Vaccine-Preventable **Diseases**

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Disclosure Statement

· I have no conflicts of interest to disclose relevant to today's presentation

Vaccine-Preventable **Diseases**

- Anthrax
- Cholera
- Diphtheria
- **Hepatitis A**
- **Hepatitis B**
- H. influenzae type B
- HPV
- Seasonal influenza

- **Japanese**
- encephalitis
- Meningococus

Shingles

Smallpox

Tetanus

Varicella

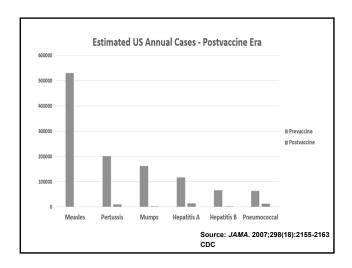
· Yellow fever

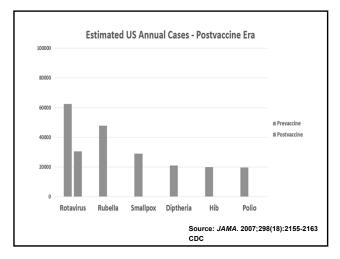
Typhoid fever

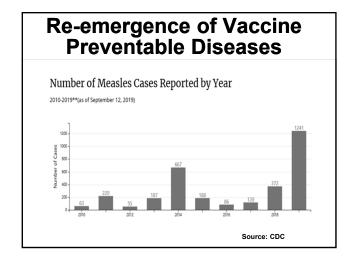
- Mumps
- Pertussis
- Pneumococcus
- Polio
- Rabies
- Rotavirus Rubella

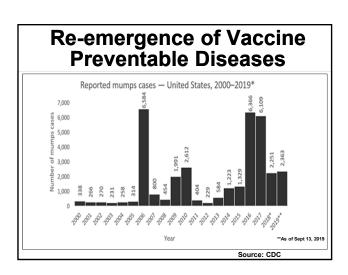
Life Before Vaccines

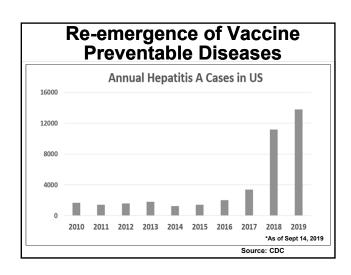
- · Diphtheria and smallpox outbreaks
- Summer infantile paralysis epidemics
- · Near universal infection with measles and pertussis during childhood
- · Congential rubella syndrome
- · Invasive Haemophilus influenzae type B disease
- · The list goes on...

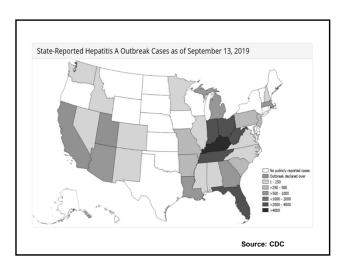


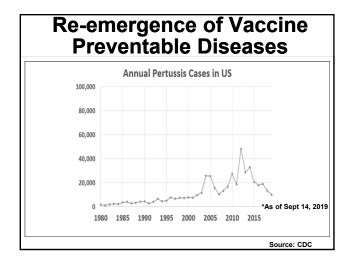












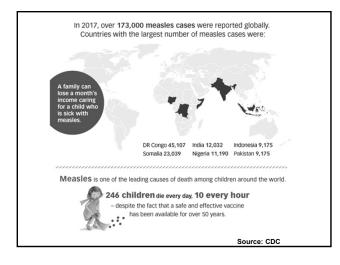
Re-emergence of Vaccine Preventable Diseases

- Contributing Factors:
 - · Decreased vaccination rates
 - · Endemic transmission
 - · Increased international travel
 - · Waning vaccine-mediated immunity

Outline

- · Notable VPDs in the clinic setting:
 - Measles
 - Mumps
 - · Hepatitis A
 - Influenza addressed in separate webcast
- Common vaccine questions from patients

