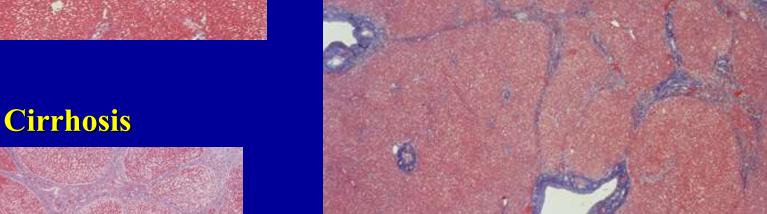
Cirrhosis and Portal hypertension

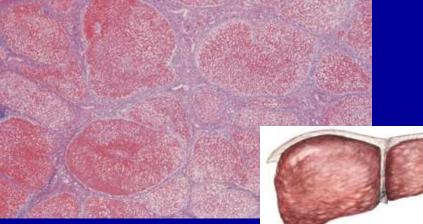


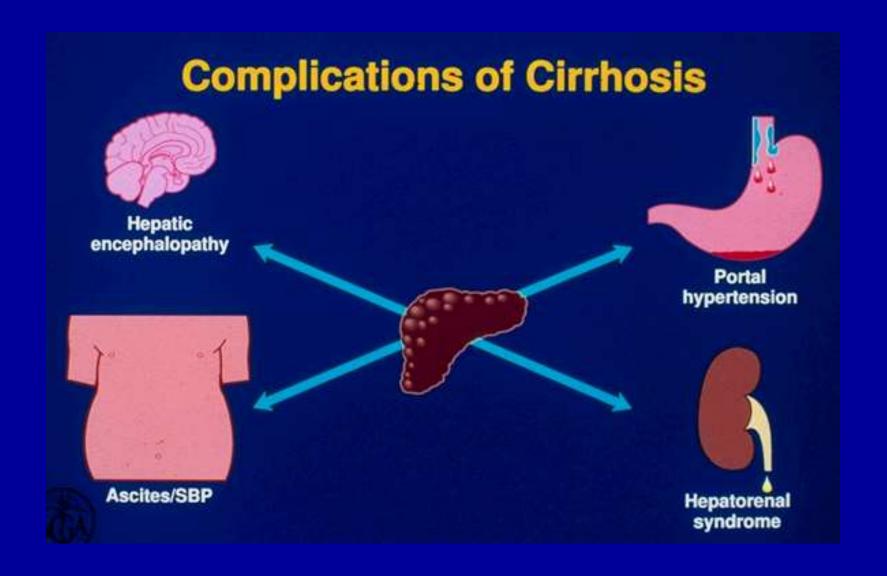




Liver Fibrosis











Jaundice

Accumulation of bilirubin in the blood stream causing yellowish discoloration of plasma and heavily perfused tissues









Spider Angiomas

Small, centrally raised bumps (papules) caused by a dilated arteriole (small artery). A network of dilated capillaries (tiny blood vessels) radiate from the arteriole. Pressing on the lesion causes the redness to disappear briefly, and there is a rapid return of redness once the pressure is lifted.



Finger Clubbing

a condition where there is enlargement of the terminal end of the digit over the distal phalanx.

It is usually symmetrical and affects the fingers







Gynecomastia

Breast development in men





Dupuytren's Contractures

Joint contractures





Distended and engorged umbilical veins which are seen radiating from the umbilicus across the abdomen to join systemic veins.















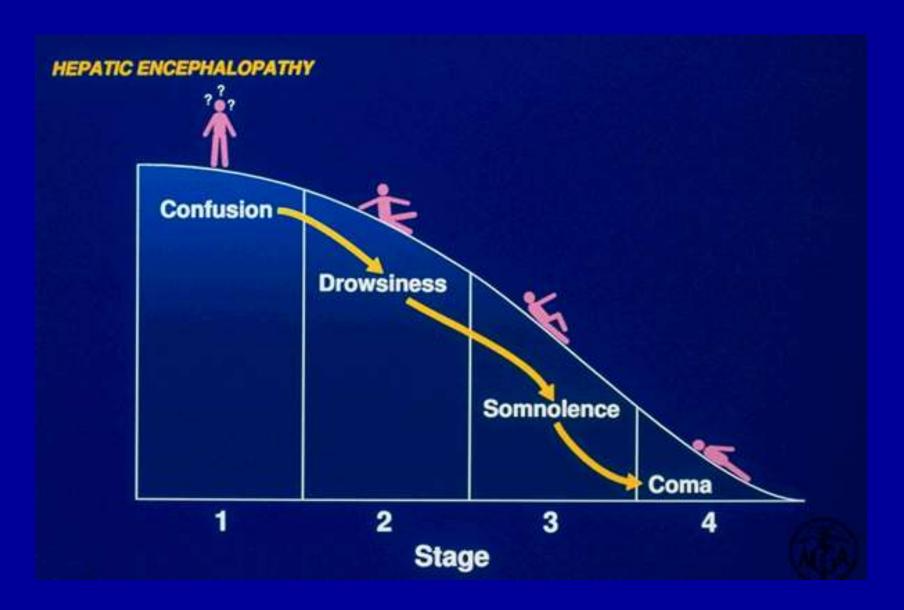
















Astraxia

Flapping tremors, quick arrythmic movement in back ground tonic muscle contracion



