Bioterrorist Threats: Sources, Recognition, & Safety

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This Presentation is Approved for 1 CRCE Credit Hour

Learning Objectives

Describe the etiology, manifestations, management, & safety precautions related to likely bioterrorist threats



History

- Book of Exodus: 5th & 6th plagues of Egypt (boils)
- > 1600s: "Black Bane" kills cattle in Europe
- > 1880: Immunization of cattle
- > 1915: First used as a bioweapon against cattle
- > 1950s-60s: U.S. develops bioweapons

History

- > 1969: U.S. ends bioweapons program
- > 1970: Anthrax vaccine is FDA approved
- > 1972: International convention outlaws biological weapons
- > 1995: Iraq admits to producing 8,500 L of anthrax weapon
- > 2001: Letter containing anthrax is mailed to NBC

Etiology

- Causative organism: bacillus anthracis
 - & Gram-positive, spore-forming rod
 - * Spore-forming \rightarrow very tough organism
 - Occurs globally, esp. in developing countries
 Primarily infects herbivores

 - * Produces lethal toxin



Routes for Transmission

- > Cutaneous: most common
- Gastrointestinal: ingestion of poorly cooked meat from infected animals
- > Inhalation of dust that contains spores woolsorter's disease

Cutaneous Anthrax

- Etiology & pathogenesis
 - Introduced via skin or mucus membrane through cut or abrasion
 - * Spores germinate & multiply
- > Manifestations: skin lesion
 - * Develops 12 36 H after infection
 - * Resembles bug or spider bite
 - * Black eschar develops

Cutaneous Anthrax



See links below to view more cutaneous anthrax

Cutaneous Anthrax

> Manifestations

- * Proximal lymphedema develops
- * Infection disseminates
- Septicemia
- Meningitis
- * Frequently fatal, if untreated

Gastrointestinal Anthrax

> Manifestations: inflammation of GI tract

- Nausea
- * Hematemesis
- * Fever
- * Acute abdomen abdominal pain
- * Severe diarrhea
- * Sepsis
- > High mortality rate

Inhalational (Pulmonary) Anthrax

- > Etiology: inhalation of spores * Special processing for deposition * 1 - 5 micron
 - Too large: upper airway deposition
 Too small: exhaled

Inhalational (Pulmonary) Anthrax

- Incubation period: generally 3-5 D, depends on germination rate
- > Manifestations early
 - * Fever, chills * Dyspnea
 - * Cough
 - * Headache
 - * Nausea & vomiting
 - * Chest pain

Inhalational (Pulmonary) Anthrax

- Manifestations: fulmination
- * Fever
- * Dyspnea
- * Stridor: mediastinal enlargement
- * Diaphoresis

Inhalational (Pulmonary) Anthrax

Manifestations: fulmination

- * Fever
- * Dyspnea
- * Stridor: mediastinal enlargement
- * Diaphoresis
- * Shock
- * Hemorrhagic meningitis: delirium
- * Hypoxemia

Inhalational (Pulmonary) Anthrax

Chest x-ray: mediastinal widening





Anthrax

- Diagnosis * Sputum exams are NOT useful * Standard blood culture: growth in 6 - 24 H
- > Pathology: hemorrhagic, necrotizing pneumonic lesion

Anthrax

- Management
 - * Antibiotics: susceptible to
 - Ciproflaxin
 - Doxycycline
 Penicillin
 - Amoxacillin
 - Chloramphenicol
 - Rifampin
 - * NOT susceptible to cephalosporins

Anthrax

Management

- Supplemental oxygen
- * Mechanical ventilation
- * Vasopressors for shock
- $\ensuremath{\diamond}$ Other supportive measures

Anthrax

Prevention

- * Direct, person-to-person spread is unlikely * Universal precautions for patient care: no special barriers
- * Antibiotics for suspected exposure (60 D)

Anthrax

Prevention: vaccination

- * Human live attenuated vaccine
 - Three injections, two weeks apart
 - Three injections at 6, 12, 18 mo.

Anthrax

- Prevention: vaccination
 - * Adverse reactions
 - Soreness, edema at injection site
 - Fever, nausea, headaches (5 35%)
 Serious events 1:50,000 doses

Anthrax

Decontamination

 Bleach
 Sandia foam: new, safe
 Formaldehyde
 Nanoemulsion

Anthrax

- Why anthrax?
- > It is tough
 - * Sunshine kills spores
 - * Heat does not kill
- \star Explosion does not kill \rightarrow can be dispersed by explosive shells



History

10,000 BC: Believed to have appeared in Africa

1350 BC: First recorded epidemic in Egypt

180 AD: Major epidemic coincides with fall of Roman empire

1500-1800 AD: Introduction of smallpox to New World decimates native population

History

1763: Biological warfare by placing smallpox scabs in blankets given to Native Americans

