

Dr. Salwa Tayel & Prof. Ashry Gad Mohamed Depart. Family & Community Medicine College of Medicine November, 2017

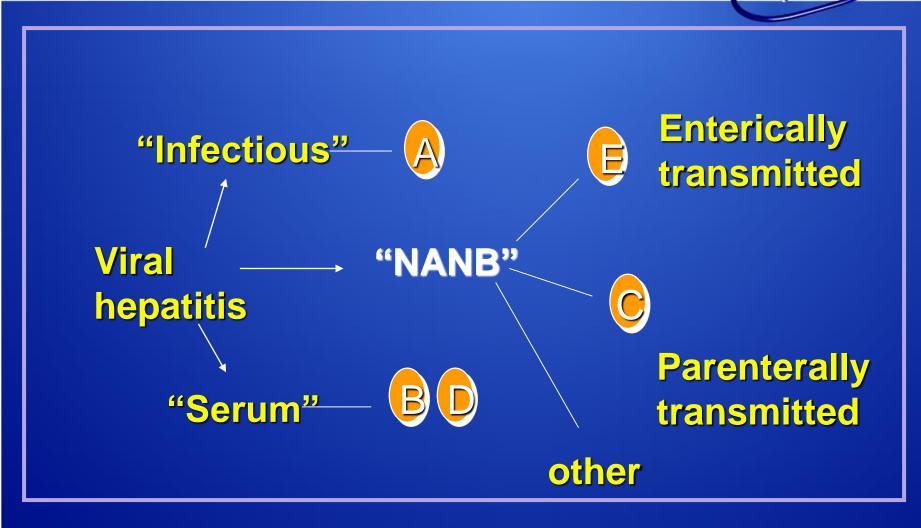
November, 2017

Objectives

At the end of the lecture students should be able:

- Understand Classification of viral hepatitis.
- Recognize the magnitude of viral hepatitis infections.
- Understand modes of transmission of different serotypes.
- Understand measures of prevention and control of different serotypes of viral hepatitis.

Viral Hepatitis – Classification &Historical Perspective



Hepatitis A

Clinical presentation:

- Abrupt onset.
- Fever
- Malaise
- Anorexia
- Abdominal discomfort
- Jaundice



Hepatitis A



- More than 90% are asymptomatic
- Seroprevalence increases with age.
- At age 15, 95% are seropositive.
- Case fatality rate (CFR)= 0.3%.
- If age > 40 years CFR=2%.
- Studies in KSA:
 - 1997 25%
 - 1999 25% Taif 10-82% Jazan (1-12 years)

Chain of infection



- Agent: RNA virus
- Reservior : Human (Clinical & subclinical cases)
- Incubation period: 15-45 days (median one month).

Chain of infection

- 6
- Period of communicability : Last two weeks of I.P. + one week of illness.

Modes of transmission:
 Fecal-oral route.
 Common source outbreaks.
 Blood transfusion (rare).

Prevention and Control



Good sanitation & personal hygiene. "Careful hand washing"
Day- Care centers Hand washing after every diaper change and before eating.

Shellfish

heat 85-90C4 minutes.steam90 seconds.

Prevention and Control



- Inactivated hepatitis A vaccine
- Schedule 2 doses after 6 months interval.
- Intramuscularly.
- Protection after one month.
- Lasting immunity at least 10 years.

Hepatitis A patient:

