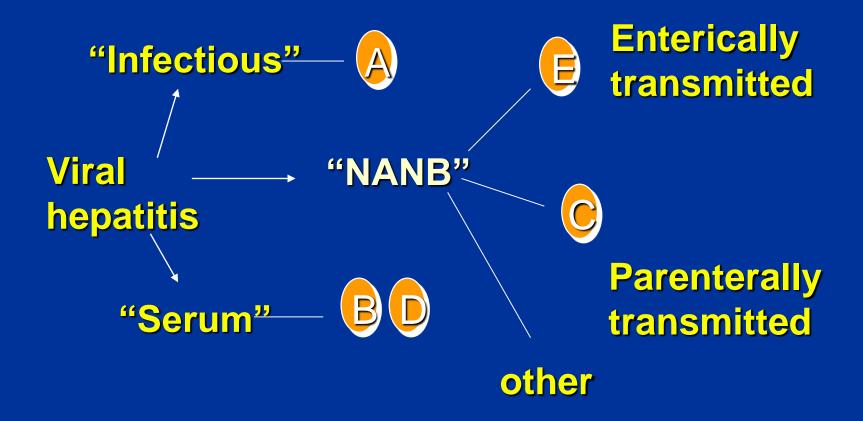
VIRAL HEPATITIS A

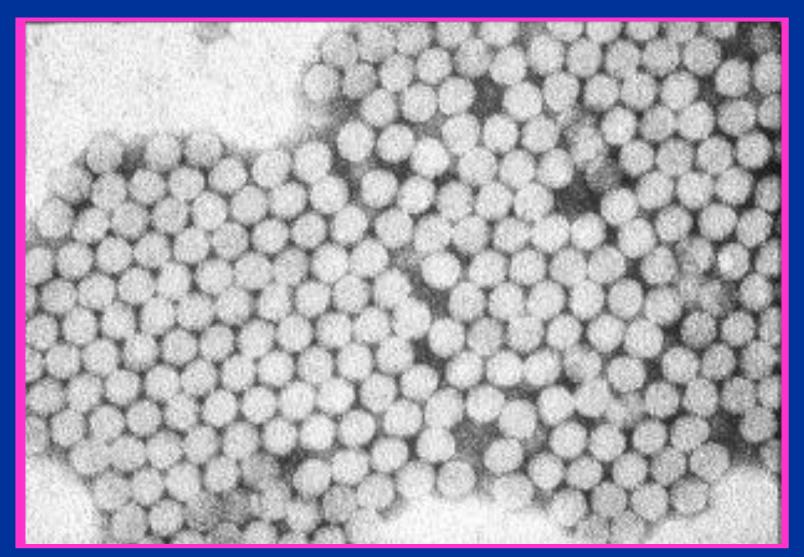


VIRAL HEPATITIS HISTORICAL PERSPECTIVE





HEPATITIS A VIRUS





HEPATITIS A VIRUS

- RNA Picornavirus
 - Single serotype worldwide
 - Acute disease and asymptomatic infection
- No chronic infection
 - Protective antibodies develop in response to infection - confers lifelong immunity