1600: Chinese introduced variolation, an early vaccination

1796: Jenner uses cowpox extract to vaccinate against smallpox

History

1967: World Health Organization campaign to eradicate smallpox

1972: Routine vaccination ceased

- 1980: WHO recommends cessation of vaccination
- 1980: Soviet government initiates program to produce large quantities of smallpox



Etiology

- Causative organism: variola virus
 - DNA virusVery infectious
 - * Related to
 - Cowpox
 - Monkeypox
 - Vaccinia virus
 - * Variola major: more virulent form
 - * Variola minor: less virulent

Pathogenesis

- > Transmission mode: person-to-person via droplet nuclei
- > Virus implants on oropharyngeal or respiratory mucosa
- > Only few varions are required to produce disease
- Viruses migrate & multiply in regional lymph nodes, spleen, & bone marrow
- > Incubation: about 12 D

Manifestations: Variola Major

- > Fever
- > Malaise
- > Headache, backache
- Maculopapular rash
 Oropharyngeal mucosa
 - * Face * Forearms
 - * Trunk
 - & Legs

Manifestations: Variola Major - Smallpox rash

Manifestations: Variola Major

Smallpox rash



Manifestations: Variola Major

- Rash becomes pustular
- > Large amount of virus in saliva: most infectious phase
- > Scabs develop
- > Toxemia
- > Encephalitis
- Mortality (30%): 5th or 6th day after onset of rash

Variola: Alternate Forms

- Malignant
 Abrupt onset
 Frequently fatal
- Hemorrhagic
 Rash hemorrhages
 Frequently fatal

Variola: Alternate Forms

Variola minor
 Fewer constitutional symptoms

- * Sparser rash
- > Partially immune victims: similar to variola minor

Diagnosis

- ightarrow One suspected case ightarrow international health emergency
- > Characteristic rash
 - * Centrifugal distribution
- * Same stage of development at each location
- Palmar & plantar location
- Confirmed by laboratory analysis

Diagnosis

Management

- * Strict isolation
- * Supportive care
- * Antibiotics for secondary bacterial infection
- * Antiviral agents
 - Currently, none are approved
 - Agents for HIV have potential

Prevention

Post-exposure control

- * All face-to-face contacts with victim
- Vaccinated
- Surveillance for fever, rash
- * Home care recommended for victims
- Vaccination of healthcare workers, police, transit workers, etc.

Hospital Infection Control

- Smallpox spreads easily by droplets
- > Rooms: negative pressure with HEPA
- > Vaccination of employees, patients
- > Laundry & waste- biohazards

Botulism

History

- First identified as poison from sausage (botulus = sausage)
- > 1735: First case described
- > 1897: Botulism toxin identified
- > 1930s: Japanese used as weapon
- > 1991: Iraq admits to producing 19,000 L of botulism toxin

Etiology

- Causative organism: clostridium botulinum bacterium
 - * Widespread, soilborne
 - * Obligate anaerobe * Spore-forming
 - Produces botulinum neurotoxin
 - Colorless
 - Odorless, tasteless
 - Inactivated by heat

Forms

 Food-borne: ingestion of toxin in foods that have not been canned or preserved properly

Forms

- Wound botulism, systemic spread of toxin produced by organisms inhabiting wounds, associated with
 - * Trauma
 - * Surgery
 - * Subcutaneous heroin injection
 - * Sinusitis from intranasal cocaine abuse

Forms

- > Infant botulism
 - Intestinal colonization of organisms in infants younger than 1 year
 - $\boldsymbol{\ast}$ Associated with ingestion of honey by infants

Modes of Toxin Transmission

- Food: almost all types
- > Aerosol: bioterrorism
- > Water supply: unlikely because water treatment deactivates toxin

Manifestations

- Incubation: 2 H to 8 D after exposure, ingestion
- > Diplopia
- > Blurred vision
- > Dysphonia
- > Dysphagia
- > Dysarthria
- > Loss of gag reflex

Manifestations

Paralysis

- * Loss of head control
- * Generalized weakness
- * Diaphragm & accessory ventilatory muscles
- * Recovery in weeks to months

Manifestations

Pathognomonic

- Symmetric, descending paralysis
- * Afebrile patient
- * Normal sensorium

Diagnosis

- Differential diagnosis rule out
 - * Guillain-Barre syndrome
 - * Myasthenia gravis
 - * Poliomyelitis

Laboratory tests: available only at CDC
 Blood

- * Gastric aspirates
- * Stool

Management

- > Botulism is NOT an infection
- > Evaluate airway & breathing
 \$ Loss of gag reflex → intubation
 \$ Loss of ventilatory muscles → ventilation

Management

Botulism antitoxin: STAT

 Minimizes severity
 Does not reverse existing paralysis

Prevention

> Botulism toxoid: immunization

Botulism antitoxin
 Post-exposure prevention
 Scarce

Prevention

> Decontamination: usual procedures

Infection control
 No isolation necessary
 Universal precautions



History

Naturally occurring plague: ancient

- > 425 BC: first recorded epidemic in Athens
- > 540 AD: first recorded pandemic
- > 1340 AD: pandemic from China to Europe, killing 1/3 of Europeans
- > 1400s AD: used as biological weapon by Tatars
- > 1665 AD: great plague of London

