

# ACUTE VIRAL HEPATITIS

- CLINICAL PRESENTATION.
- DIGNOSIS.
- EPEDEMOLOGY OF VIRAL HEPATITIS INFECTION A,B,C IN KSA.
- MANAGEMENT.

# Viral Hepatitis - Overview

## Type of Hepatitis

	<b>A</b>	<b>B</b>	<b>C</b>	<b>D</b>	<b>E</b>
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

# Diagnosis of hepatitis

- Patient history
- Physical examination
- Liver function tests
- Serologic tests

# Symptoms and Signs

## ■ Pre-icteric phase

1. Anorexia
2. Fatigue
3. Nausea
4. Vomiting
5. Arthralgia
6. Myalgia
7. Headache
8. Photophobia
9. Pharyngitis
- 10.
- 11.

## ■ Icteric phase::

1. Enlarged liver
2. Tender upper quadrant
3. Discomfort
4. Splenomegaly (10-20%)
5. General adenopathy

## ■ Post-icteric phase

# Lab Findings

1. L FT increase >5-10 times of normal
2. Markers of hepatitis B or C or A might be positive

# DD:

1. Infectious Mononucleosis
2. Drug Induced Hepatitis
3. Chronic Hepatitis.
4. Alcohol Hepatitis
5. Cholecystitis, Cholelithiasis
- 6-Auto-immun hepatitis

# MARKERS OF VIRAL HEPATITIS

- HBV MARKERS
- HCV MARKERS
- HAV MARKERS

# Hepatitis B Markers

- anti-HBc → exposure (IgM = acute)
- HBsAg → infection (carrier)
- anti-HBs → immunity
- HBeAg → viral replication
- anti-HBe → seroconversion
- HBV-DNA → viral replication:

# Hepatitis C Markers

- ANTI -HCV
- PCR-RNA HCV

# Hepatitis A Markers

- HAV igM
- HAV igG

# Hepatitis E Markers

- HEV igM
- HEV igG
- HEV RNA PCR

# AUTOIMMUN HEPATITIS MARKERS

- ANF
- ANTI MITOCHONDRIAL AB
- ANTI SMOOTH MUSCLES ABS.

# AUTOIMMUN HEPATITIS MARKERS

- ANA)
- ANTI MITOCHONDRIAL
- ANTI SMOOTH MUSCLES ABS

# INCIDENCE OF ACUTE HEPATITIS IN 5 HEPATOLOGY CLINICS IN KSA 2013

Causes of Hepatitis	HAV	HBV	HCV	AIH	DILI
KKUH	7	3	1	11	?
NGH	10	5	2	8	8
AMC	1	0	0	3	3
KFH	1	2	0	3	?
DAMMAM UN.	2	1	0	1	5
TOTAL	21	11	4	26	16

# Complications

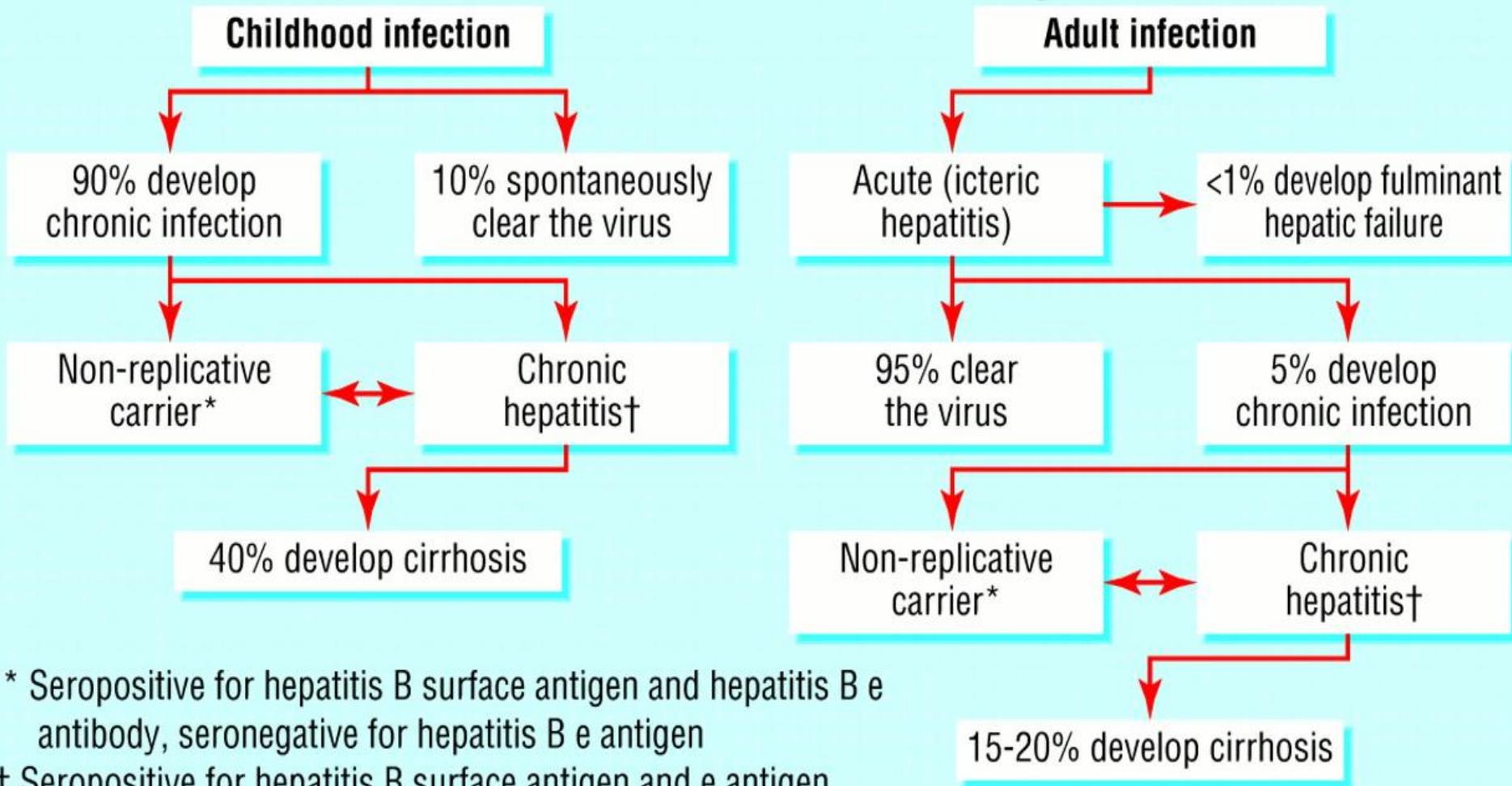
1. Chronic hepatitis → cirrhosis- HCC

2. Fulminant hepatitis

# FULMINANT HEPATITIS

- Definition: Hepatic Failure Within 8 Weeks Of Onset Of Illness.
- Manifestation: Encephalopathy and Prolonged PT
- Histopathology: Massive Hepatic Necrosis.