• Enteric precaution for the Period of communicability

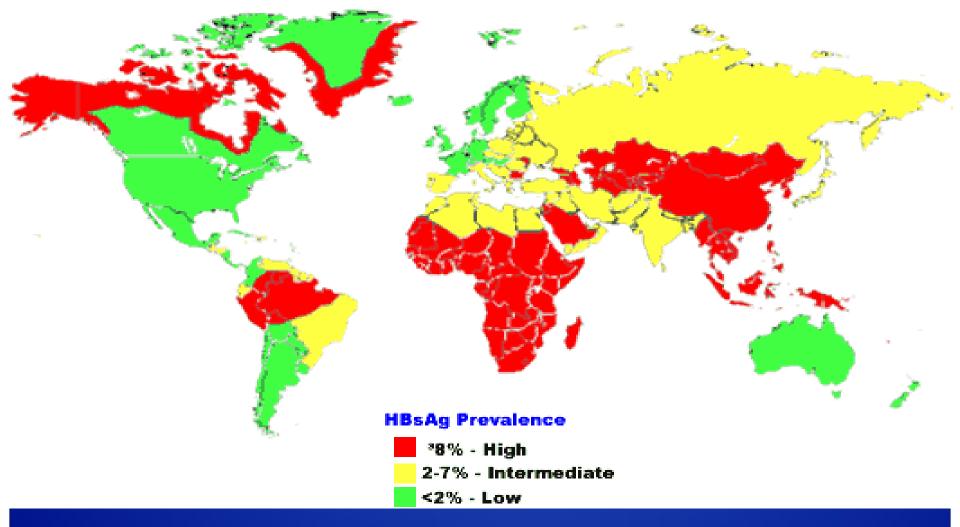
Hepatitis B

Clinical presentation:

- Insidious onset.
- Anorexia.
- Abdominal discomfort.
- Nausea.
- Vomiting.
- Arthralgia.
- Jaundice.



Geographic Distribution of Chronic HBV Infection

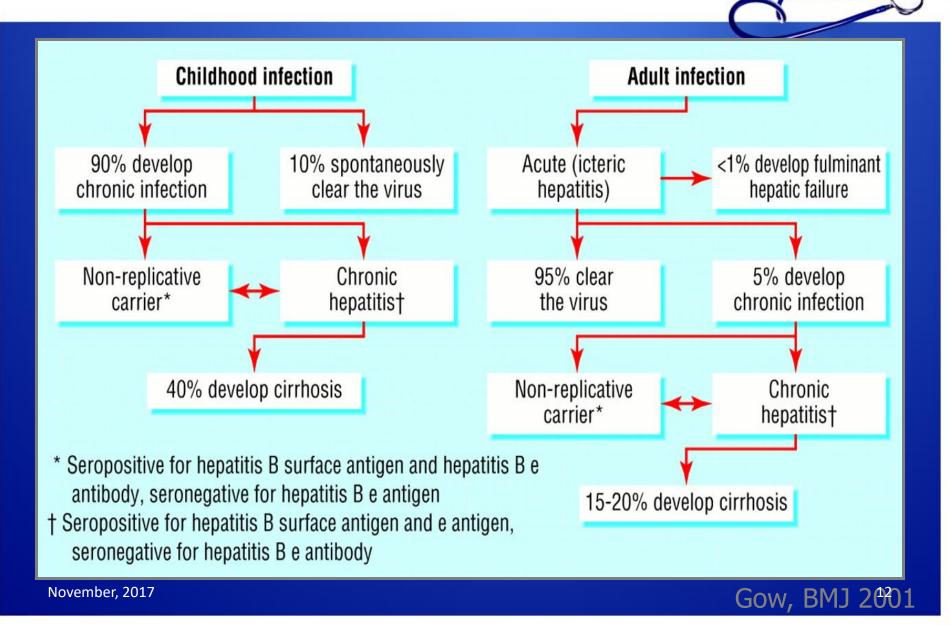


More than 500,000 death/year

2 billion people infected

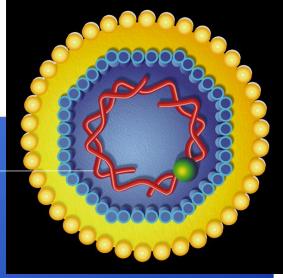
360 million CHB

Natural History



Hepatitis B Virus

HBsAg



HBcAg

HBeAg

Double-Stranded

DNA

The presence of HBsAg indicates active infection or chronic carrier. Antibody to HBsAg, from either disease or vaccine, indicates immunity.

