (and other arthropods) is much higher than until recently known. Additional studies are needed to further map this diversity, allowing a better understanding of tick-associated virus evolution. A better characterization of the virus diversity in tick populations can also facilitate studying the spread of known and novel zoonotic viruses, allowing more targeted pre- and post-exposure countermeasures for tick-borne diseases to be taken where needed.

In conclusion, we present the first study looking at virus diversity in I. ricinus from Belgium. We show that Belgian ticks carry a wide variety of viruses and that several of these viruses are found in different locations and different tick developmental stages. We also managed to reconstruct (near)-complete genome sequences for several novel viruses, including a new strain of the pathogenic Eyach virus, a new coltivirus species, and a highly divergent reovirus that remains to be classified. This research further illustrates that many tick-associated viruses remain to be discovered and that additional studies are needed to expand our knowledge of virus diversity in ticks and to further characterize these novel viruses.

Data availability
Virus sequences generated in this study have been submitted to GenBank under accession numbers MW874081-MW874117.

Supplementary data
Supplementary data is available at Virus Evolution online.

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