# Natural History



\* Seropositive for hepatitis B surface antigen and hepatitis B e antibody, seronegative for hepatitis B e antigen

† Seropositive for hepatitis B surface antigen and e antigen, seronegative for hepatitis B e antibody

# Hepatitis B Virus

## Modes of Transmission

- Sexual
- Parenteral
- Perinatal

# Concentration of Hepatitis B Virus in Various Body Fluids

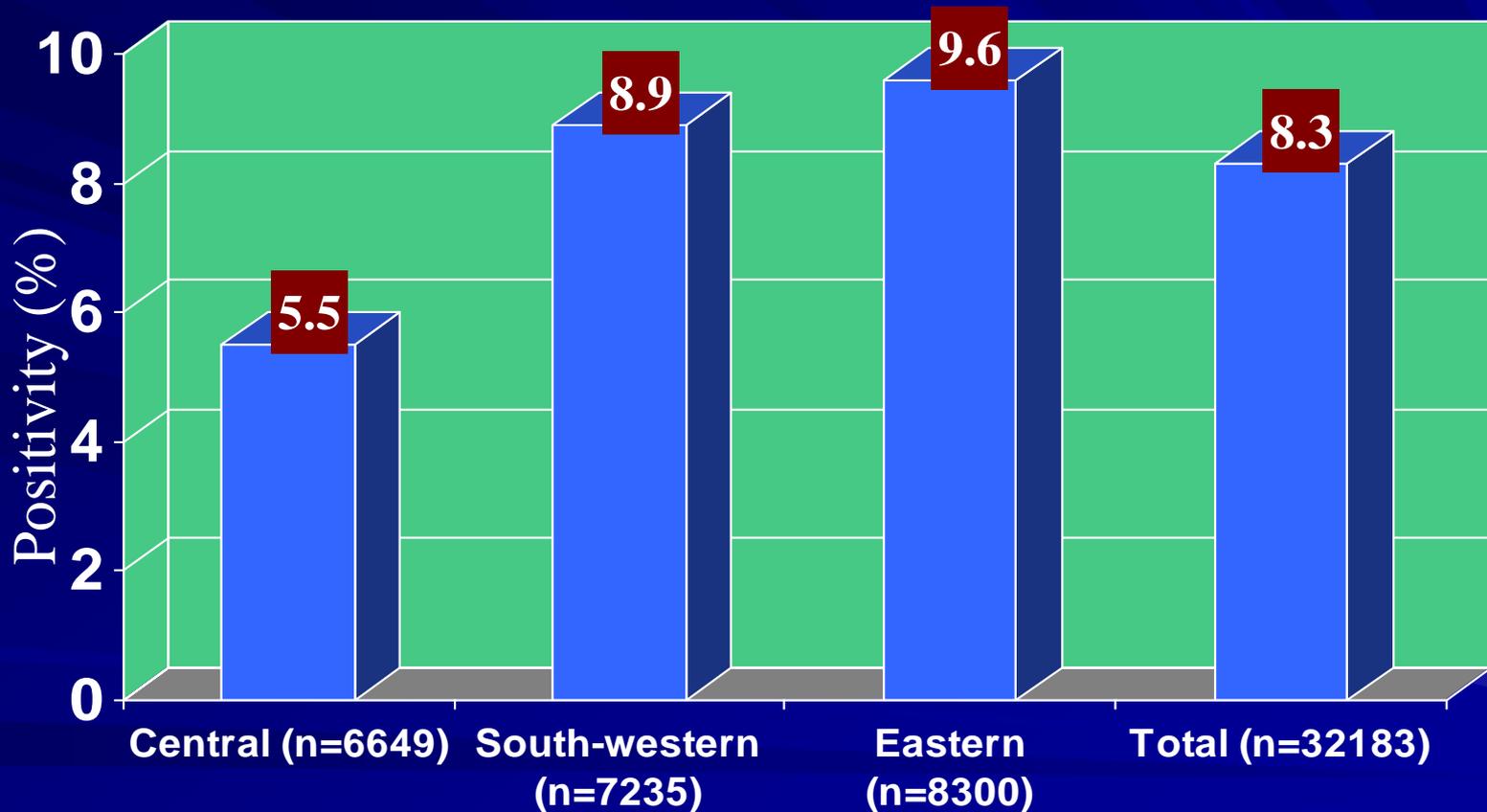
<b>High</b>	<b>Moderate</b>	<b>Low/Not Detectable</b>
blood	semen	urine
serum	vaginal fluid	feces
wound exudates	saliva	sweat
		tears
		breastmilk

# Possible transmission route of HBV in KSA

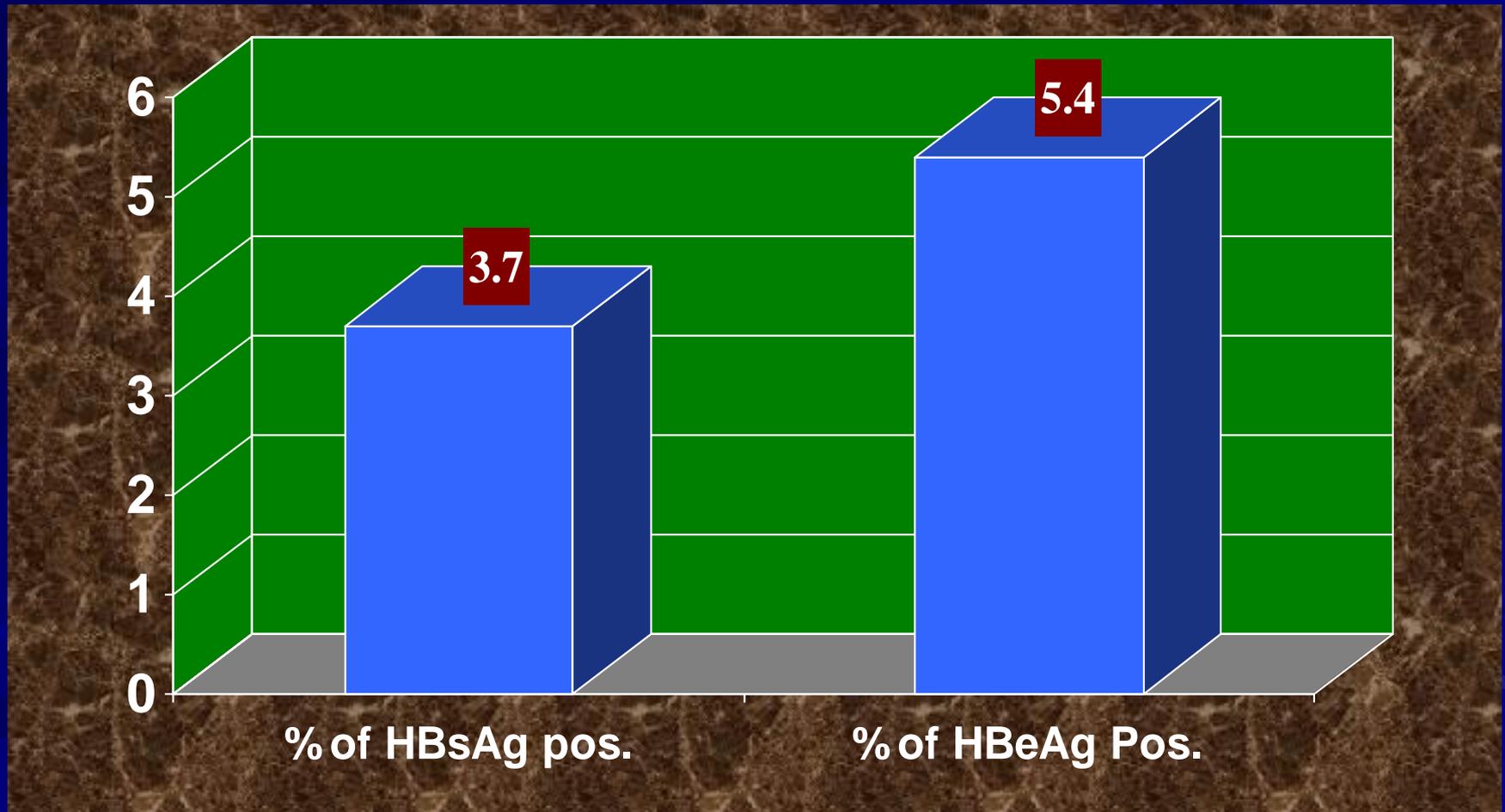
- 1-Horizontal transmission (person to person) is the main transmission route
- 2-Perinatal transmission (positive HBSAG mothers) especially if they are HBEAG positive
- 3- Heterosexual transmission
- 4-Illegal injection drug use
- 5- Contaminated equipment used for therapeutic injections and other health care related procedures
- 6- Folk medicine practice
- 7-Blood and blood products transfusion without prior screening

