

# FACT SHEET

## Recombinant Sendai Viral Vectors

The following provides information on the use and containment of recombinant Sendai viral vectors. Investigators should use these guidelines as part of their risk assessment when planning experiments with these vectors and preparing applications to the Institutional Biosafety Committee (IBC). Note the listed containment levels are the minimum that should be employed with these vectors: some experiments, such as the expression of toxins or oncogenes, may require higher levels of containment. The appropriateness of the containment should be considered as part of the investigator's risk assessment and will be reviewed by the IBC.

---

**NIH Risk Group**

RG2

---

**Biocontainment Level**

BSL-2

Sendai virus (SeV) causes respiratory disease in rodents and sometimes swine. There is limited evidence of zoonotic transmission to humans. However, the virus is capable of infecting human cell lines (<http://jvi.asm.org/content/84/22/11718.full>) and is similar to human parainfluenza virus type 1. For these reasons, SeV work is usually classified as BSL-2.

Recombinant constructs expressing oncogenes or toxins should be handled at BSL-2 enhanced

---

**Infectious to Humans/Animals**

Mice

---

**Route of Transmission**

SeV is responsible for a highly transmissible respiratory tract infection in mice, hamsters, guinea pigs, rats, and occasionally pigs, with infection passing through both air and direct contact routes.

---

**Laboratory Hazards**

No reported cases of laboratory acquired disease but inhalation of aerosolized droplets, mucous membrane contact, parenteral inoculation, or ingestion are possible routes of infection.

---

**Disease**

Respiratory disease. Infections of mice are usually associated with a high mortality rate although latent infections can occur.

---

**Treatment/Prophylaxis**

Antivirals may reduce shedding

---

**Pathogenesis**

The respiratory infection of Sendai virus in mice is acute. Virus may first be detected in the lungs 48 to 72 hours following exposure. As the virus replicates in the respiratory tract of an infected mouse, the concentration of the virus grows most quickly during the third day of infection. After that, the growth of the virus is slower but consistent. Typically, the peak concentration of the virus is on the sixth or seventh day, and rapid decline follows that by the ninth day. A fairly vigorous immune response mounted against the virus is the cause of this decline.

---

**Replication Competent**

Yes

**RCV Testing**

No

**Disinfection**

Effective disinfectants require a minimum of 20 minutes contact time. Use one of the following:

- RECOMMENDED: Sodium hypochlorite (0.5%: use 1:10 dilution of fresh bleach)
  - 5% Phenol
  - 70% Ethanol or Isopropanol
- 

**Animals**

- ABSL-3: Animal cages must be labeled with a biohazard sign. Note there are currently no ABSL-3 suites at the University of Utah.
- 

Sources:

[http://web.stanford.edu/dept/EHS/prod/researchlab/bio/docs/Working\\_with\\_Viral\\_Vectors.pdf](http://web.stanford.edu/dept/EHS/prod/researchlab/bio/docs/Working_with_Viral_Vectors.pdf)

[http://www.dartmouth.edu/~ehs/biological/biosafety\\_docs/110\\_1\\_ibc\\_viral\\_vector\\_policy.pdf](http://www.dartmouth.edu/~ehs/biological/biosafety_docs/110_1_ibc_viral_vector_policy.pdf)



125 South Fort Douglas Blvd, Salt Lake City, UT 84113

801.581.6590 | [ehs.utah.edu](http://ehs.utah.edu)