

# ANTHRAX INVESTIGATION FORM

**STOP: PRIOR TO CREATING THIS INVESTIGATION, YOU MUST NOTIFY & CONSULT WITH CENTRAL OFFICE  
(800) 338-8374 (24-HOUR COVERAGE)**

## BASIC DEMOGRAPHIC DATA

Last Name: \_\_\_\_\_ First Name: \_\_\_\_\_ Middle Name: \_\_\_\_\_

DOB: \_\_\_\_/\_\_\_\_/\_\_\_\_ Age: \_\_\_\_ ☐ years ☐ months Current Sex: ☐ Female ☐ Male ☐ Unknown

Is the patient deceased? ☐ No ☐ Unknown ☐ Yes Date of Death: \_\_\_\_/\_\_\_\_/\_\_\_\_

Street Address 1: \_\_\_\_\_ Street Address 2: \_\_\_\_\_

City: \_\_\_\_\_ State: \_\_\_\_\_ Zip Code: \_\_\_\_\_ County: \_\_\_\_\_

Home Phone: (\_\_\_\_) - \_\_\_\_ - \_\_\_\_ Cell Phone: (\_\_\_\_) - \_\_\_\_ - \_\_\_\_ Work Phone: (\_\_\_\_) - \_\_\_\_ - \_\_\_\_ Ext. \_\_\_\_\_

Ethnicity: ☐ Hispanic or Latino ☐ Not Hispanic or Latino ☐ Unknown

Race: ☐ American Indian/Alaska Native ☐ Asian ☐ Black/African American ☐ Native Hawaiian/Other Pacific Islander ☐ White ☐ Unknown

## INVESTIGATION SUMMARY

Investigation Start Date: \_\_\_\_/\_\_\_\_/\_\_\_\_ Investigation Status: ☐ Open ☐ Closed Investigator: \_\_\_\_\_

## REPORTING SOURCE

Date of Report: \_\_\_\_/\_\_\_\_/\_\_\_\_ Reporting Source: \_\_\_\_\_

## CLINICAL

Physician's Name: \_\_\_\_\_ Phone Number: (\_\_\_\_) - \_\_\_\_ - \_\_\_\_ Ext. \_\_\_\_\_

Was patient hospitalized for this illness? ☐ No ☐ Unknown ☐ Yes If yes: Hospital Name: \_\_\_\_\_

Admission Date: \_\_\_\_/\_\_\_\_/\_\_\_\_ Discharge Date: \_\_\_\_/\_\_\_\_/\_\_\_\_ Duration of Stay \_\_\_\_\_ day(s)

Diagnosis Date: \_\_\_\_/\_\_\_\_/\_\_\_\_ Illness Onset Date: \_\_\_\_/\_\_\_\_/\_\_\_\_ Illness End Date: \_\_\_\_/\_\_\_\_/\_\_\_\_

Age at Onset: \_\_\_\_\_ ☐ days ☐ hours ☐ minutes ☐ months ☐ unknown ☐ weeks ☐ years

Did the patient die from this illness? ☐ No ☐ Unknown ☐ Yes Date of Death: \_\_\_\_/\_\_\_\_/\_\_\_\_

## EPIDEMIOLOGIC

Is this patient associated with a day care facility? ☐ No ☐ Unknown ☐ Yes Is this patient a food handler? ☐ No ☐ Unknown ☐ Yes

Is this case part of an outbreak? ☐ No ☐ Unknown ☐ Yes If yes, outbreak name: \_\_\_\_\_

Case Status: ☐ Confirmed ☐ Not a Case ☐ Probable ☐ Suspect ☐ Unknown MMWR Week: \_\_\_\_\_ MMWR Year: \_\_\_\_\_

## ADMINISTRATIVE

General Comments: \_\_\_\_\_

## PHA4 SUPERVISOR REVIEW

Date Due: \_\_\_\_/\_\_\_\_/\_\_\_\_ Investigation ready for supervisor review: ☐ Reviewed (Complete) ☐ Reviewed (Incomplete)

Date investigation ready for supervisor review: \_\_\_\_/\_\_\_\_/\_\_\_\_ ☐ Reviewed (Not a case) ☐ Yes

Review comments (completed by supervisor): \_\_\_\_\_

**CONTACT ATTEMPTS**

Physician Contact Date(s):

1<sup>st</sup> Attempt: \_\_\_/\_\_\_/\_\_\_\_\_2<sup>nd</sup> Attempt: \_\_\_/\_\_\_/\_\_\_\_\_3<sup>rd</sup> Attempt: \_\_\_/\_\_\_/\_\_\_\_\_

Patient Contact Date(s):