**HBV INFECTION  
before and after  
vaccination program**

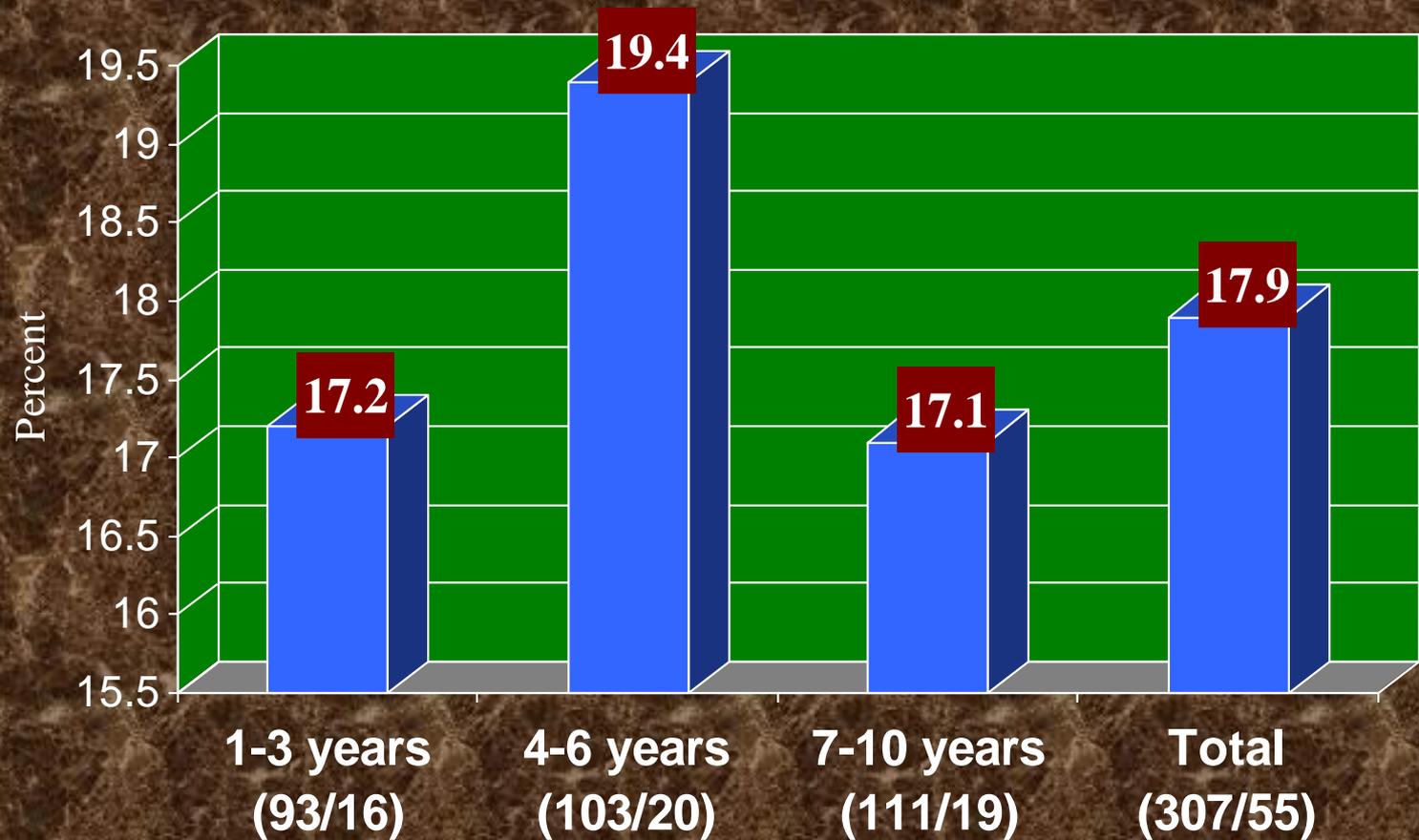
# OVERALL PREVALENCE OF HBsAg AMONG SAUDIS IN THE 80'S ACCORDING TO REGIONS