Hepatitis A-E Viruses

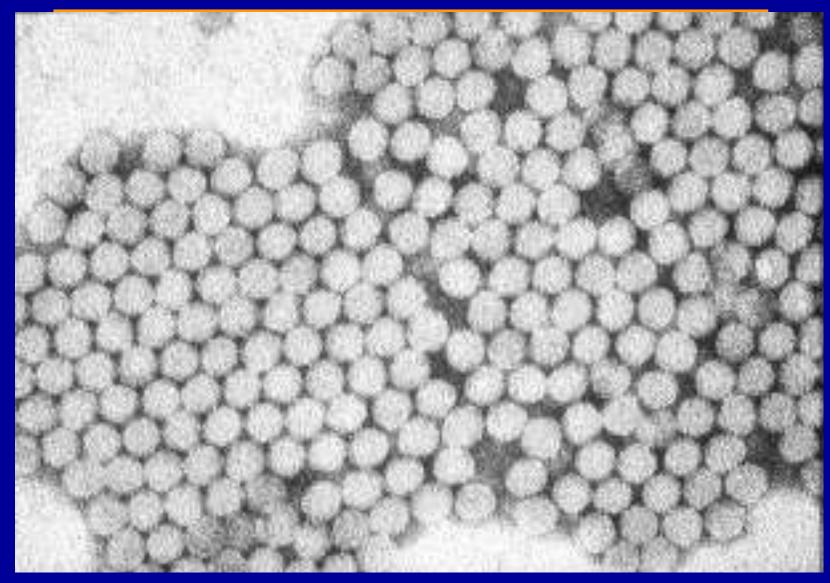
An Overview



Type of Hepatitis

	A	В	С	D	E
Source of virus	feces	blood/ blood-derived body fluids	blood/ blood-derived body fluids	blood/ blood-derived body fluids	feces
Route of transmission	fecal-oral	percutaneous permucosal	percutaneous permucosal	percutaneous permucosal	fecal-oral
Chronic infection	no	yes	yes	yes	no
Prevention	pre/post- exposure immunization	pre/post- exposure immunization	blood donor screening; risk behavior modification	pre/post- exposure immunization; risk behavior modification	ensure safe drinking water

Hepatitis A Virus





Hepatitis A - Clinical Features

- Incubation period:
- Jaundice by age group:
- Complications:

Chronic sequelae:

Average 30 days Range 15-50 days

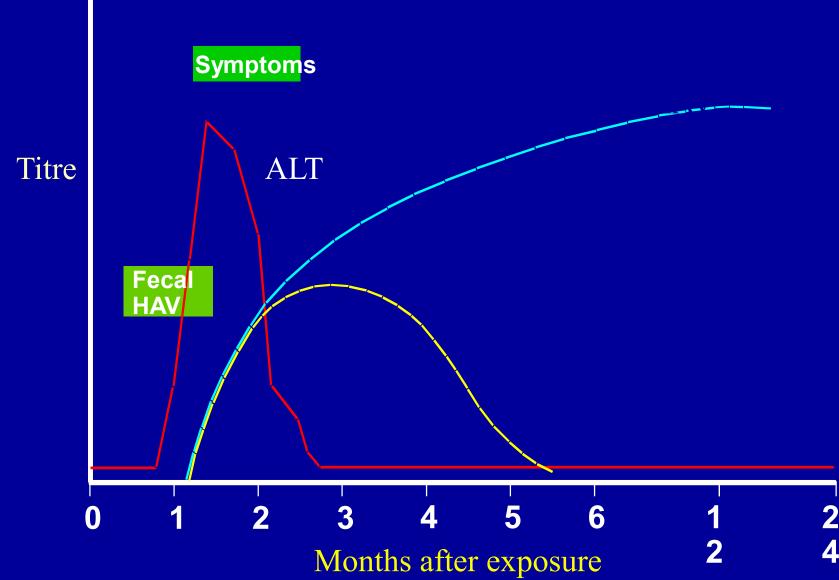
<6 yrs, <10% 6-14 yrs, 40%-50% >14 yrs, 70%-80%

Fulminant hepatitis
Cholestatic hepatitis
Relapsing hepatitis

None



Hepatitis A Infection Typical Serological Course





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, raw shellfish)
- Blood exposure (rare)
 (e.g., injecting drug use, transfusion)



Laboratory Diagnosis

- Acute infection is diagnosed by the detection of HAV-IgM in serum by EIA.
- Past Infection i.e. immunity is determined by the detection of HAV-IgG by EIA.