November, 2017

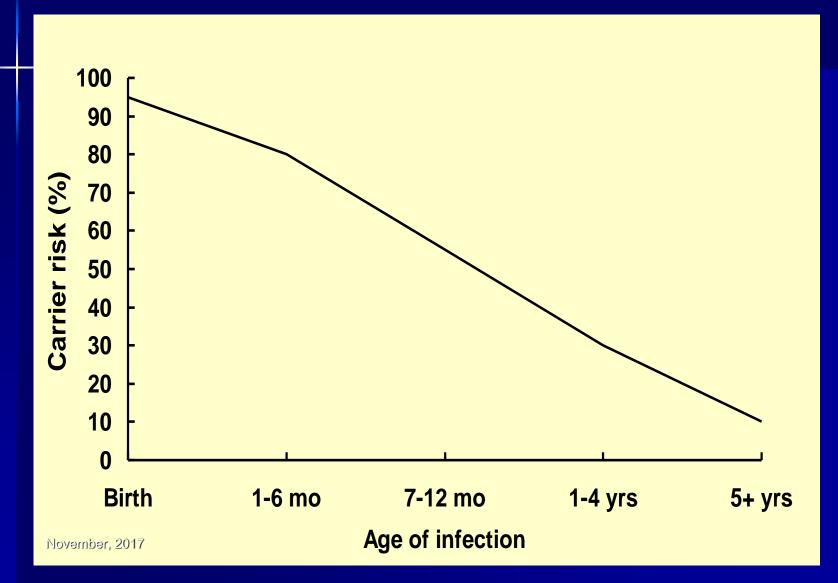
Chain of infection

• Agent: Double strand DNA.

Serotypes adw, ayw, adr, ayr.

- **Reservior**: Human (case + carrier).
- I.P. 2-3 months.
- P.C. One week of I.P. + illness period + carriage.
- Carriage depends on age at infection;
 - <5 yrs, 30%-90% chronicity
 - >5 yrs, 2%-10% chronicity

Risk of Chronic HBV Carriage by Age of 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

Parenteral

Sexual

Perinatal



November, 2017

Modes of transmission:

 Percutaneous and permucosal exposure to: -infective body fluids Blood transfusion -Organs transplants -Sharing needles - Haemodialysis - Needlestick -Tattooing – Razors & toothbrushes.

Modes of transmission:



- Sexual transmission.
- Perinatal transmission especially when HBs Ag carrier mothers are also HBe Ag positive.

Prevention and control



Hepatitis B Vaccine

Subunit recombinant HBs Ag IM in the deltoid region.

- 3 dose series, typical schedule 0, 1, 6 months no maximum time between doses (no need to repeat missed doses or restart)
- Wide scale immunization of infants (revise compulsory vaccination schedule).
- Immunization of high risk persons.

Haemodialysis patients.Bleeding disorders.Susceptible households.Health care personnel.

Prevention and control

Blood banks:

Avoid donors from risky groups. Education & history taking. Testing for HBs Ag.

Discourage:

Tattooing, Drug abuse, Extramarital sexual relations.

Needle stick

Single dose of HBIG (24 hours). Vaccine series.

Prevention and control



Sexual exposure

- Single dose of HBIG (14 days) and
- Vaccination.

Infants to HBsAg +ve mothers.

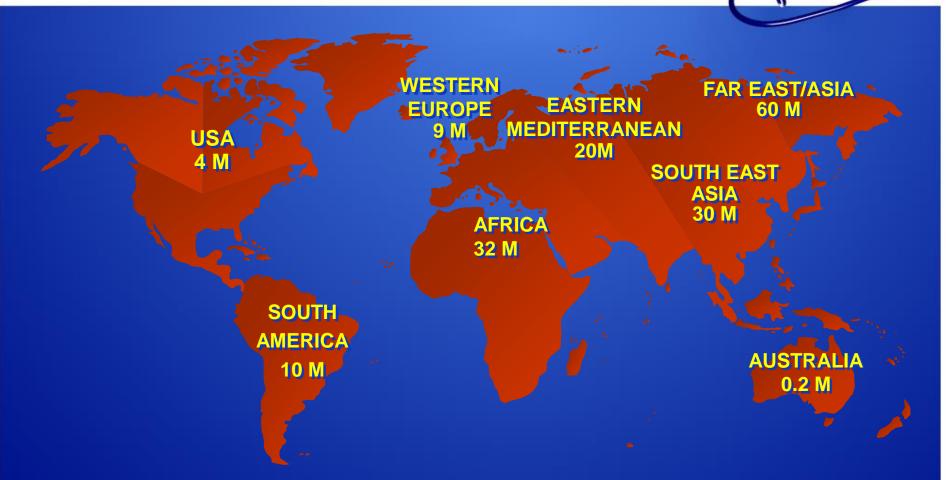
- 0.5 ml HBIG (IM).
- First dose of the vaccine.
- 2nd & 3rd doses at 1 & 6 months later.
- Health care personnel.
 Nole haiversal precautions