# PREVALENCE OF HBeAg AMONG HBsAg POSITIVE SAUDIS PREGNANT WOMEN (n = 20920)



# FREQUENCY OF HBeAg AMONG HBsAg POSITIVE SAUDI CHILDREN (n=307)

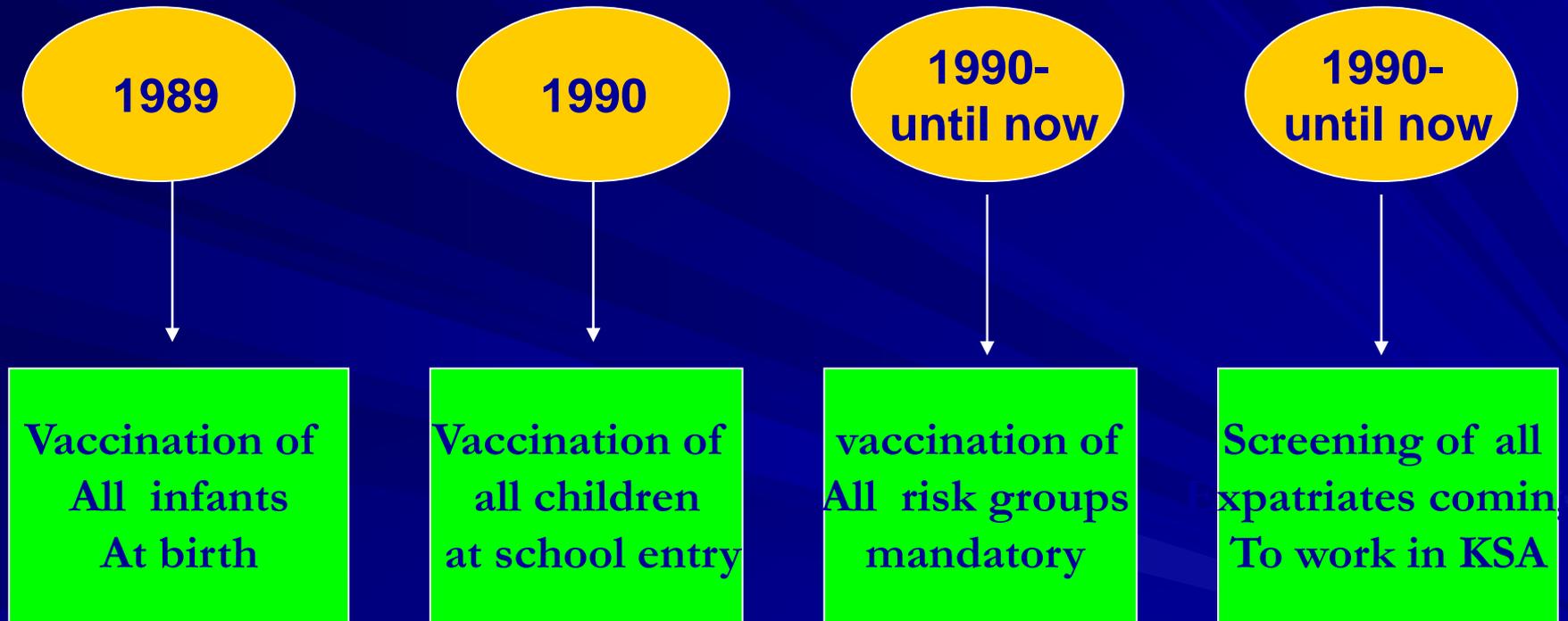


# PREVENTION STRATEGIES OF MINISTRY OF HEALTH IN KSA

Introducing HBV vaccine in EPI program; and

- Mandatory screening of blood donors and expatriates.
- Vaccination of risk groups.
- Health education especially among medical personnel.

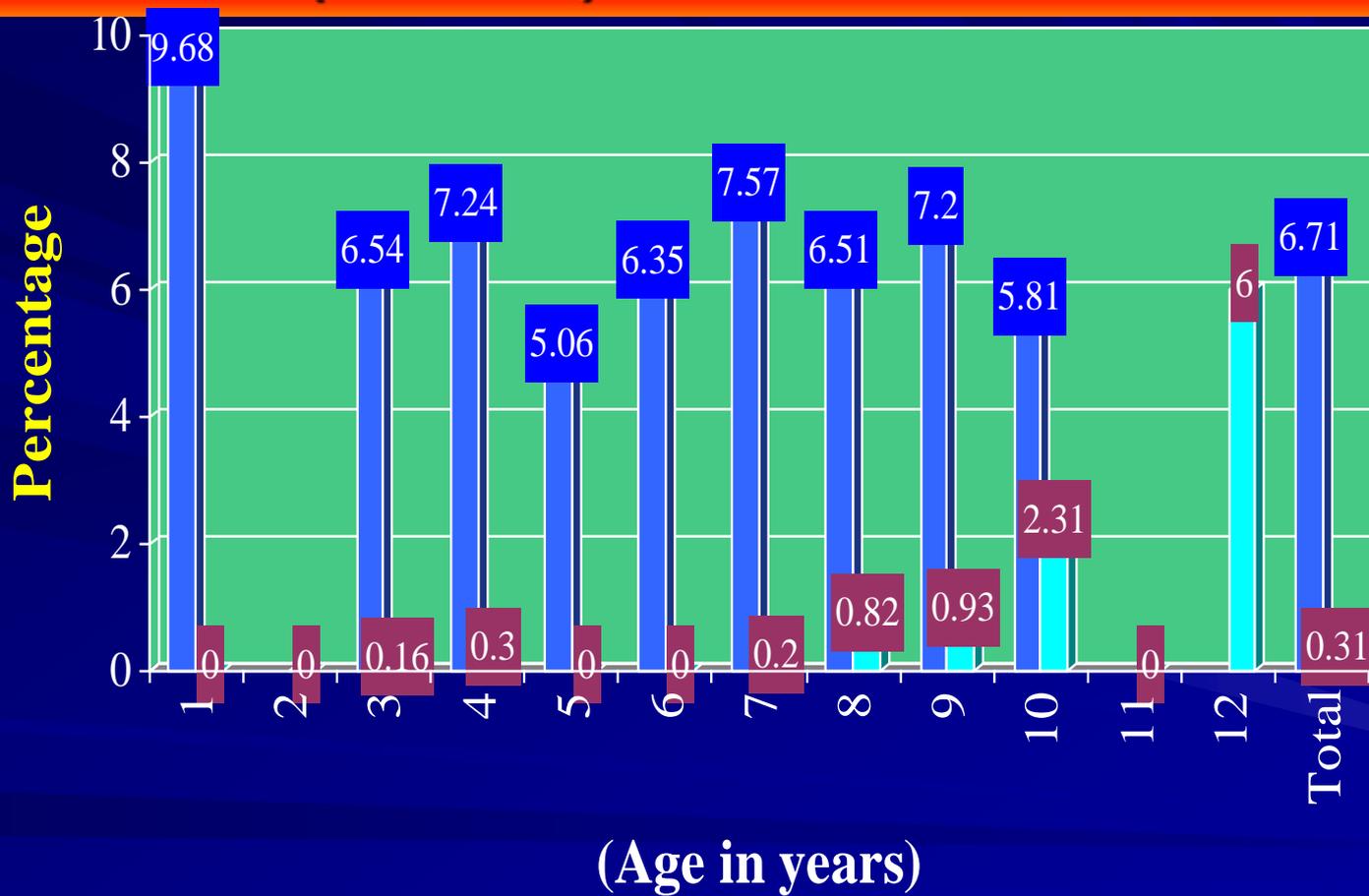
# History of HBV infection control in KSA



