Boginia virus, a newfound hantavirus harbored by the Eurasian water shrew (*Neomys fodiens*) in Poland

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Abstract

**Background:** Guided by decades-old reports of hantaviral antigens in the Eurasian common shrew (*Sorex araneus*) and the Eurasian water shrew (*Neomys fodiens*) in European Russia, we employed RT-PCR to analyze lung tissues of soricine shrews, captured in Boginia, Huta Dłutowska and Kurowice in central Poland during September 2010, 2011 and 2012.

**Findings:** In addition to Seewis virus (SWSV), which had been previously found in Eurasian common shrews elsewhere in Europe, a genetically distinct hantavirus, designated Boginia virus (BOGV), was detected in Eurasian water shrews captured in each of the three villages. Phylogenetic analysis, using maximum likelihood and Bayesian methods, showed that BOGV formed a separate lineage distantly related to SWSV.

**Conclusions:** Although the pathogenic potential of BOGV and other recently identified shrew-borne hantaviruses is still unknown, clinicians should be vigilant for unusual febrile diseases and clinical syndromes occurring among individuals reporting exposures to shrews.

**Keywords:** Hantavirus, Neomys, Sorex, Shrew, Phylogeny, Poland

Findings

Although hantaviral antigens were detected by the indirect immunofluorescent antibody test and enzyme immunoassay in tissues of the Eurasian common shrew (*Sorex araneus*), Eurasian pygmy shrew (*Sorex minutus*) and Eurasian water shrew (*Neomys fodiens*) in European Russia and the former Yugoslavia more than two decades ago [1-3], shrews (order Soricomorpha) had been dismissed as being unimportant in the ecology and evolution of hantaviruses. Guided by these long-ignored reports, and aided by access to archival tissue collections, an opportunistic search, employing reverse transcription-polymerase chain reaction (RT-PCR), demonstrated a genetically distinct hantavirus, named Seewis virus (SWSV), in a Eurasian common shrew captured in Graubünden, Switzerland [4].

Subsequent studies have indicated that SWSV is widespread across the vast distribution of its soricid reservoir, in Austria, Czech Republic, Slovakia, Finland, Germany, Hungary and Russia [5-7]. SWSV has also been detected in the Siberian large-toothed shrew (*Sorex daphaenodon*) and tundra shrew (*Sorex tundrensis*) in Siberia [6]. And in a comprehensive study on the phylogeography of SWSV in central Europe, SWSV was shown to exhibit distinct geographic-specific clustering in Eurasian common shrews and Eurasian pygmy shrews [7]. The overlapping geographic ranges of the Eurasian common shrew and other shrew species in central Poland provided an opportunity to investigate the existence of SWSV and other soricine shrew-borne hantaviruses.

Shrews were captured, using live traps and pitfalls traps, placed 5 m apart and baited with raw bacon or beef, during the month of September in 2010, 2011 and/or 2012 in Boginia (51°20′26.80 N, 19°36′41.36 E), Huta Dłutowska (51°35′49.51 N, 19°22′46.80 E) and/or...
Kurowice (51°39′48.03 N, 19°42′20.92 E), three closely located villages within Łódź East County, in central Poland (Figure 1). All trapping and experimental procedures on shrews were approved by the General Directorate for Environmental Protection (DOP-OZGiZ. 4200/N2732/10/JRO, DOP-OZGiZ.6401.05.25.2011kp.3 and DOP-OZGiZ.6401.05.28.2011kp.1).

Total RNA, extracted from lung tissues preserved in RNAlater® RNA Stabilization Reagent (Qiagen) from 40 shrews (Table 1), was analyzed for hantavirus RNA by RT-PCR, using oligonucleotide primers designed from soricomorph-borne hantaviruses [5,8-10]. Using Clustal W [11], pair-wise alignment and comparison of the 795- and 782-nucleotide regions of the M and L segments, respectively, revealed a genetically distinct hantavirus, named Boginia virus (BOGV), in 3 of 6 Eurasian water shrews captured in Boginia, Huta Dłutowa and Kurowice (Tables 1 and 2). The high prevalence of BOGV infection in Eurasian water shrews captured in three separate villages in central Poland suggested a well-established reservoir host-hantavirus relationship.

The L-segment genetic variation between BOGV strains 2073, 2074 and 2177 was 3.1–15.0% at the nucleotide level, but there was 100% amino acid homology. The observation that hantaviruses exhibit high sequence conservation at the amino acid level, despite considerable divergence of more than 15% at the nucleotide level, has been made previously [5-7,12]. This has been attributed to the long-standing co-adaptation between virus and reservoir host, as well as the strong selection pressure to preserve function of the three gene products.