Hepatitis C

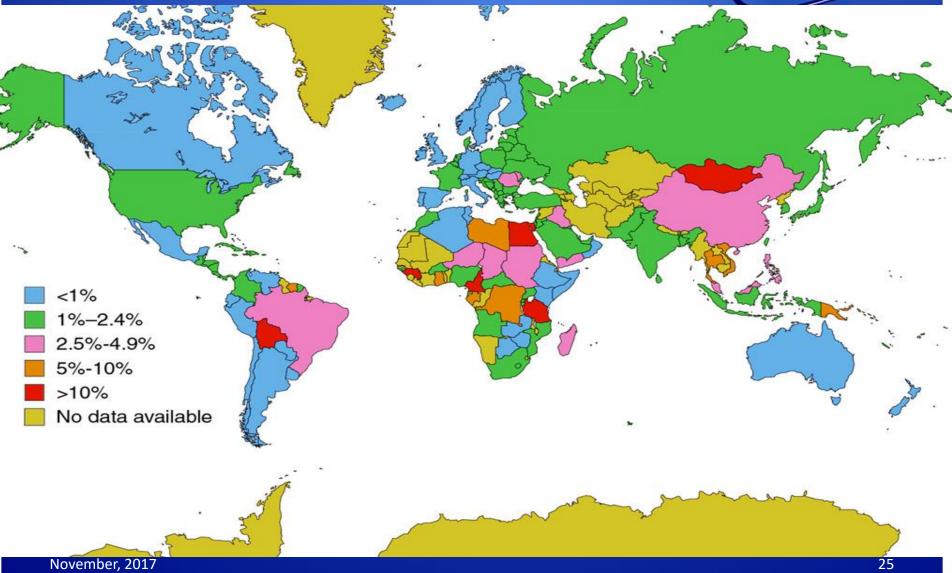
November, 2017

Hepatitis C



170 Million Hepatitis C virus (HCV) carriers 3-4 MM new cases / year

Hepatitis C



AGE SPECIFIC PREVALENCE OF ANTIBODY TO HCV/ANTI-HCV AMONG HEALTHY SAUDIS

Age Group		Community Based Study			
(years)	No. tested	Anti-HCV Pos. (%)	Location		
1 – 10	1214	0.6	Central Province		
	490	0.0	Eastern Province		
	677	0.4	North-Western Province		
	1096	0.9	South-Western Province		
	1019	1,9	Southern Province		
10 – 19	504	6 (1.2)	Gizan		
20 – 29	361	4 (1.1)	Gizan		
30 - 39	290	6 (2.1)	Gizan		
40 – 49	183	6 (3.3)	Gizan		
> 50	144	5 (3.5)	Gizan		
Total	1482	27 (1.8)	Gizan		

November, 2017

Al-Faleh et al, Hepatology Vol. 14(2), 1991²⁶

PREVALENCE OF ANTIBODY TO HCV TO SAUDI HIGH RISK GROUPS

High Risk Group	No. Tested	No. Pos.	%	Location
Hemophiliacs	28	22	78.6	KKUH, Riyadh
Thalassaemia and sickle cell disease	78	26	33.3	KKUH, Riyadh
β-thalassaemia major	20	14	70.0	KKUH, Riyadh*
Sickle cell anaemia	55	10	18.2	KKUH, Riyadh*
Patients with sexually transmitted diseases	220	35	15.9	KKUH, Riyadh*

2nd-generation anti-HCV tests and confirmation were only done November, 2017 in this study.