HEPATITIS A - CLINICAL FEATURES

•Jaundice by <6 yrs <10%

age group: 6-14 yrs 40%-50%

>14 yrs 70%-80%

•Rare complications: Fulminant hepatitis

Cholestatic hepatitis

Relapsing hepatitis

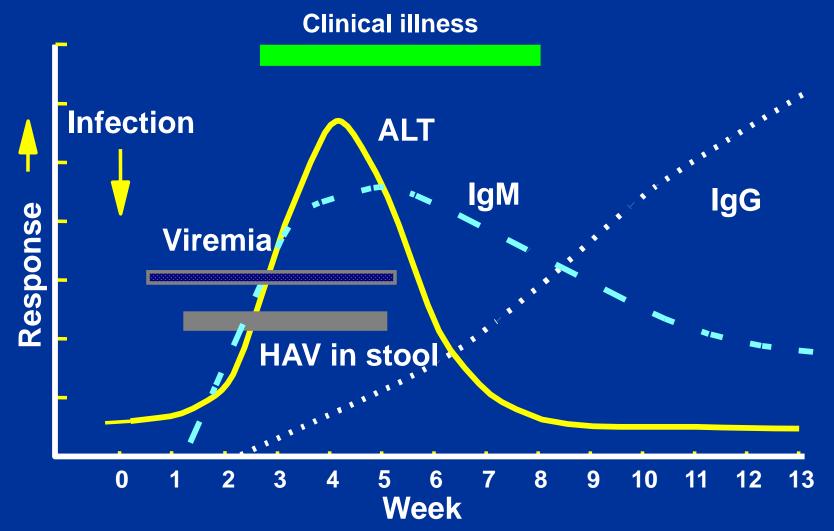
Incubation period: Average 30 days

Range 15-50 days

Chronic sequelae: None

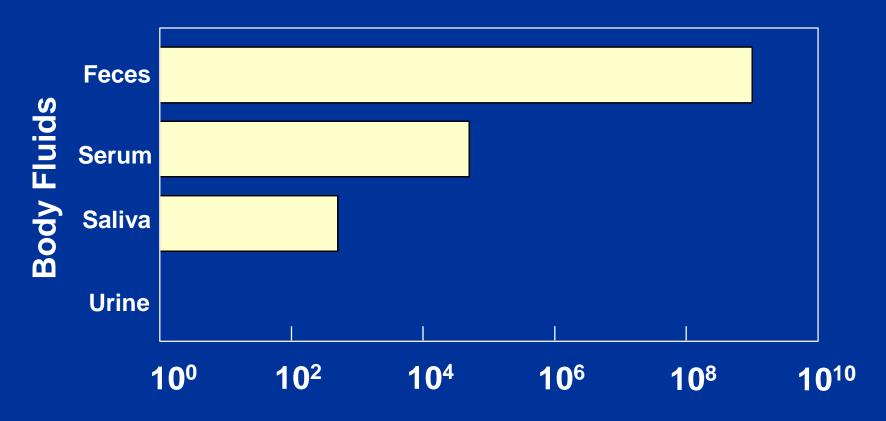


EVENTS IN HEPATITIS A VIRUS INFECTION





CONCENTRATION OF HEPATITIS A VIRUS IN VARIOUS BODY FLUIDS



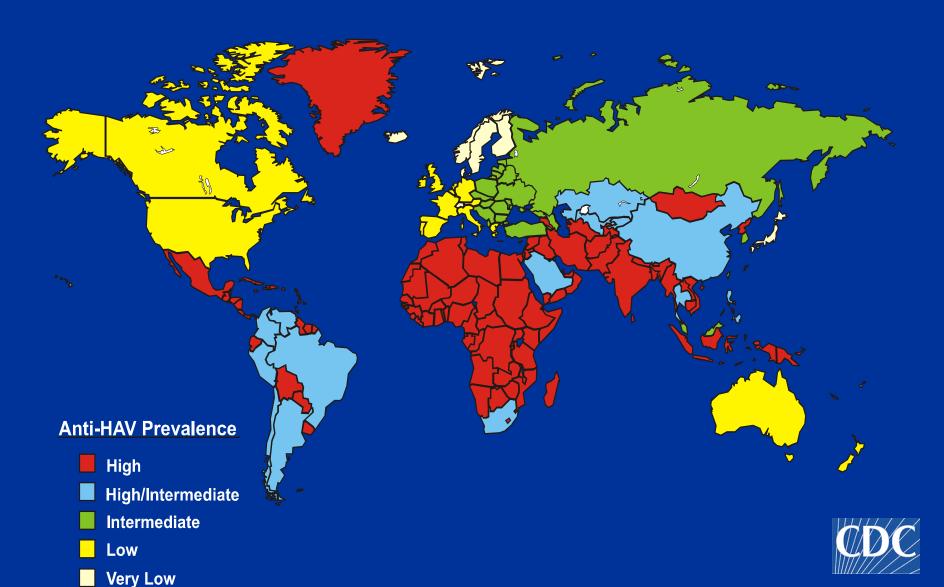
Infectious Doses per mL

Source: Viral Hepatitis and Liver Disease 1984;9-22

J Infect Dis 1989;160:887-890



GEOGRAPHIC DISTRIBUTION OF HEPATITIS A VIRUS INFECTION



ACUTE HEPATITIS A CASE DEFINITION FOR SURVEILLANCE

Clinical criteria

An acute illness with:

