

Name of Hospital: _____

Location of Hospital: _____

Dates in Hospital: ___/___/___ to ___/___/___

Record Number: _____

Did the patient have any of the following?

Fever >101 F or >38.3 C:	Yes	No	Unk.	Highest Fever: _____
Thrombocytopenia (platelets <150,000 mm ³):	Yes	No	Unk.	Lowest platelet count: _____
Elevated Hematocrit (Hct):	Yes	No	Unk.	Highest Hct: _____
Elevated creatinine:	Yes	No	Unk.	Highest creatinine: _____
WBC: _____ Lymphocytes: _____ (%)		Total Neutrophils: _____ (%)	Banded Neutrophils: _____ (%)	
CXR with unexplained bilateral interstitial infiltrates or suggestive of ARDS?	Yes	No	Unk.	Date: ___/___/___
Respiratory compromise requiring supplemental oxygen?	Yes	No	Unk.	
Oxygen saturation <90% at any time?	Yes	No	Unk.	
Was the patient intubated?	Yes	No	Unk.	Date: ___/___/___
Has the patient received ribavirin?	Yes	No	Unk.	

History of any relevant underlying medical conditions (i.e. COPD, malignancy, immunosuppression, diabetes)?

Other possible explanations for acute illness (i.e. sepsis, burns, trauma)?

Outcome of illness?	Alive	Dead	Unk.	If deceased, date of death: ____/____/____
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Was an autopsy performed?	Yes	No	Unk.	
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If yes, was exam compatible with non-cardiogenic pulmonary edema?	Yes	No	Unk.	
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Are tissue specimens (fresh-frozen or paraffin blocks) available for testing?	Yes	No	Unk.	
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Is serum/blood specimen available for testing for hantavirus infection?	Yes	No	Unk.	
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Has a specimen been tested for hantavirus infection at another laboratory?	Yes	No	Unk.	
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If yes, where? _____	Type of specimen? _____	Results (i.e. titer, OD)? _____
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History of any rodent exposure in 6 weeks prior to onset of illness?	Yes	No	Unk.
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If yes, date of contact: ____/____/____

Mouse Rat Other: Unk.

Type of rodent:

Place of Contact (town, county, state): _____

Comment:

State Health Dept. reporting case: _____

State/local ID Number:

Date form completed: ___/___/___

Person completing report: _____

Phone number: (____)____-_____

Name of patient's physician: _____

Phone number: (____)____-_____

Public reporting burden of this collection of information is estimated to average 20 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Reports Clearance Officer; 1600 Clifton Road NE, MS D-24, Atlanta, GA 30333; ATTN: PRA (0920-0009). Centers for Disease Control and Prevention
Revised August 2002