ANTI-HCV IN <u>HAEMODYLYSIS</u> PATIENTS IN SAUDI POPULATION

Author	No. of Persons	Type of Test	%
Fakunle et al	895	ELISA I	53.7
Al-Mugeriren et al	20 Children	ELISA I	45.0
Ayoola et al	74	ELISA I	41.9
Huraib et al	22 HD Centre 1147 Persons	ELISA II	68.8

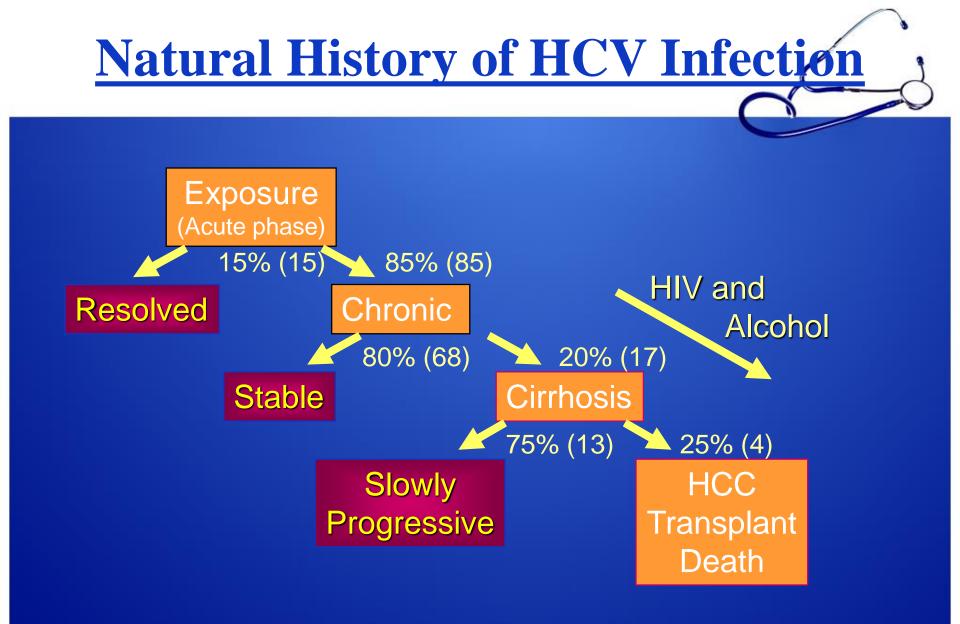
Hepatitis C Virus Genotypes



11 (6 major) with many subtypes and quasispecies

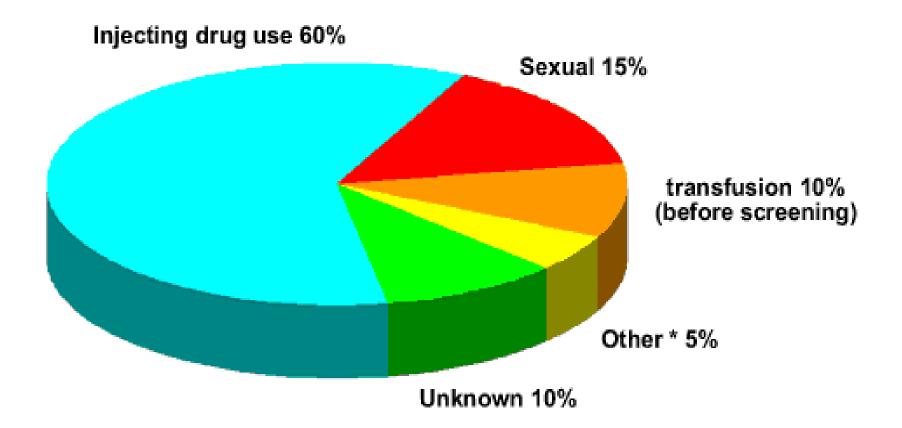
The predominate genotype in Saudi is Genotype 4 (62.9%)

- Europe & America Genotype 1→ 75 (24.8) %) → severe disease
- Genotype 2 = 10.8 (7.4) %
- Genotype 3 = 5.8 (5.9) %
- Genotype 1 & $4 \rightarrow$ Poor response to therapy



MJ Semin Liver Dis 1995; 15: Management of Hepatitis C NIH Consensus Statement 1997; March 24-26:15(3).