# THE CURRENT EPI IN THE KINGDOM OF SAUDI ARABIA

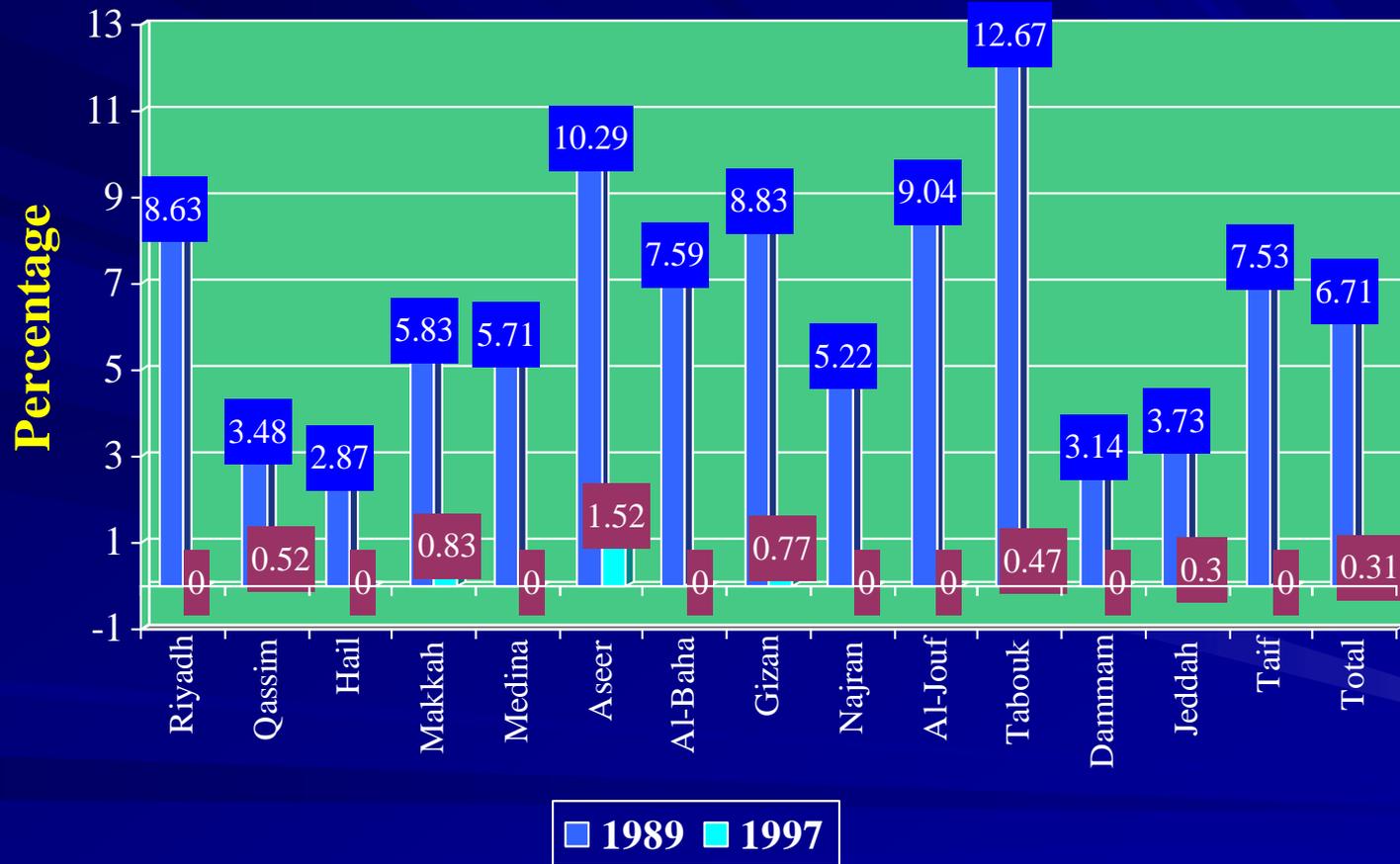
- |    |              |             |           |
|----|--------------|-------------|-----------|
| 1. | At birth     | BCG +       | HB1       |
| 2. | At 6 weeks   | DPT1 + OPV1 | Hb2       |
| 3. | At 3 months  | DPT2 + OPV2 |           |
| 4. | At 5 months  | DPT3 + OPV3 |           |
| 5. | At 5 months  | Measles     | HB3       |
| 6. | At 12 months | MMR         |           |
| 7. | At 18 months | (DPT + OPV) | Booster 1 |
| 8. | At 4-6 years | (DPT + OPV) | Booster 2 |

# COMPARISON OF PREVALENCE OF HBsAg AMONG SAUDI CHILDREN IN 1989 (n=4575) AND 1997 (n=5355) – ACCORDING TO AGE



■ 1989 ■ 1997

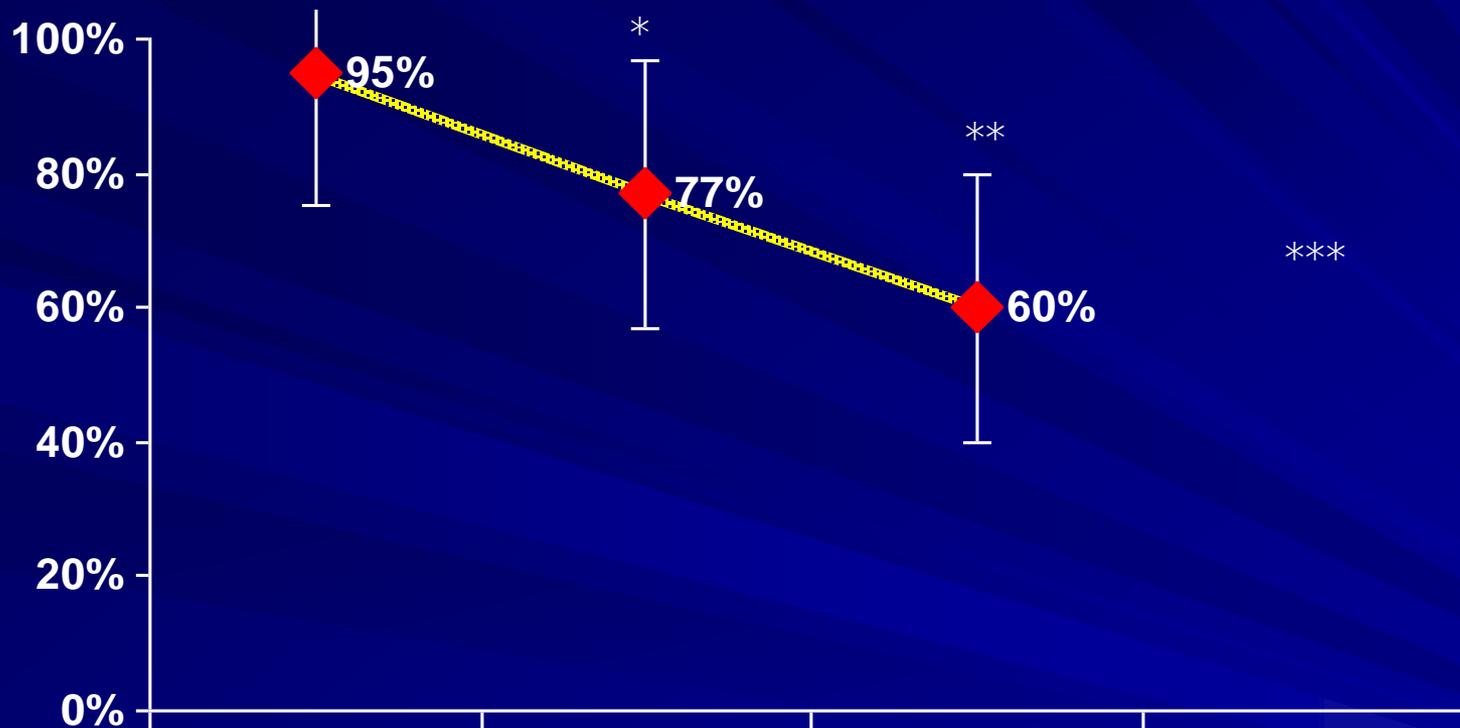
# COMPARISON OF PREVALENCE OF HBsAg AMONG SAUDI CHILDREN IN 1989 (n=4575) AND 1997 (n=5355) – ACCORDING TO REGION