Hepatitis B Virus





Hepatitis B - Clinical Features

- Incubation period:
- Clinical illness (jaundice):
- Acute case-fatality rate:
- Chronic infection:
- Premature mortality from chronic liver disease:

Average 60-90 days Range 45-180 days

<5 yrs, <10% 5 yrs, 30%-50%

0.5%-1%

<5 yrs, 30%-90% 5 yrs, 2%-10%

15%-25%

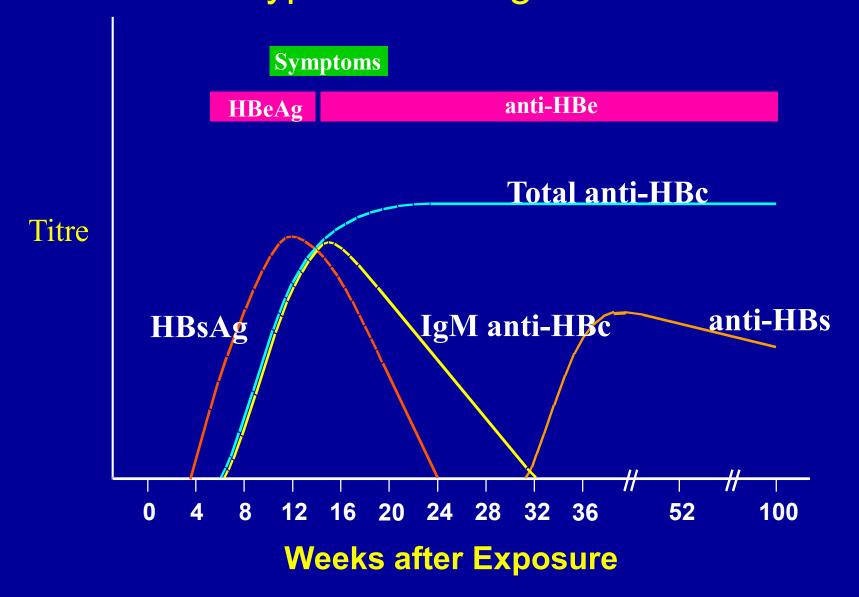


Spectrum of Chronic Hepatitis B Diseases

- 1. Chronic Persistent Hepatitis asymptomatic
- 2. Chronic Active Hepatitis symptomatic exacerbations of hepatitis
- 3. Cirrhosis of Liver
- 4. Hepatocellular Carcinoma



Acute Hepatitis B Virus Infection with RecoveryTypical Serologic Course





Outcome of Hepatitis B Virus Infection by Age at Infection 1007 100 80 80 Chronic Infection (%) Symptomatic **60** -60 **Chronic Infection** Infection (% 40 -40 **Chronic Infection (%)** 20 -20 **Symptomatic Infection** 1-6 months 1-4 years **Older Children Birth** 7-12 months and Adults Age at Infection

Concentration of Hepatitis B Virus in Various Body Fluids

High	Moderate	Low/Not Detectable
blood	semen	urine
serum	vaginal fluid	feces
wound exudates	saliva	sweat
		tears
		breastmilk



Hepatitis B Virus Modes of Transmission

- Sexual sex workers and homosexuals are particular at risk.
- Parenteral IVDA, Health Workers are at increased risk.
- Perinatal Mothers who are HBeAg positive are much more likely to transmit to their offspring than those who are not. Perinatal transmission is the main means of transmission in high prevalence populations.



Diagnosis

- A battery of serological tests are used for the diagnosis of acute and chronic hepatitis B infection.
- HBsAg used as a general marker of infection.
- HBsAb used to document recovery and/or immunity to HBV infection.
- anti-HBc IgM marker of acute infection.
- anti-HBcIgG past or chronic infection.
- HBeAg indicates active replication of virus and therefore infectiveness.
- Anti-Hbe virus no longer replicating. However, the patient can still be positive for HBsAg which is made by integrated HBV.
- HBV-DNA indicates active replication of virus, more accurate than HBeAg especially in cases of escape mutants. Used mainly for monitoring response to therapy.