- discrete onset of symptoms (e.g. fatigue, abdominal pain, loss of appetite, intermittent nausea, vomiting), and
- jaundice or elevated serum aminotransferase levels

Laboratory criteria

IgM antibody to hepatitis A virus (anti-HAV) positive

Case Classification

 Confirmed. A case that meets the clinical case definition and is laboratory confirmed or a case that meets the clinical case definition and occurs in a person who has an epidemiologic link with a person who has laboratory-confirmed hepatitis A (i.e., household or sexual contact with an infected person during the 15-50 days before the onset of symptoms).



HEPATITIS A VIRUS TRANSMISSION

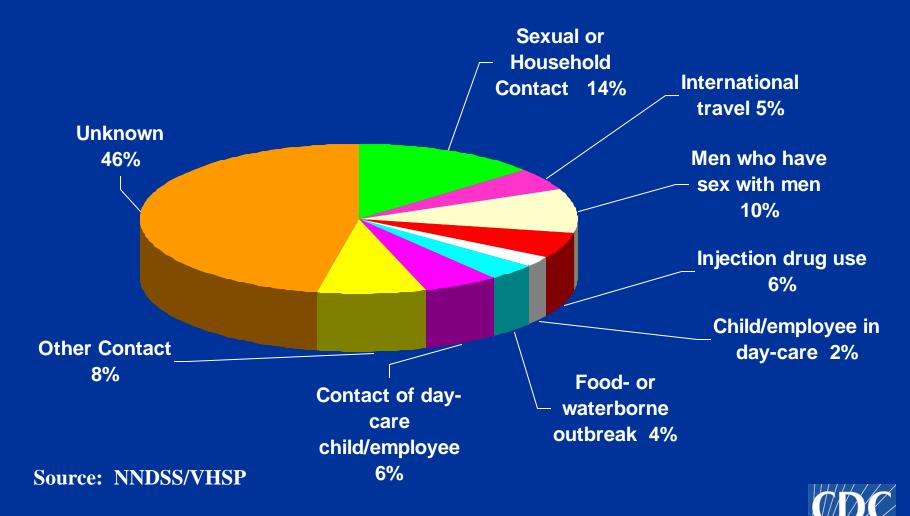
- Close personal contact

 (e.g., household contact, sex
 contact, child day-care centers)
- Contaminated food, water (e.g., infected food handlers)
- Blood exposure (rare)

 (e.g., injection drug use, rarely by transfusion)



RISK FACTORS ASSOCIATED WITH REPORTED HEPATITIS A, 1990-2000, UNITED STATES



PREVENTING HEPATITIS A

- Hygiene (e.g., hand washing)
- Sanitation (e.g., clean water sources)
- Hepatitis A vaccine (pre-exposure)
- Immune globulin (pre- and postexposure)



PREPARATION OF INACTIVATED HEPATITIS A VACCINES

 Cell culture adapted virus grown in human fibroblasts

- Purified product inactivated with formalin
- Adsorbed to aluminum hydroxide adjuvant



HEPATITIS A VACCINES

- Highly immunogenic
 - 97%-100% of children, adolescents, and adults have protective levels of antibody within 1 month of receiving first dose; essentially 100% have protective levels after second dose
- Highly efficacious
 - In published studies, 94%-100% of children protected against clinical hepatitis A after equivalent of one dose