# Prevalence Of HBsAg Among Saudi Population Before & After Vaccination over 18 y



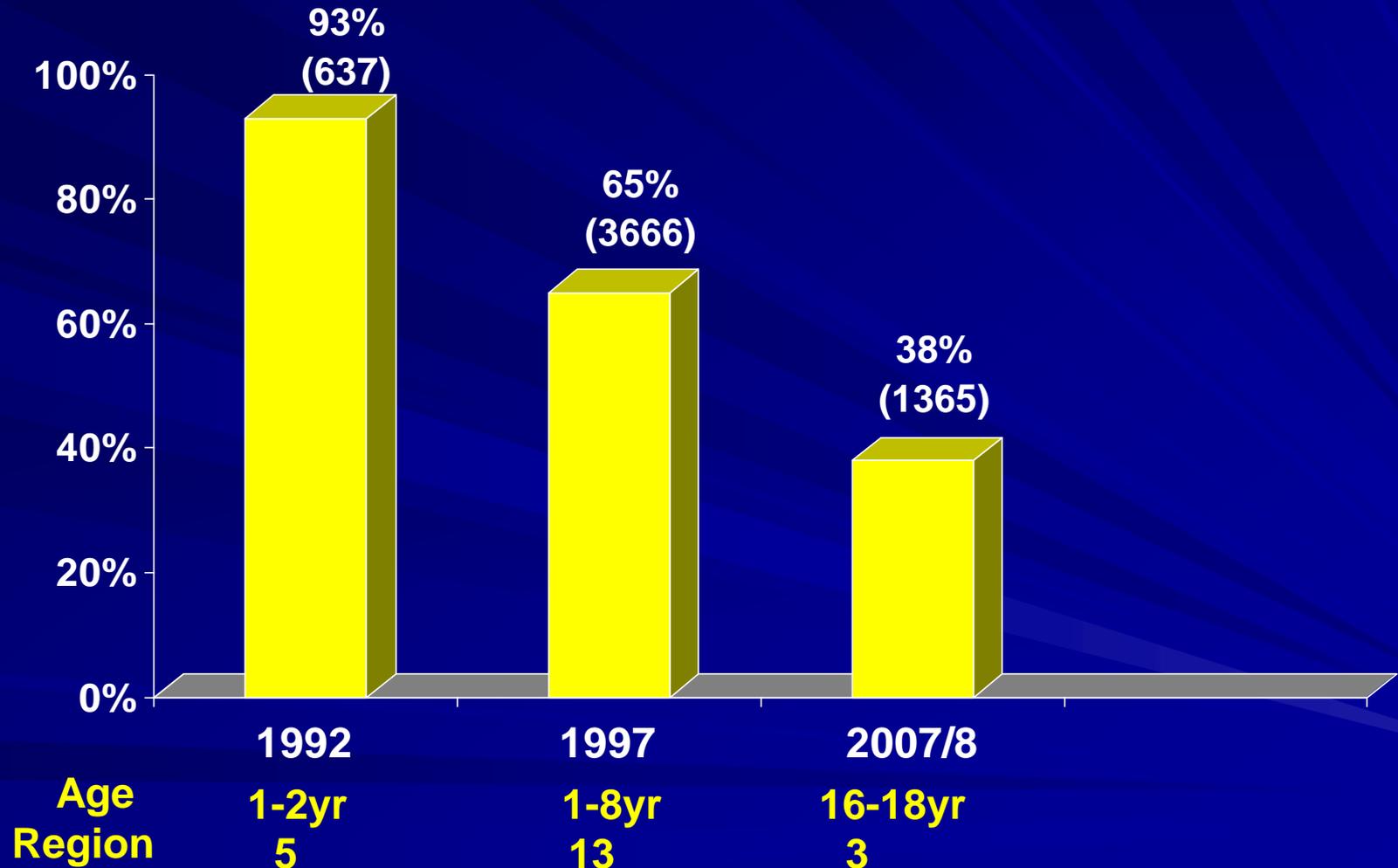
# Long Term Seroconversion Rate Over 18 Years (Anti-HBS)



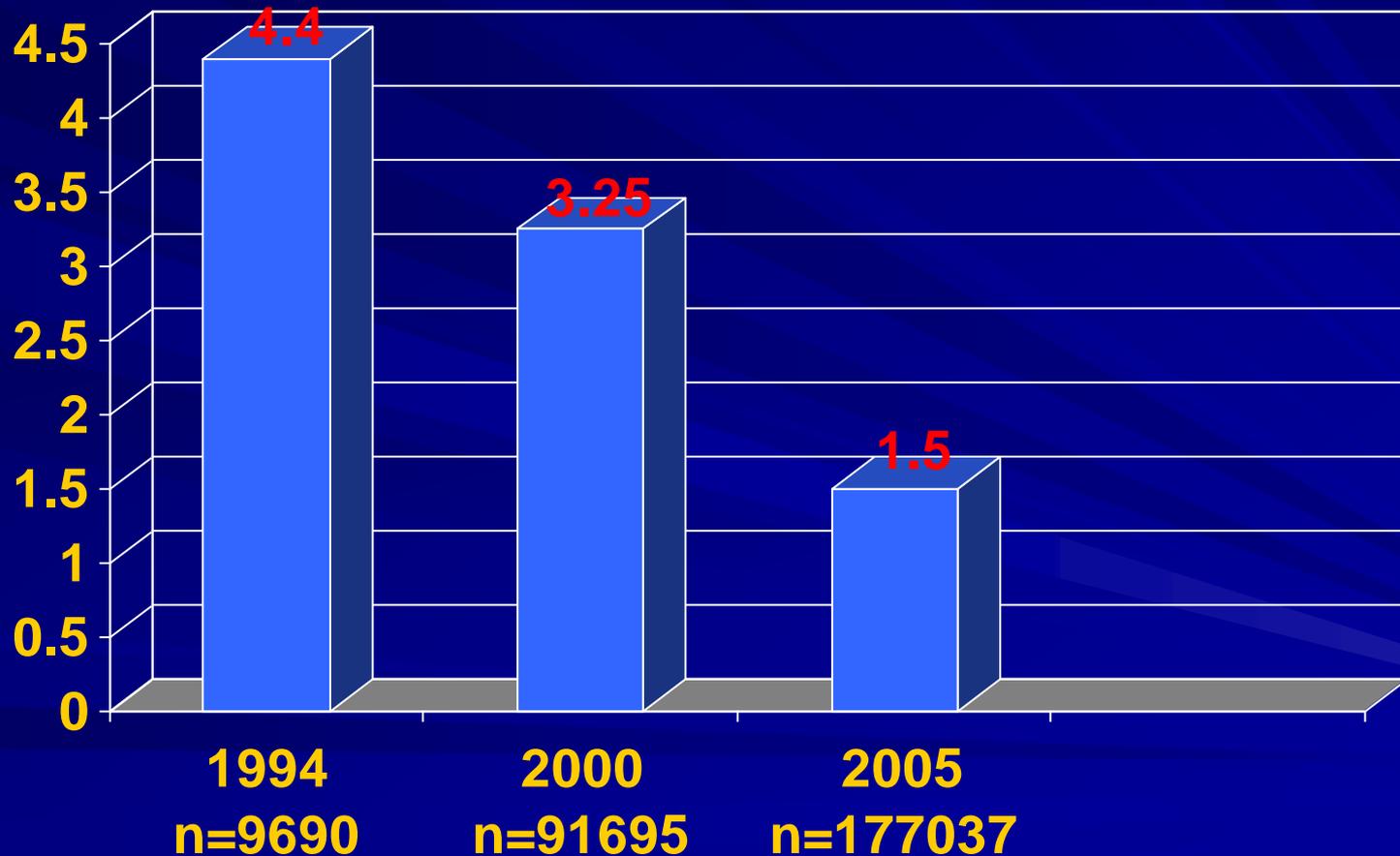
Age	1-2yr	1-12yr	16-18yr
N	637	3666	1365

