

## Virginia Department of Health Plague: Overview for Healthcare Providers

Organism	Yersinia pestis are small gram-negative, rod-shaped bacteria (safety pin appearance)
	belonging to the family Enterobacteriaceae
Reporting to Public Health	Suspected or confirmed cases require <u>immediate</u> notification to the local health department
Infectious Dose	(LHD). See https://www.vdh.virginia.gov/health-department-locator/
	<100 colony-forming units by inhalation
Occurrence	Plague occurs worldwide, but primarily in sub-Saharan Africa      In the United States and successful and the United States and successful and the United States and successful and the United States and St
	• In the United States, plague is rare (1–17 cases/year), with most cases reported in the
	southwest and western regions
	Plague occurs naturally in the Western U.S. (e.g., Arizona, Colorado, New Mexico, and     Idada a naturally in the Western U.S. (e.g., Arizona, Colorado, New Mexico, and
Noticed December	Utah); it does not occur naturally in Virginia
Natural Reservoir	Primarily rodents (e.g., ground squirrels, prairie dogs, chipmunks, wood rats, deer mice and value) and their floor.
	voles) and their fleas
Davida of Infantion	Rabbits, hares, wild carnivores, or domestic cats can also be infection sources
Route of Infection	Bite of infected flea
	Unprotected handling of infected animal tissues or body fluids, or laboratory specimens
	Respiratory droplets from people or animals with plague pharyngitis or pneumonic plague
	Aerosolized bacteria could be used for bioterrorism attack, resulting in pneumonic plague
Communicability	Infected fleas can remain infectious for months under suitable environmental conditions
	Bubonic plague and septicemic plague are not usually transmitted from person to person
	unless there is direct contact with infected body fluids or tissues
	Pneumonic plague is usually highly communicable by respiratory droplets within a close
	distance (< 6 feet) when patient is symptomatic and has had less than 48 hours of
	appropriate antibiotic therapy. Patients are usually no longer infectious after 48 hours of
	appropriate antibiotic treatment with clinical improvement.
	Domestic cats can develop pneumonic plague by eating infected rodents and pose a risk of
Diala fa atawa	transmitting infectious plague droplets to their owners or veterinarians
Risk factors	Traveling to plague-endemic areas (e.g., sub-Saharan Africa)
	Handling infected animals (e.g., veterinarians, hunters, pet owners) or plague cultures (e.g.,
	laboratorians)
	Camping or hiking in areas where plague-infected animals or fleas reside
Case-Fatality Rate	Untreated bubonic plague is 50–60%; septicemic and pneumonic plague invariably fatal
	• ~11% in U.S. with treatment. Fatality rate might be higher in developing countries.
Incubation Period	1–3 days for pneumonic plague, 2–8 days for bubonic plague, poorly defined for septicemic
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Clinical Description	There are 3 main clinical forms of plague: bubonic, pneumonic, and septicemic
	Bubonic plague: acute onset of fever, headache, chills, weakness, and painful swollen
	lymph nodes (buboes), most commonly in the inguinal, axillary, or neck region. Nausea,
	vomiting, and diarrhea are common. Untreated bubonic plague can lead to septicemic
	plague and pneumonic plague.
	Pneumonic plague: fever, headache, weakness, and productive cough (possibly bloody).      Desired to the second of the seco
	Rapid development of pneumonia with dyspnea, chest pain, stridor, cyanosis, and
	respiratory failure.
	Septicemic plague: fever, chills, extreme weakness, abdominal pain, shock, and bleeding     into the abit and other property of the second party into the abit and other property.
	into the skin and other organs. Respiratory distress and gastrointestinal symptoms (nausea,

	vomiting, and diarrhea) might be present. Can progress rapidly to septic shock,
	intravascular coagulopathy, meningitis, or coma.
	Rare forms include pharyngeal, meningitic, cutaneous, ocular, and gastrointestinal
Differential	Bubonic: cat scratch disease (Bartonella), ulceroglandular tularemia, adenitis due to
Diagnosis	staphylococcal, streptococcal, or filarial infection, tuberculosis, nontuberculosis mycobacterial infection, lymphogranuloma venereum, <i>Capnocytophaga canimorsus</i> infection, chancroid, strangulated inguinal or femoral hernia, lymphadenopathy due to nonspecific infections, appendicitis, cellulitis
	<ul> <li>Pneumonic: Other bacterial pneumonia (<i>Mycoplasma, Legionella, Staphylococcus, Streptococcus, Haemophilus, Klebsiella</i>) and viral pneumonia (influenza, respiratory syncytial virus, hantavirus, severe acute respiratory syndrome), Q fever, inhalation anthrax, tularemia</li> <li>Septicemic: Other gram-negative sepsis and gram-positive sepsis (<i>Staphylococcus</i>),</li> </ul>
	meningococcemia, rickettsial infections, malaria
Radiography	Pulmonary infiltrates or consolidation on chest radiograph for pneumonic plague
Specimen Collection	If plague is suspected, notify LHD immediately to discuss the case and laboratory testing
and Laboratory	• Specimens may be sent to Division of Consolidated Laboratory Services (DCLS) after LHD
Testing	has been consulted and testing has been approved by VDH
	• Pre-treatment specimens should be collected, if possible, but treatment should not be delayed.
	• Do not wait for diagnostic test results if plague is suspected; confirmatory diagnosis can be established later with specialized lab tests. Disease can progress rapidly to death without appropriate antibiotic therapy.
	<ul> <li>Specimens to test depend on clinical presentation but could include culture isolates submitted on slants; lymph node aspirate (≥1 mL) for culture; lower respiratory tract specimen (&gt;1 ml for induced sputum, tracheal aspirate, BAL, or pleural fluid) for culture; acute and convalescent serum (&gt;2 ml each) for serology.</li> <li>For questions about specimen collection, contact the DCLS Emergency Officer 24/7 at 804-</li> </ul>
	335-4617
Treatment	<ul> <li>Begin appropriate therapy as soon as plague is suspected</li> <li>Gentamicin and fluoroquinolones are first-line treatments in the United States</li> <li>Information on choice of drugs, dosing, and duration of treatment for adults, children, and pregnant women is available at <a href="https://www.cdc.gov/plague/healthcare/clinicians.html">https://www.cdc.gov/plague/healthcare/clinicians.html</a></li> <li>For additional information on dosing, please consult the drug package inserts</li> </ul>
Preexposure	If standard and droplet precautions can be maintained, no need for PrEP for individuals
Prophylaxis (PrEP)	caring for pneumonic plague patients
	<ul> <li>In cases of mask shortages, patient overcrowding, poor ventilation, or other situations,</li> <li>PrEP might be warranted if there are sufficient antimicrobial supplies</li> </ul>
	Information on PrEP for children and adults is available at
	https://www.cdc.gov/plague/healthcare/clinicians.html
Postexposure Prophylaxis (PEP)	PEP is indicated in persons with known exposure to plague, such as close (<6 ft), sustained contact with a pneumonic plague patient, or direct contact with infected body fluids or tissues
	<ul> <li>Information on PEP for children and adults is available at https://www.cdc.gov/plague/healthcare/clinicians.html</li> </ul>
Vaccine	A vaccine for plague is not commercially available in the United States
Infection Control	Use Standard Precautions for all types of plague
infection Control	<ul> <li>Ose standard Precautions for all types of plague</li> <li>Patients with pneumonic signs should also be isolated and placed on Droplet Precautions until patient has received at least 48 hours of effective antibiotic therapy.</li> </ul>
	and patient has received at reast to hours of effective untilbiotic therapy.