Seoul Virus in Rats
(Rattus norvegicus),
Hyesan, North
Korea, 2009–2011

To the Editor: Seoul virus (SEOV), a member of the family Bunyaviridae, genus Hantavirus, is primarily carried by Rattus norvegicus rats. Because members of Rattus species are widely distributed, SEOV has the potential to cause human disease worldwide. It has been reported that SEOV causes a milder form of hemorrhagic fever with renal syndrome than Hantaan virus and Dobrava-Belgrade virus and is responsible for 25% of cases of hemorrhagic fever with renal syndrome in Asia (1). Although it is well known that SEOV is endemic to China (2) and South Korea (3), little is known about its distribution in North Korea (4).

In September 2009, June and September 2010, and September 2011, a total of 89 R. norvegicus rats were trapped in the city of Hyesan (128°30′E, 41°30′N) during the operation of a cooperative rodent surveillance program of China and North Korea. The captured rodents were euthanized with barbiturate (100 mg/kg), weighed, measured, classified by sex, and then autopsied. Lung samples were probed for the large segment of SEOV by reverse transcription PCR by using the RT primer P14 (5), the primary PCR primers HAN-L-F1 and HAN-L-R1, and the nested PCR primers HAN-L-F2 and HAN-L-R2 (6). PCR products were sequenced by using an ABI 3730 sequencer (Applied Biosystems, Foster City, CA, USA).

A high rate of SEOV infection was detected in R. norvegicus rats; 15 (16.8%) of 90 rodent samples tested positive for SEOV by reverse transcription PCR. Infection rates at each surveillance time were 26.7% (4/15) in September 2009, 7.5% (3/40) in June 2010, 28.6% (6/21) in September 2010, and 15.4% (2/13) in September 2011. All infected R. norvegicus rats were adults; 9 were male and 6 were female. The rate of nucleotide substitution in these 15 SEOV amplicons (330 bp; GenBank accession nos. KC576788–KC576802, JX853574) was calculated by Bayesian Markov chain Monte Carlo analysis using BEAST 1.74 (7). The mean substitution rate, calculated by using the uncorrelated lognormal distribution relaxed molecular clock model and a Bayesian skyline model for the large segment of SEOV, was 8.27 × 10⁻³ substitutions/site/year, with a 95% high posterior density interval that ranged from 1.02 × 10⁻² to 1.79 × 10⁻². This substitution rate is about 3 times greater than that for middle and small segments (2).

Phylogenetic relationships were assessed by using the uncorrelated lognormal distribution relaxed molecular clock model with the SRD06 substitution model (8) in BEAST 1.74. The Hantaan virus strain AA57 (GenBank accession no. AB620033) sequence was used as the outgroup. The resulting phylogenetic tree (Figure) showed that SEOV strains in the city of Hyesan shared >97.3% identity and were all clustered in their own lineages, subdivided into 2 co-existing sublineages. Although the geographic distance from Hyesan to northeastern China (e.g., Liaoning Province) is much less than that between northeastern and southeastern China (e.g., Zhejiang Province) or central China (e.g., Hubei Province), the phylogenetic distance between SEOV strains in North Korea and those in each location in China in clade A, calculated by using MEGA5.1 (9), was 0.03, but was only 0.01–0.02 between locations in China.

One possible explanation for this discrepancy in phylogenetic and geographic distances between SEOV strains in China and those in North Korea may be differences in the extent of human contact. Although human interactions among different regions of China are extensive, by comparison, those between China and North Korea are considerably reduced for political reasons. In addition, combining with small segment (GenBank accession no. HQ992815) sequence analysis (data not shown), the fact that SEOV strain L0199 from Laos were not clustered in clade A-D(2) showed that Laos was another possible area of origin for SEOV.

Our work contributes to the known epidemiology of exposure to the SEOV pathogen in Hyesan. Hyesan adjoins Changbai County in Jilin Province of China. However, SEOV was not detected in Changbai County during the surveillance program (data not shown), which was consistent with previous research (10). This study further highlights the need for long-term surveillance.

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Schmallenberg Virus Infection in Dogs, France, 2012

To the Editor: In 2011, Schmallenberg virus (SBV) emerged in Europe (1); the virus spread into France in January 2012 (2). During January–March 2012, a total of >1,000 cases were reported in France, mainly in stillborn and newborn lambs with congenital malformations.

In March 2012, neurologic disorders were detected in five 15-day-old puppies (Belgian shepherd) from a dog breeding kennel in northwestern France (Orne). We report data suggesting that these puppies were infected with SBV.

In June 2012, the kennel veterinarian contacted a veterinary school (Unité de Médecine de l'Elevage et du Sport, Maisons-Alfort, France) after neurologic signs of ataxia, exotropia, and imbalance were observed in five 15-day-old puppies. The puppies were born to a 16-month-old female Belgian shepherd dog and were housed in a separate room from adult dogs that did not show any signs of pathology.

The puppies were initially observed by veterinarians at 6 weeks of age. At 16 weeks of age, they were examined by a veterinarian who noted ataxia and exotropia. The puppies were euthanized, and a necropsy was performed.

Histologic examination of the brain revealed inflammation and gliosis in the cerebellar cortex, especially in the granular layer. Immunohistochemical staining revealed the presence of SBV in the cerebellum.

The results of this study suggest that SBV can cause neurologic disorders in puppies. Further studies are needed to determine the long-term effects of SBV infection on the development of the central nervous system in puppies.

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