\* Al Faleh et al Annals of Saudi meds 1993  
\*\* Al Faleh et al Journal of infection 1999  
\*\*\* AlFaleh et al journal of infection2008

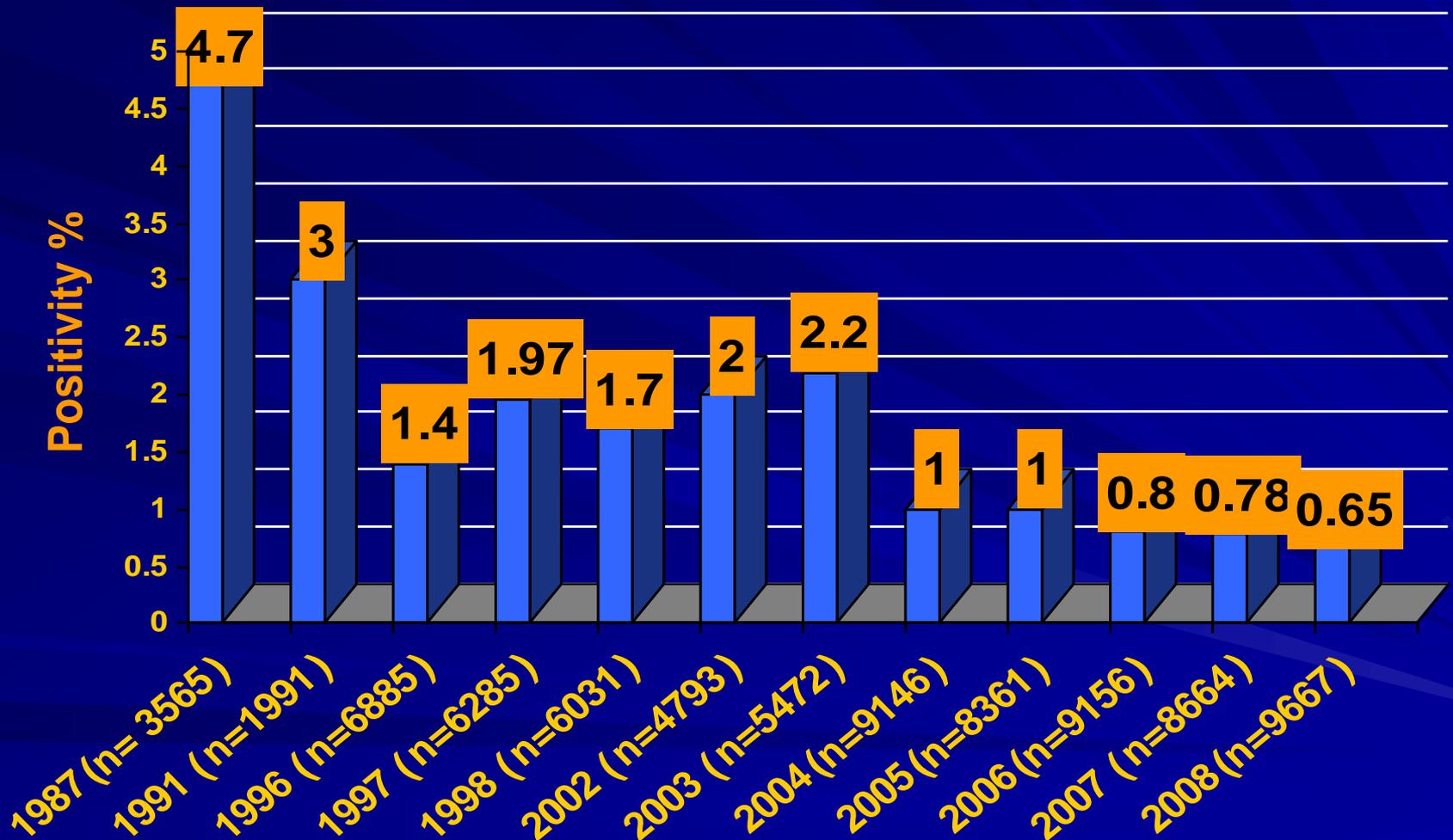
# Long-Term protection of HB- vaccine over 18 years ( anti-HBS>10IU/L)(n=1355)



# CHANGING PATTERNS OF HBsAg POSITIVITY AMONG BLOOD DONORS IN MOH,CENTRAL BLOOD BANK 1994-2005



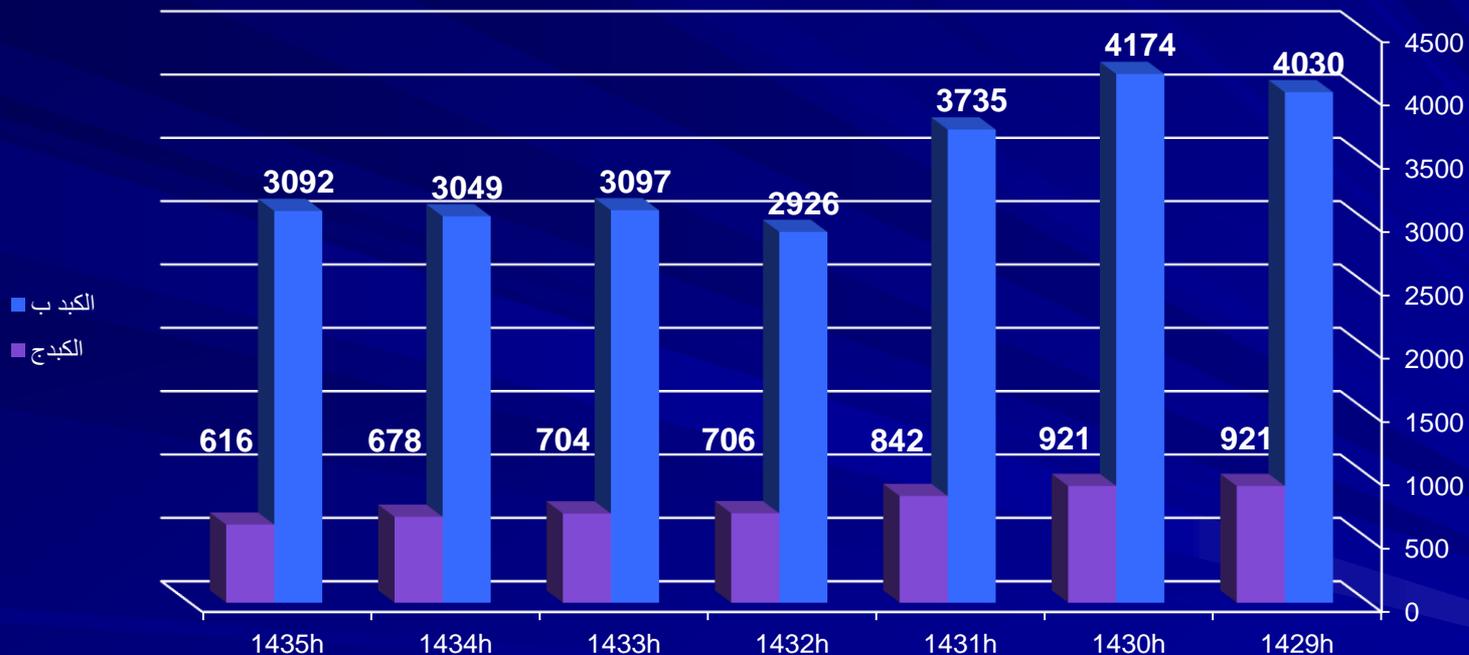
# PREVALENCE OF HBsAg POSITIVITY AMONG BLOOD DONORS IN KKHU FROM 1987 TO 2008