History

- > 1894: causative organism identified by Yersin, 'yersinia pestis'
- > Present day
 - * Natural epidemics recur * Organism present in rodents, worldwide, including Western U.S.
- > WWII: used by Japan as biological weapon
- Soviet Union developed large quantities of weapon-grade plague

Etiology

> Causative organism * Yersinia pestis * Gram-negative bacillus



Etiology

- Causative organism: yersinia pestis
- > Insect vector: x. cheopis flea
- > Animal reservoir: rodents
 - * Rats
 - * Mice
 - * Prairie dogs
 - * Ground squirrels

Forms

- > Bubonic: buboes are infected lymph glands
- > Pneumonic: pulmonary infection
- > Septicemic: disseminated to blood

Transmission Modes

Bites of infected fleas: bubonic form

> Aerosol

- * Pneumonic * Biological weapon

Manifestations: Bubonic

- Incubation: bubonic 2-10 D
- > Malaise
- > High fever
- > Lymph glands * Swollen & tender * May progress to buboes
- > Leukocytosis
- > Mortality 50%, if untreated

Manifestations: Bubonic



Manifestations: Bubonic

- Incubation 2-3 D
- ≻ Malaise
- > High fever, chills
- > Headache
- > Hemoptysis
- > Leukocytemia

Manifestations: Pneumonic

- Rapidly progressive bronchopneumonia
- > Dyspnea
- > Stridor
- > Hypoxemia
- > Mortality: 100% if untreated

Diagnosis

- > Index of suspicion: sudden outbreak of severe pneumonia & sepsis
- > Gram stain: sputum or blood, gram negative bipolar rod

Management

> Antibiotics: initiate STAT

- * Streptomycin: drug of choice
- * Gentamycin
- * Doxycycline
- * Tetracycline
- * Chloramphenicol
- * Trimethoprim-sulfamethoxazole
- * NOT cephalosporins

Management

> Supportive measures

* Oxygen * Mechanical ventilation

Prevention

- > Post-exposure antibiotics: seven days post-exposure * Tetracycline * Doxycycline
 - * TMP-SMT
 - * Chloramphenicol

Prevention

Isolation

- * Respiratory isolation of patient for first 48 hours * Close contacts who refuse chemoprophylaxis
- > Vaccine: limited availability
- > Decontamination: usual measures

Additional Bioterrorist Threats

- > Tularemia: extremely infectious bacterium
- > Ebola: rapidly fatal virus
- > Aflatoxin: carcinogen
- > Clostridium perfringens: gangrene
- > Ricin: slow poison

Summary & Review

Anthrax

- * Antracis bacillus
- * Cutaneous, gastrointestinal, pulmonary * Pulmonary manifestations
- * Management
- * Prevention: immunization, chemoprophylaxis
- Oniversal precautions

Summary & Review

Smallpox

- * Variola major
- * Communication: droplet nuclei
- * Primary manifestation: centrifugally distributed rash
- * Management
 - Supportive
 - Isolation
 - Home care

Summary & Review

Smallpox

- * Prevention: vaccination
- * Precautions
 - Strict isolation • Biohazardous waste

Summary & Review

> Botulism

- * Clostridium botulinum: produces neurotoxin
- * Sources
- Manifestation: descending paralysis
 Management
 - May require intubation, ventilation
- Antitoxin
- Prevention: immunization (botulinum toxoid)
- * Universal precautions

Summary & Review

> Plague

- * Yersinia pestis: gram negative rod
- * Insect vector (flea)
- * Infected rodents
- $\boldsymbol{\ast}$ Types: bubonic, pneumonic, septicemic
- Manifestations: buboes, pneumonia
 Management: antibiotics, etc.
- Prevention: immunization, chemoprophylaxis
- Precautions: isolation first 48 hours

References

- http://www.newadvent.org/cathen/12143a.htm
- http://www.bact.wisc.edu/Bact330/lectureanthrax
- > http://www.cdc.gov/ncidod/dbmd/diseaseinfo/anthrax_g.htm
- http://www.cdc.gov/mmwr/preview/mmwrhtml/mm5041a2.htm
- http://jama.ama-assn.org/issues/v281n18/ffull/jst80027.html
- http://www.sandia.gov/media/cbwfoam.htm
- http://38.232.17.254/journals/annals/15oct97/smallpox.htm
- http://jama.ama-assn.org/issues/v281n22/ffull/jst90000.html

References

- http://www.acponline.org/journals/annals/01oct96/botuedit.htm
- http://jama.ama-assn.org/issues/v285n8/ffull/jst00017.html
- > http://www.cdc.gov/ncidod/dvbid/plague/index.htm