HEPATITIS A VACCINES

Recommended Dosages of Hepatitis A Vaccines

Schedule Vaccine	Age		Volume	2-Dose
	(yrs)	<u>Dose</u>	<u>(mL)</u>	(<u>mos)</u>
HAVRIX ® #	1-18	720 (EL.U.*)	0.5	0, 6-12
	>18	1,440	1.0	0, 6-12
VAQTA ®##	1-18	25 (U**)	0.5	0, 6-18
	>18	50	1.0	0, 6-18

has no preservative



^{*} EL.U. – Enzyme-linked immunosorbent assay (ELISA) units

^{**} Units

[#] has 2-phenoxyethanol as a preservative

SAFETY OF HEPATITIS A VACCINE

- Most common side effects
 - Soreness/tenderness at injection site -50%
 - Headache 15%
 - Malaise 7%
- No severe adverse reactions attributed to vaccine
- Safety in pregnancy not determined risk likely low
- Contraindications severe adverse reaction to previous dose or allergy to a vaccine component
- No special precautions for immunocompromised persons



DURATION OF PROTECTION AFTER HEPATITIS A VACCINATION

- Persistence of antibody
 - At least 5-8 years among adults and children
- Efficacy
 - No cases in vaccinated children at 5-6 years of follow-up
- Mathematical models of antibody decline suggest protective antibody levels persist for at least 20 years
- Other mechanisms, such as cellular memory, may contribute



FACTORS ASSOCIATED WITH DECREASED IMMUNOGENICITY TO HEPATITIS A VACCINE

- Decreased antibody concentration:
 - Concurrent administration of IG
 - Presence of passively-transferred maternal antibody
 - Age
 - Chronic liver disease
- Decreased seroconversion rate:
 - HIV infection
 - May be related to degree of immunosuppression
 - Liver transplantation



USE OF HEPATITIS A VACCINE FOR INFANTS

- Safe and immunogenic for infants without maternal antibody
- Presence of passively-acquired maternal antibody blunts immune response
 - all respond, but with lower final antibody concentrations
- Age by which maternal antibody disappears is unclear
 - still present in some infants at one year
 - probably gone in vast majority by 15 months



COMBINED HEPATITIS A HEPATITIS B VACCINE

- Approved by the FDA in United States for persons ≥18 years old
- Contains 720 EL.U. hepatitis A antigen and 20 µg. HBsAg
- Vaccination schedule: 0,1,6 months
- Immunogenicity similar to single-antigen vaccines given separately
- Can be used in persons ≥ 18 years old who need vaccination against both hepatitis A and B
- Formulation for children available in many other countries



PRE-VACCINATION TESTING

Considerations:

- cost of vaccine
- cost of serologic testing (including visit)
- prevalence of infection
- impact on compliance with vaccination
- Likely to be cost-effective for:
 - persons born in high endemic areas
 - ◆ Older U.S. born adults
 - Older adolescents and young adults in certain groups (e.g., Native Americans, Alaska Natives, Hispanics, IDUs)



POST-VACCINATION TESTING

Not recommended:

- High response rate among vaccinees
- Commercially available assay not sensitive enough to detect lower (protective) levels of vaccine-induced antibody



HEPATITIS A PREVENTION IMMUNE GLOBULIN

- Pre-exposure
 - travelers to intermediate and high HAV-endemic regions
- Post-exposure (within 14 days)

Routine

household and other intimate contacts

Selected situations

- institutions (e.g., day-care centers)
- common source exposure (e.g., food prepared by infected food handler)



HEPATITIS A VACCINATION RECOMMENDATIONS: GUIDING PRINCIPLES

- Need comprehensive strategy to reduce overall rates
 - Routine vaccination of children likely to be most effective
- Need creative approaches
 - Formulation not available that would allow integration into infant schedule



ACIP RECOMMENDATIONS PERSONS AT INCREASED RISK OF INFECTION, 1996

- Men who have sex with men
- Illegal drug users
- International travelers
- Persons who have clotting factor disorders
- Persons with chronic liver disease