Sources of Infection for Persons with Hepatitis C



*Nosocomial: Health-care work; Perinatal

Source: Centers for Disease Control and Prevention



Blood transfusion



1:100,000 in US

November, 2017

IV drug abuse



80% infected in first year

Un-common HCV Transmission Modes



?

Household transmission



Vertical transmission mother - Child

1-5%

Needle stick injury

November, 2017

3%

HCV Counseling

Other Transmission Issues

- HCV not spread by kissing, hugging, sneezing, coughing, food or water, sharing eating utensils or drinking glasses, or casual contact
- Do not exclude from work, school, play, childcare or other settings based on HCV infection status



Features of Hepatitis C Virus Infection

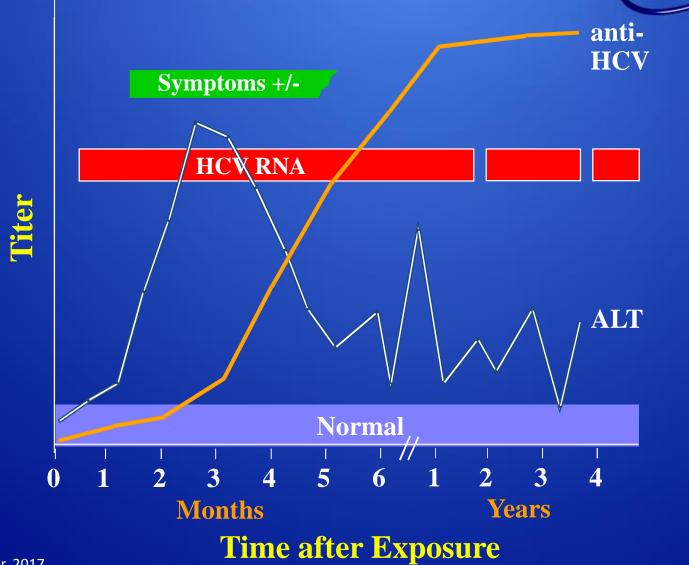
Incubation periodAverage 6-7 weeks Range 2-26 weeks **Acute illness (jaundice)** Mild (<20%) **Case fatality rate** Low 60%-85% **Chronic infection Chronic hepatitis** Age-related 10%-70% **Cirrhosis**-<5%-20%**Mortality from CLD** 1%-5%

Chronic Hepatitis C Factors Promoting Progression or Severity

- Increased alcohol intake
- Age > 40 years at time of infection
- HIV co-infection
- Other

 Male gender
 Chronic HBV co-infection

Serologic Pattern of Acute HCV Infection with Progression to Chronic Infection





November, 2017

Perinatal Transmission of HCV

- Transmission only from women HCV-RNA positive at delivery
 - Average rate of infection 6%
 - Higher (17%) if woman co-infected with HIV
 - Role of viral titer unclear
- No association with
 - Delivery method
 - Breastfeeding
- Infected infants do well
 Severe hepatitis is rare

Sexual Transmission of HCV

Case-control, cross sectional studies

 Infected partner, multiple partners, early sex, non-use of condoms, other STDs, sex with trauma, Partner studies

 Low prevalence (1.5%) among long-term partners
 infections might be due to common percutaneous exposures (e.g., drug use), BUT
 Male to female transmission more efficient
 more indicative of sexual transmission

Household Transmission of HCV

- Rare but not absent
- Could occur through percutaneous/mucosal exposures to blood
 - Contaminated equipment used for home therapies
 - IV therapy, injections
 - Theoretically through sharing of contaminated personal articles (razors, toothbrushes)

Public Health Service Guidelines for Anti-HCV-Positive Persons

Anti-HCV-positive persons should:

- Be considered potentially infectious
- Keep cuts and skin lesions covered
- Be informed of the potential for sexual transmission
- Be informed of the potential for perinatal transmission
 - no evidence to advise against pregnancy or breastfeeding

Anti-HCV-positive persons should not:

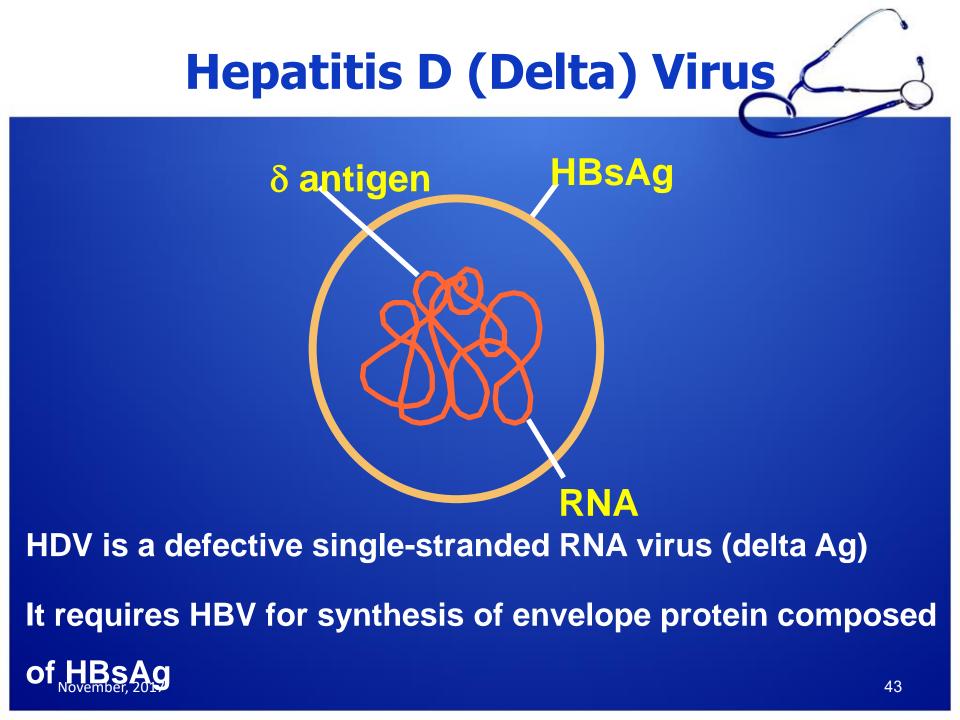
- Donate blood, organs, tissue, or semen
- Share household articles (e.g., toothbrushes, razors)

CAU



Hepatitis D

November, 2017



Hepatitis D - Clinical Features

Coinfection with HBV

- severe acute disease
- low risk of chronic infection

Superinfection on top of chronic HBV

- usually develop chronic HDV infection
- high risk of severe chronic liver disease

Hepatitis D Virus Modes of Transmission

Percutanous exposures

 injecting drug use

 Permucosal exposures

 sex contact

Hepatitis D - Prevention

- HBV-HDV Coinfection
 - Pre or postexposure prophylaxis to prevent HBV infection (HBIG and/or Hepatitis B vaccine)
- HBV-HDV Superinfection
 - Education to reduce risk behaviors among persons with chronic HBV infection



Hepatitis E

November, 2017

Hepatitis E - Clinical Features

- Incubation period:
- Case-fatality rate:
- Illness severity:Chronic sequelae:

Average 40 days Range 15-60 days Overall, 1%-3% Pregnant women, 15%-25% Increased with age None identified

Hepatitis E - Epidemiologic Features

- Most outbreaks associated with fecally contaminated drinking water
- Minimal person-to-person transmission

Geographic Distribution of Hepatitis E

Outbreaks or Confirmed Infection in >25% of Sporadic Non-ABC Hepatitis



Viral Hepatitis - Overview

Type of Hepatitis

	A	B	C	D	Ε
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	ensure safe drinking water
November, 2017				modification	CENTERS FOR DISEASE CONTROL AND PREVENTION 51



References

1-Nelson KE, Thomas L. Viral hepatitis. In: Infectious disease Epidemiology, theory and Practice. 2nd edition . Edited by Nelson KE and Williams CM 2007. Published by Jones & Bartlett. Toronto Pages895-939.