BOGV and other representative soricine shrew-borne hantaviruses exhibited M- and L-segment sequence similarity of only 71.7–76.7% and 75.1–87.7% at the nucleotide and amino acid levels, respectively. The taxonomic identities of the BOGV-infected Eurasian water shrews were verified by mitochondrial DNA sequence analysis (GenBank accession no. KC537796 for BOGV 2073; KC537797 for BOGV 2074; and KC537798 for BOGV 2177).

Eurasian common shrews, captured in Boginia and Huta Dłutowa, as well as a Eurasian pygmy shrew from Kurowice, were infected with SWSV (Table 1), indicating simultaneous circulation of genetically distinct hantaviruses in syntopic shrew species captured in the same trap sites. Recently, SWSV was detected in Eurasian pygmy shrews in Germany and the Czech Republic [7], but the low prevalence is suggestive of spillover. Nevertheless, carriage of the same hantavirus in two soricine shrew species in central Poland parallels findings of host sharing of SWSV in other Sorex shrew species [6,7], as well as of rodent-borne hantaviruses, such as Tula virus, in several arvicolid rodent species [12-14].

Phylogenetic trees, generated by maximum likelihood and Bayesian methods, implemented in PAUP* (Phylogenetic Analysis Using Parsimony, 4.0b10) [15],

### Table 1 RT-PCR detection of hantavirus RNA in tissues of soricine shrews from Poland

| Genus species     | Capture site     | Capture dates       | Number tested | Number positive |
|-------------------|------------------|---------------------|---------------|-----------------|
| *Sorex araneus*   | Boginia          | September 20–22, 2010 | 3             | 1               |
|                   |                  | September 20–22, 2011 | 6             | 0               |
|                   | Huta Dłutowa     | September 15–18, 2011 | 9             | 2               |
| *Sorex minutus*   | Boginia          | September 21–22, 2011 | 2             | 0               |
|                   | Huta Dłutowa     | September 16–18, 2011 | 4             | 0               |
|                   | Kurowice         | September 7–14, 2012 | 10            | 1               |
| *Neomys fodiens*  | Boginia          | September 22, 2011  | 1             | 1               |
|                   | Huta Dłutowa     | September 22, 2011  | 1             | 1               |
|                   | Kurowice         | September 9–14, 2012 | 4             | 1               |
RAxML Blackbox webserver [16] and MrBayes 3.1 [17], under the best-fit GTR + I + Γ model of evolution established using jModeltest 0.1.1 [18], showed that BOGV formed a separate lineage distantly related to SWSV (Figure 2), in keeping with the evolutionary relationship of their soricid hosts. As expected, phylogenetic trees showed SWSV to segregate generally along geographic-specific lineages (Figure 2), as reported previously [5-7].

As for several other shrew-borne hantaviruses, the sequence divergence of BOGV presented insurmountable difficulties in obtaining the full genome sequence. Sequencing efforts were also severely constrained by the limited availability of tissues, and poor-quality tissue RNA thwarted efforts at employing next-generation sequencing technology. Efforts, now underway, to rapidly freeze tissues collected from Eurasian water shrews captured in central Poland will be used for virus isolation attempts.

The detection of a novel, genetically distinct hantavirus in the Eurasian water shrew captured in central Poland confirms decades-old reports that this shrew species might serve as a reservoir [2]. Whether other Neomys species, such as the Mediterranean water shrew (Neomys anomalus), also harbor a BOGV-like hantavirus, rather than SWSV, is unknown. Nevertheless, neither physical proximity with sharing of habitats nor genetic relatedness of shrew host species allows accurate prediction of the hantavirus species in a particular reservoir. That said, this study provides clear evidence of the co-existence of two genetically distinct hantaviruses in their soricomorph reservoir host species inhabiting the same ecological niche. To what extent hantavirus spillover occurs between reservoir shrews and rodents is currently under investigation.

Unlike most soricine shrew species, the Eurasian water shrew is comparatively large, measuring approximately 10 cm in body length, with a long tail. It is highly terrestrial and lives a solitary life near freshwater and other wetland habitats, including rivers, streams, marshes, bogs and damp grasslands and meadows, at sea level to elevations over 2,500 m [19,20]. Eurasian water shrews forage almost exclusively underwater, using to great effect their venomous saliva to feast on aquatic invertebrates, including insects, gastropods and crustaceans, as well as occasionally small vertebrates, such as amphibians and fish. Some terrestrial insects, such as dipteran larvae, are also consumed. The vast distribution of the Eurasian water shrew [19,20], spanning from the United Kingdom and Scandinavia throughout Europe and much of Asia, including far eastern Russia and Sakhalin, where other syntopic soricomorphs and rodents reside, provides rich opportunities to investigate the genetic diversity and phylogeography of BOGV.