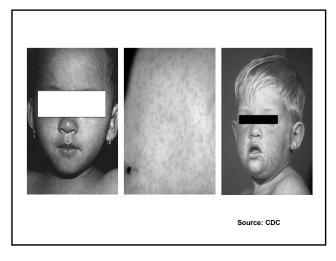


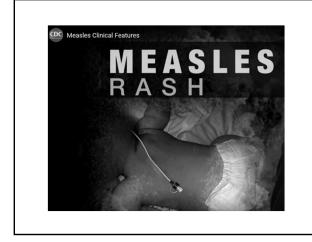


Measles Clinical Features

- Incubation Period: 8-12 days (range: 7-21 days)
- Symptoms arise as two distinct phases:
 - Prodrome
 - 2-4 days prior to rash onset
 - Fever & "the 3 C's"
 - Koplik spots
 - Rash
 - Cephalocaudal progression
 - Confluence
 - Fading with desquamation
- Infectious 4 days prior to 4 days after rash onset







Measles Complications Measles Can Be Serious · Acute Otitis Media Diarrhea Febrile Seizures Pneumonia Encephalitis 1 out of every 1,000 1 to 3 out of 1,000 Post-infectious people with measles will develop brain people with measles will die, even with people who get measles encephalomyelitis swelling due to infection the best care. (encephalitis), which may Subacute sclerosing lead to brain damage. panencephalitis Source: CDC

Measles Diagnosis

- RT-PCR
 - · Nasopharyngeal or throat swab specimen
 - · Highest sensitivity during first 3 days of rash
- Serology
 - IgM
 - Acute specimens may have false negative
 - · False positives may occur with other viral infections
 - IgG
 - · Usually positive by 1-2 weeks after rash

Measles Treatment and Prophylaxis

- Treatment
 - Supportive
 - · Vitamin A for hospitalized children
- **Post-Exposure Prophylaxis**
 - MMR Vaccine within 72 hours
 - · Immunoglobulin within 6 days
 - · Intramuscular immune globulin -Infants
 - Intravenous immune globulin
 - -Immunocompromised children and adults
 - -Pregnant women without evidence of immunity

Measles Prevention

- Prevention
 - · 2 dose MMR series in childhood
 - Other indications:
 - · Students at post-high school educational institutions
 - · Adults born during or after 1957
 - · Prior to international travel
 - · Healthcare personnel



Mumps

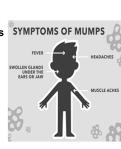
- · Highly contagious Spread via respiratory droplets
- · Endemic transmission ongoing in US





Mumps Clinical Features

- Incubation period 16-18 days (range: 12-25 days)
- Non-specific prodromal symptoms
- Tender unilateral or bilateral parotitis
 - Symptoms peak in 1-3 days
 - Resolve over 1 week
- · Clinical presentation may vary
 - Asymptomatic
 - Non-specific respiratory symptoms
- Infectious 2 days before to 5 days after parotitis onset







Source: CDC

Ear Protrusion &
Obscuring of the angle of the jaw common



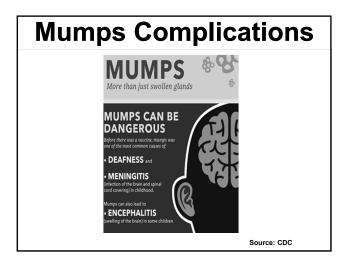
Source: CDC

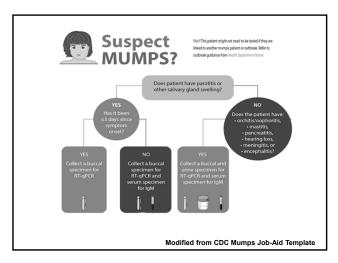
Mumps Complications

- · Occur less commonly in vaccinated patients
- · Complications are more common in adults
- Orchitis
- Oophoritis
- Mastitis



Source: CDC





Mumps Testing Considerations

- · Previously vaccinated patients:
 - Obtain PCR specimens within 1-3 days after onset
 - May have transient or undetectable IgM
 - IgG during acute phase usually very high

Mumps Treatment and Prophylaxis

- Treatment
 - Supportive
- Post-exposure prophylaxis
 - None

Mumps Prevention

- Prevention
 - 2 dose MMR series in childhood
 - Other indications:
 - Students at post-high school educational institutions
 - Adults born during or after 1957
 - Prior to international travel
 - · Healthcare personnel
 - 3rd dose for high risk groups during outbreak



