

**Hantavirus Pulmonary Syndrome (HPS)
Specimen Submittal Instructions**

Guidelines for Submission

- Fill out as completely as possible:
 - VRDL GENERAL PURPOSE SPECIMEN SUBMITTAL form
 - HPS CASE HISTORY FORM
 - Fax both forms to the Medical Epidemiology Liaison Section (MELS) at (510) 307-8599 AND send a copy with the specimen(s) to avoid delays in testing.
- Collect two tubes and send on cold packs (**It is very important to use an overnight delivery service because the EDTA samples will begin to degrade within three days**)
 - One 5 ml tube in EDTA (purple top)
 - One 10 ml whole clotted blood (red top).
- Since the incidence of HPS is rare in California, we recommend that you also submit a respiratory specimen (nasopharyngeal swabs or washes, tracheal aspirates, bronchoalveolar lavage, and/or pleural fluid) for viral isolation and/or respiratory PCR assays to test for other agents that may be causing your patient's illness.
- Save all specimens (including hematology differential slides) from the patient until HPS serology has been completed. Additional samples may be tested if the patient is deceased.
 - Paraffin embedded lung and kidney tissues- Ship and store at ambient temperature
 - Fresh or frozen lung and kidney- Ship and store at -70°C

HPS Consultation

- If you would like to consult about a possible HPS patient, call the Infectious Disease Branch at (916) 552-9730 or call the Medical and Epidemiology Liaison Section (MELS) for the VRDL at (510) 307-8585. If neither is available, local health departments may contact the Duty Officer at (510) 620-3434.
- Clinical consultations for patient management are available from the staff at the University of New Mexico Medical School. Call 1-888-866-7257 and request a HPS consultation.
- In cases where clinical presentation is not consistent with VRDL HPS test results, or VRDL HPS results are equivocal, specimens may be forwarded to a reference laboratory for further testing.

**Screening Criteria for Hantavirus Pulmonary Syndrome
in Persons with Unexplained Respiratory Illness ***

* MMWR October 28, 1993 pp 816-820

Potential case-patients must have one of the following:

- A febrile illness (temperature ≥ 101 F or ≥ 38.3 C) occurring in a previously healthy person characterized by unexplained adult respiratory distress syndrome (ARDS)
- Bilateral interstitial pulmonary infiltrates developing within one (1) week of hospitalization with respiratory compromise requiring supplemental oxygen
- Unexplained respiratory illness resulting in death in conjunction with an autopsy examination demonstrating non-cardiogenic pulmonary edema without an identifiable specific cause of death
- Thrombocytopenia along with elevated hematocrit and high WBC with immunoblasts in the smear is characteristic of patients suspected to be infected with hantavirus.

Potential case-patients are to be excluded if they have any of the following:

- An acute illness that provides a likely explanation for the respiratory illness (unless there is history of recent potential rodent exposure) such as:
 - Recent major trauma, burn, surgery, recent seizures or history of aspiration
 - Bacterial sepsis
 - Another respiratory disorder such as respiratory syncytial virus in young children, influenza, or legionella pneumonia

Confirmed case-patients must have the following:

- Compatible clinical history of illness

AND

- Detection of Immunoglobulin M (IgM) antibodies or a significant (i.e., fourfold or greater) rise in hantavirus-specific Immunoglobulin G (IgG) antibody titers
OR
- Detection of hantavirus-specific nucleic acid amplification testing (NAAT) in an appropriate clinical specimen
OR
- Detection of hantavirus antigen by immunohistochemistry (IHC)

Hantavirus Pulmonary Syndrome Case History Form

Patient Id. (assigned by State Lab)

Please return with Specimen Submittal Form to:

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|---|---|--|--|--|--|--|--|

Viral and Rickettsial Disease Laboratory

ATTN: Specimen Receiving

850 Marina Bay Parkway

Richmond, CA 94804

Phone (510) 307-8585

Fax (510) 307-8599

--FIPS-- --YR-- -----CA # -----

| | | | |
|----------------------------------|------|--|---------------------------|
| Patient's Last Name, First Name | | Middle Name: | Patient's Mailing Address |
| Date of Birth: ____/____/____ | Age: | Sex: M F | Occupation: |
| County Health Jurisdiction: | | Race/Ethnicity: <input type="checkbox"/> White <input type="checkbox"/> Black <input type="checkbox"/> Asian/Pacific Islander <input type="checkbox"/> American Indian/Alaska Native <input type="checkbox"/> Hispanic <input type="checkbox"/> Non-Hispanic <input type="checkbox"/> Unknown | |

Date of Onset and Hospitalization History

| | |
|-----------------------|--|
| Onset Date: | Was patient hospitalized for this illness? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unk |
| Name of Hospital: | |
| Location of Hospital: | |
| Dates in Hospital: | ____/____/____ to ____/____/____ |
| MR# | |

Clinical Signs, Symptoms and Laboratory Values

| | | | | |
|---|-------------------------|--------------------------|-------------------------|-------------------------------|
| Did the patient have any of the following? (Circle) | | | Additional Information: | |
| Fever > 101F or > 38.3 C: | Yes | No | Unk | Highest fever: |
| Thrombocytopenia (platelets ≤ 150,000 mm): | Yes | No | Unk | Lowest platelet count: |
| Elevated hematocrit (Hct): | Yes | No | Unk | Highest Hct: |
| Elevated creatinine: | Yes | No | Unk | Highest creatinine: |
| CXR with unexplained bilateral interstitial infiltrates or Suggestive of ARDS? | Yes | No | Unk | Date Performed: |
| Oxygen saturation < 90% at any time? | Yes | No | Unk | |
| Was patient intubated? | Yes | No | Unk | Date Performed: |
| Has patient received ribavirin? | Yes | No | Unk | |
| WBC: | Total Neutrophils: % | Banded neutrophils: % | Lymphocytes: % | Atypical Lymphocytes: % |
| History of any relevant underlying medical conditions (i.e. COPD, malignancy, immunosuppression, diabetes)? | | | | |
| Other possible explanations for acute illness (i.e. sepsis, burns, trauma)? | | | | |
| History of rodent exposure in 6 weeks prior to illness? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unk | | | | |
| Date of Exposure to known direct or indirect contact with rodents or their excreta : ____/____/____ | | | | |
| Type of Rodent: _____ | | | | |
| Place of Exposure: _____ | | | | |
| Outcome of Illness? <input type="checkbox"/> Alive <input type="checkbox"/> Dead (if deceased, date of death) ____/____/____ <input type="checkbox"/> Unk | | | | |
| If deceased, was an autopsy performed? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | |
| Evidence of non-cardiogenic pulmonary edema? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | |
| Available Samples: Serum/blood <input type="checkbox"/> Yes (date collected ____/____/____) <input type="checkbox"/> No | | | | |
| Fresh frozen or paraffin tissue blocks <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | |
| Has a specimen been tested for hantavirus infection at another lab? <input type="checkbox"/> Yes <input type="checkbox"/> No | | | | |
| If yes then Name of lab and append a copy of the results: | | | | |
| Comments: | | | | |