That distinct diseases or clinical syndromes have yet to be attributed to shrew-borne hantaviruses is not particularly surprising, given that most hantaviruses are nonpathogenic. Neotomine and sigmodontine rodents were known to harbor hantaviruses long before the terrifying outbreak of a rapidly progressive, frequently fatal disease (now known as hantavirus cardiopulmonary syndrome) signaled the highly pathogenic nature of these hantaviruses [21-23]. Similarly, while the pathogenic potential of BOGV and other recently identified, still-orphan soricid-borne hantaviruses is still unknown, active vigilance by physicians and public health workers for unusual febrile syndromes, occurring among individuals reporting either contact with shrews or exposure to wetlands that may be contaminated by the Eurasian water shrew, would be crucial to uncover an etiological association. Intensive investigations are underway to isolate and further characterize BOGV to ascertain its impact on human health.

### Table 2 Partial S-, M- and/or L-segment sequences of hantaviruses detected in lung tissues of soricine shrews captured in central Poland

| Virus strain | Capture site  | Capture date | S segment (bp) | M segment (bp) | L segment (bp) | GenBank accession number |
|-------------|-------------|-------------|--------------|--------------|--------------|------------------------|
| SWSV 1107   | Boginia     | 22-Sep-2010 | 690 (435–1025) | JX990921     |              |                        |
| SWSV 2048   | Huta Dłutowa | 16-Sep-2011 | 647 (1520–2100) | JX990967     | 806 (2520–3400) | JX990936               |
| SWSV 2049   | Huta Dłutowa | 17-Sep-2011 |              |              | 756 (2541–3296) | JX990944               |
| SWSV 2121   | Kurowice    | 12-Sep-2012 |              |              | 357 (2900–3300) | JX990945               |
| BOGV 2073   | Huta Dłutowa | 22-Sep-2011 |              |              | 316 (2998–3313) | KC537794               |
| BOGV 2074   | Boginia     | 22-Sep-2011 |              |              | 394 (2520–2935) | JX990964               |
| BOGV 2177   | Kurowice    | 9-Sep-2012  |              |              | 795 (1520–2355) | JX990966               |
|             |             |             |              |              | 783 (2541–3323) | JX990965               |
|             |             |             |              |              | 408 (2520–2935) | KC537795               |

Abbreviations: BOGV, Boginia virus; SWSV, Seewis virus; nt, nucleotides.

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Phylogenetic tree generated by the maximum-likelihood method, using the GTR + I + Γ model of evolution, based on the alignment of L-segment sequences of BOGV strains 2073 (JX990964), 2074 (JX990965) and 2177 (KC537795). The phylogenetic positions of BOGV are shown in relationship to representative soricine shrew-borne hantaviruses, including Cao Bang virus (CBNV CBN-3, EF543525), Ash River virus (ARRV MSB73418, EF619961), Jemez Springs virus (JMSV MSB144475, FJ593501) and Qiandao Lake virus (QDLV YN05-284, GU566021). Representative strains of Seewis virus (SWSV) from Poland and other widely separated geographic regions are indicated by their country of origin and GenBank numbers. Also shown are crocidurine shrew-borne hantaviruses, including Thottapalayam virus (TPMV VRC6412), Imjin virus (MJIN 05–11, EF641806), Tanganya virus (TGNV Tan826, EF050454), Azagury virus (AZGV KBM15, JF276228) and Jeju virus (JJUV 10–11, HQ834697); and mole-borne hantaviruses, including Asama virus (ASAV N10, EU29078), Nova virus (NVAV MSB95703, FJ593498) and Oxbow virus (OXBV Ng1453, FJ593497). Rodent-borne hantaviruses include Hantaan virus (HTNV 76–118, NC_005222), Seoul virus (SeOV 80–39, NC_005238), Tula virus (TULV MS302v, NC_005226), Puuramala virus (PUUV Sotkamo, NC_005225), Prospect Hill virus (PHV PH-1, EF46763), Sin Nombre virus (SNV NMH10, NC_005217) and Andes virus (ANDV Chile9717869, AF291704). The numbers at each node are bootstrap values, expressed as percentages, based on 1,000 iterations, and the scale bar indicates nucleotide substitutions per site.
Competing interests
The authors declare that they have no competing interests.

Authors' contributions
SHG and HJK performed primer design, RNA extraction, RT-PCR and DNA sequencing reactions and phylogenetic analysis. JM and JH conducted trapping expeditions and collected shrew tissues. BS and PPL provided logistical support. RT conceived the project and provided overall scientific oversight. All authors contributed to the preparation of the final manuscript. All authors read and approved the final manuscript.

Acknowledgements
This work was supported in part by U.S. Public Health Service grants R01AI075057 from the National Institute of Allergy and Infectious Diseases and P20GM103516 from the National Institute of General Medical Sciences, National Institutes of Health, and by grant N N303 604538 from the State Committee for Scientific Research, Ministry of Scientific Research and Technology Information in Poland.

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Received: 3 March 2013 Accepted: 10 May 2013 Published: 22 May 2013

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