Source: CDC

Hepatitis A

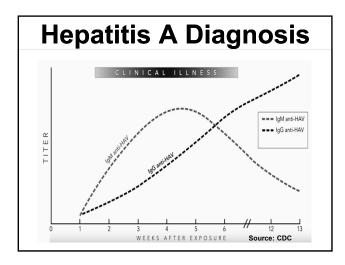
- · Transmission routes:
 - Fecal-oral
 - · Contaminated food or water
- · Risk factors:
 - · Contact with infected person
 - International travel
 - Men who have sex with men
 - · Users of injection and non-injection drugs
 - · Persons with clotting factor disorders
 - Working with NHP

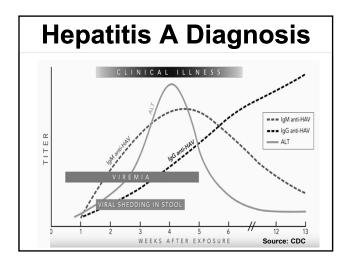
Hepatitis A Clinical Features

- Incubation period: 28 days (range: 15-50 days)
- Most children < 6 years asymptomatic
- · Older children and adults:
 - Fever
 - Fatique
 - Abdominal pain
 - Nausea and vomiting
 - Diarrhea
 - Jaundice

Hepatitis A Clinical Features

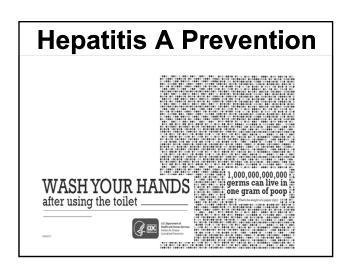
- Symptoms resolve in < 2 months
- · Prolonged or relapsing disease may occur
- · Does not cause chronic infection
- Infectious from 2 weeks before to 1 week after jaundice onset





Hepatitis A Treatment and Prophylaxis

- Treatment
 - Supportive
- · Post-Exposure Prophylaxis within 14 days of exposure
 - · Hepatitis A vaccine
 - Healthy persons aged ≥ 12 months
 - · Immune globulin & hepatitis A vaccine
 - Immunocompromised persons aged ≥ 12 months
 - · Chronic liver disease
 - Healthy persons aged > 40
 - · Immune globulin alone
 - Infants < 12 months

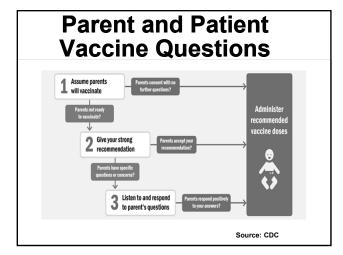


- All children at age 1 year - Travelers to countries where hepatitis A is common - Family and caregivers of adoptees from countries where hepatitis A is common - Men who have sexual encounters with other men - People who use or inject drugs - People with chronic or long-term liver disease, including hepatitis B or hepatitis C - People with clotting factor disorders - People with direct contact with others who have hepatitis A - People experiencing homelessness

Source: CDC

Hepatitis A Prevention

- · Other patient populations to vaccinate:
 - · Persons at increased risk of complications
 - Congenital or acquired immunodeficiency
 - HIV infection
 - Hemodialysis
 - · Transplant recipients
 - · latrogenic immune suppression
 - · Occupational risks
 - Nonhuman primates
 - · Working with HAV in research laboratory



Common Questions and Concerns

- Too many vaccines
 - Contrast vaccine antigens with every day exposures
- Vaccines make me sick
 - · Educate on immune response
- Vaccines contain aluminum or other metals
 - No known safety risks with amount in vaccines
 - Ingested in food and water daily
- Delayed vaccine schedule
 - · No data that delayed schedule is more safe
 - · Any time delay places at risk

Common Questions and Concerns

- · Delay for mild illness
 - · Mild febrile illnesses are not contraindications
- · VPDs don't exist anymore
 - Educate on international and US outbreaks
 - · "One plane ride away"



Source: CDC

Summary

- · One of greatest public health achievements
- VPDs still exist in US and internationally
- Maintain high level of clinical suspicion
- Encourage families and patients to vaccinate