1<sup>st</sup> Attempt: \_\_\_/\_\_\_/\_\_\_\_\_ Time: \_\_\_\_\_ ☐ AM ☐ PM2<sup>nd</sup> Attempt: \_\_\_/\_\_\_/\_\_\_\_\_ Time: \_\_\_\_\_ ☐ AM ☐ PM3<sup>rd</sup> Attempt: \_\_\_/\_\_\_/\_\_\_\_\_ Time: \_\_\_\_\_ ☐ AM ☐ PM

Regular Letter Mailed: \_\_\_/\_\_\_/\_\_\_\_\_

Certified Letter Mailed: \_\_\_/\_\_\_/\_\_\_\_\_

Was clinical information obtained from the physician or patient? ☐ Yes ☐ No**SIGNS AND SYMPTOMS**Did the patient have a fever ( $\geq 100.4^{\circ}\text{F}$ )? ☐ No ☐ Unknown ☐ YesWas the patient septic? ☐ No ☐ Unknown ☐ Yes**Cutaneous Anthrax:**☐ Eschar, location: \_\_\_\_\_ ☐ Lymphadenopathy ☐ Malaise ☐ Papule that became vesicular**Inhalation Anthrax:**☐ Acute respiratory distress ☐ Dyspnea (short of breath) ☐ Mediastinal widening ☐ Shock  
☐ Cyanosis ☐ Hypoxia ☐ Pleural effusion ☐ Viral respiratory-like illness**Intestinal Anthrax:**☐ Abdominal swelling ☐ Bloody Diarrhea ☐ Nausea ☐ Vomiting  
☐ Anorexia ☐ Hematemesis (bloody vomit) ☐ Severe abdominal pain**Oropharyngeal Anthrax:**☐ Cervical adenopathy ☐ Edema ☐ Painless oral mucosal lesion ☐ Pharyngitis**Meningeal Anthrax:**☐ Coma ☐ Convulsions ☐ Meningeal signs (e.g., meningitis)**EXPOSURES**

What is the patient's primary occupation? \_\_\_\_\_ Name and location of employer: \_\_\_\_\_

In the 7 days prior to onset of symptoms did patient have exposure to or contact with any of the following?

Livestock: ☐ No ☐ Unknown ☐ Yes Date: \_\_\_/\_\_\_/\_\_\_\_\_ Location: \_\_\_\_\_Animal skins, fur, or hair: ☐ No ☐ Unknown ☐ Yes Date: \_\_\_/\_\_\_/\_\_\_\_\_ Location: \_\_\_\_\_**CASE CLASSIFICATION**

1	Did the patient have an acute illness, or post-mortem examination, revealing a form of clinical anthrax? <input type="checkbox"/> Cutaneous - a skin lesion evolving over 2-6 days from a papule, to vesicular stage, to depressed black eschar; <input type="checkbox"/> Inhalation - a viral respiratory-like illness followed by hypoxia, dyspnea, or acute respiratory distress resulting in cyanosis and shock; <input type="checkbox"/> Gastrointestinal - severe abdominal pain with swelling, nausea, vomiting, hematemesis, bloody diarrhea, anorexia, fever, and septicemia; <input type="checkbox"/> Oropharyngeal - a painless mucosal lesion in the oral cavity or oropharynx, fever, edema, pharyngitis, cervical adenopathy, and possibly septicemia; <u>or</u> <input type="checkbox"/> Meningeal - fever, convulsions, coma, or meningeal signs (often occurs secondary to other forms of anthrax).	<input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> Yes
2	Was a confirmatory laboratory result demonstrated? <input type="checkbox"/> Culture and identification of <i>B. anthracis</i> by the Laboratory Response Network (LRN); <input type="checkbox"/> Antigen detected by immunohistochemical staining using both cell wall <u>and</u> capsule monoclonal antibodies; <input type="checkbox"/> 4-fold antibody titer increase between acute and convalescent serum or between paired convalescent sera using CDC quantitative anti-PA IgG ELISA testing; <u>or</u> <input type="checkbox"/> DNA from an affected tissue lesion or normally sterile site <u>with</u> a documented environmental exposure.	<input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> Yes
3	Was a probable laboratory result demonstrated? <input type="checkbox"/> DNA from an affected tissue lesion or normally sterile site; <input type="checkbox"/> Positive serum specimen using Quick ELISA Anthrax-PA kit; <input type="checkbox"/> Lethal Factor (LF) detected in a serum specimen by LF mass spectrometry; <u>or</u> <input type="checkbox"/> Positive RedLine Alert test.	<input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> Yes
4	Is the patient epidemiologically linked to a documented anthrax environmental exposure?	<input type="checkbox"/> No <input type="checkbox"/> Unknown <input type="checkbox"/> Yes
Confirmed: 1 & 2		Probable: 1 & 3 or 1 & 4
Suspect: 1		