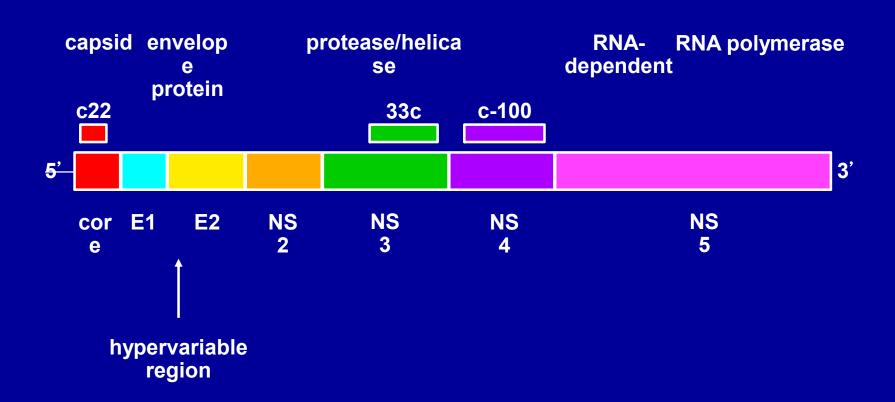


Prevention

- Vaccination highly effective recombinant vaccines are now available. Vaccine can be given to those who are at increased risk of HBV infection such as health care workers. It is also given routinely to neonates as universal vaccination in many countries.
- Hepatitis B Immunoglobulin HBIG may be used to protect persons who are exposed to hepatitis B. It is particular efficacious within 48 hours of the incident. It may also be given to neonates who are at increased risk of contracting hepatitis B i.e. whose mothers are HBsAg and HBeAg positive.
- Other measures screening of blood donors, blood and body fluid precautions.



Hepatitis C Virus





Hepatitis C - Clinical Features

Incubation period:

Average 6-7 wks

Range 2-26 wks

Clinical illness (jaundice):

30-40% (20-30%)

Chronic hepatitis:

70%

Persistent infection:

85-100%

Immunity:

No protective

antibody

response identified



Chronic Hepatitis C Infection

- The spectrum of chronic hepatitis C infection is essentially the same as chronic hepatitis B infection.
- All the manifestations of chronic hepatitis B infection may be seen, albeit with a lower frequency i.e. chronic persistent hepatitis, chronic active hepatitis, cirrhosis, and hepatocellular carcinoma.



Risk Factors Associated with Transmission of HCV

- Transfusion or transplant from infected donor
- Injecting drug use
- Hemodialysis (yrs on treatment)
- Accidental injuries with needles/sharps
- Sexual/household exposure to anti-HCV-positive contact
- Multiple sex partners
- Birth to HCV-infected mother



Laboratory Diagnosis

- HCV antibody generally used to diagnose hepatitis C infection. Not useful in the acute phase as it takes at least 4 weeks after infection before antibody appears.
- HCV-RNA various techniques are available e.g. PCR and branched DNA. May be used to diagnose HCV infection in the acute phase. However, its main use is in monitoring the response to antiviral therapy.
- HCV-antigen an EIA for HCV antigen is available. It is used in the same capacity as HCV-RNA tests but is much easier to carry out.

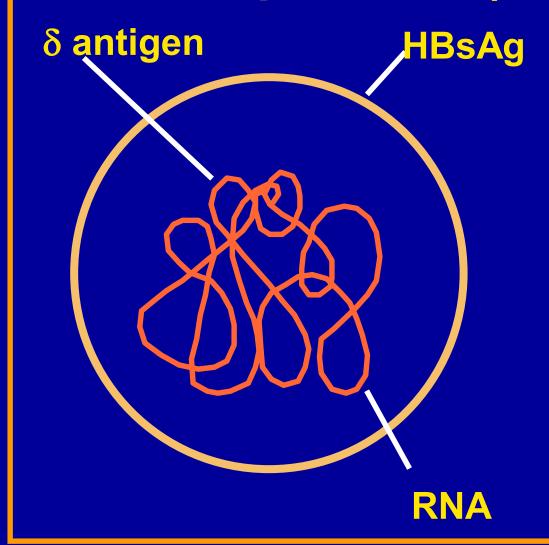


Prevention of Hepatitis C

- Screening of blood, organ, tissue donors
- High-risk behavior modification
- Blood and body fluid precautions



Hepatitis D (Delta) Virus







Hepatitis D - Clinical Features

Coinfection

- severe acute disease.
- -low risk of chronic infection.

Superinfection

- usually develop chronic HDV infection.
- high risk of severe chronic liver disease.
- may present as an acute hepatitis.



Hepatitis D Virus Modes of Transmission

- Percutanous exposures
 - injecting drug use
- Permucosal exposures
 - sex contact



Hepatitis E Virus





Hepatitis E - Clinical Features

• Incubation period:

Average 40 days Range 15-60 days

Case-fatality rate:

Overall, 1%-3%

Pregnant women,

15%-25%

Illness severity:

Increased with age

Chronic sequelae:

None identified