# PRI-MARITAL SCREENING برنامج الزواج الصحي

التهاب الكبد ب وج 1429-1435هـ			
HBV,HCV INFECTION FROM 2009-2014			
الكبد ب HBV	الكبد ج HCV	HIV	عدد المتقدمين NR.OF SCREENDS
24103	5388	512	2.131.018
1%	0.3%	%0.02	

# عدد حالات التهاب الكبد ب وج 1429-1435هـ (2009-2014)NR.OF POSITIVE HBV&HCV CASES HCV=RED



# .Case report

- **5/11/18** Sara is 35 Y/O Teacher ,living in TABUK.
- She came for marriage consultation.

# Lab. results

- 12/11/18 :
- ALT 40/L(21-72)
- AST 30 U/L (17-59)
- ALKALINE PHOSPHATASE 80.0 U/L.
- YGT 40,0U/L
- BIL.1MG/DL (0.0-1.4)
- ALB.4.6 g/l(3.5-5.0)
- INR (NORMAL)

# Lab. results

HBsAG : POSITIVE

PCR QUANTITATIVE :200IU/ML

# FINAL DIAGNOSIS

- Inactive CARRIER OF VIRAL HEPATITIS B

- Her main problem whenever she got engaged and do the marriage screening, nobody accept to proceed with the marriage.

■ How can we solve the problem ?

# HCV INFECTION

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# Transmission of HCV

- **Percutaneous**
  - Injecting drug use
  - Clotting factors before viral inactivation
  - Transfusion, transplant from infected donor
  - Therapeutic (contaminated equipment, unsafe injection practices)
  - Occupational (needlestick)
- **Per mucosal**
  - Perinatal
  - Sexual

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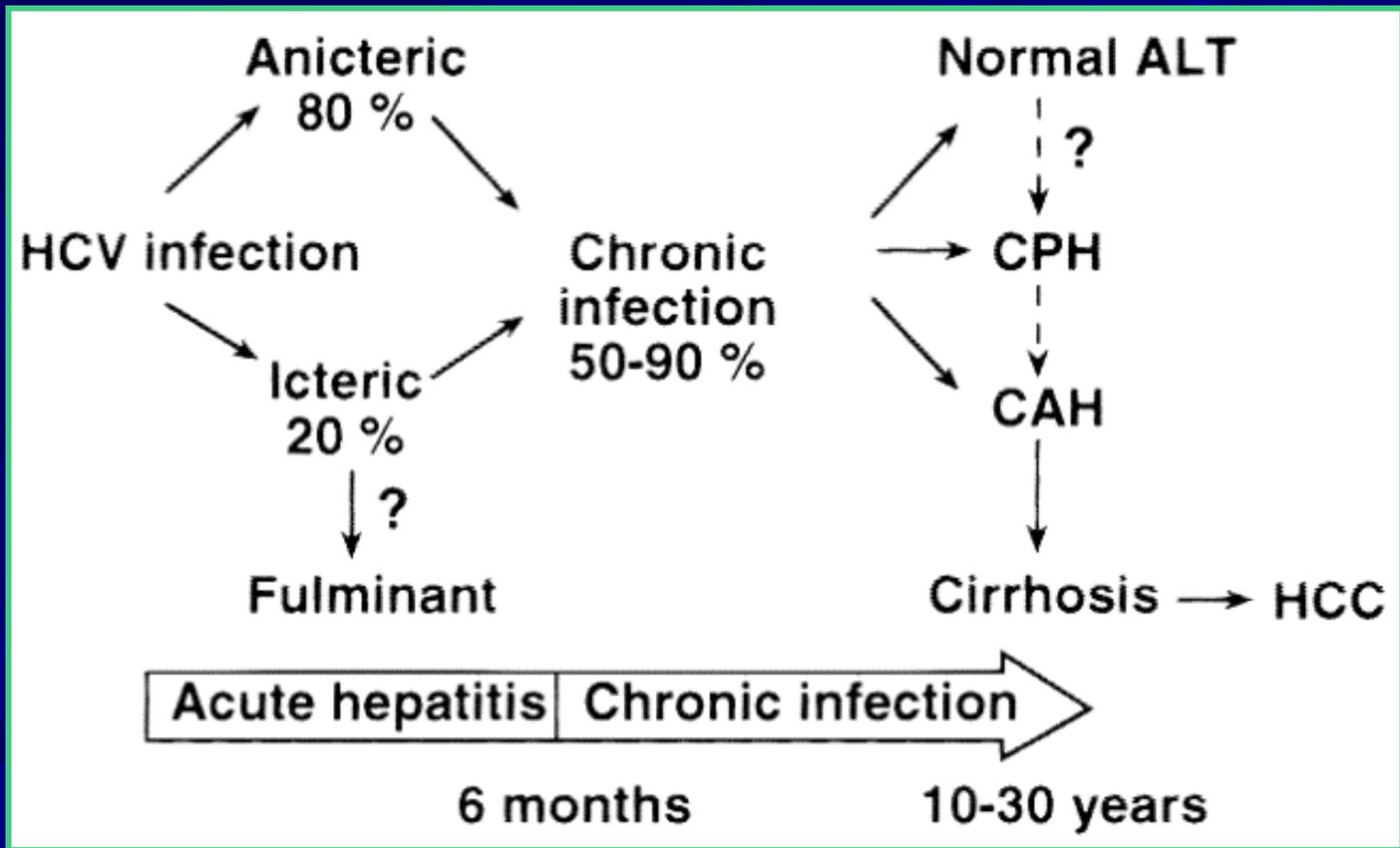
## Nosocomial Transmission of HCV

- **Recognized primarily in context of outbreaks**
- **Contaminated equipment**
  - hemodialysis\*
  - endoscopy
- **Unsafe injection practices**
  - plasmapheresis,\* phlebotomy
  - multiple dose medication vials
  - therapeutic injections

\* Reported in U.S.

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# Natural history



# Overall prevalence rate of HCV infection in KSA among children and adolescent during the last 18 yrs.

1989		1997		2008	
No. of children	Positive (%)	No. of children	Positive (%)	No. of students	Positive (%)
4496	39* (0.87%)	5350	2** (0.04%)	1357	(5)3 0.22%
Diagnostic test only by 1 <sup>st</sup> -generation EIA kit.		Diagnostic test by 3 <sup>rd</sup> -generation EIA kit and confirmatory test by RIBA kit.		Diagnostic test by PCR for anti- HCV Positive cases.	

\* ALFaleh et al. Hepatology 1991

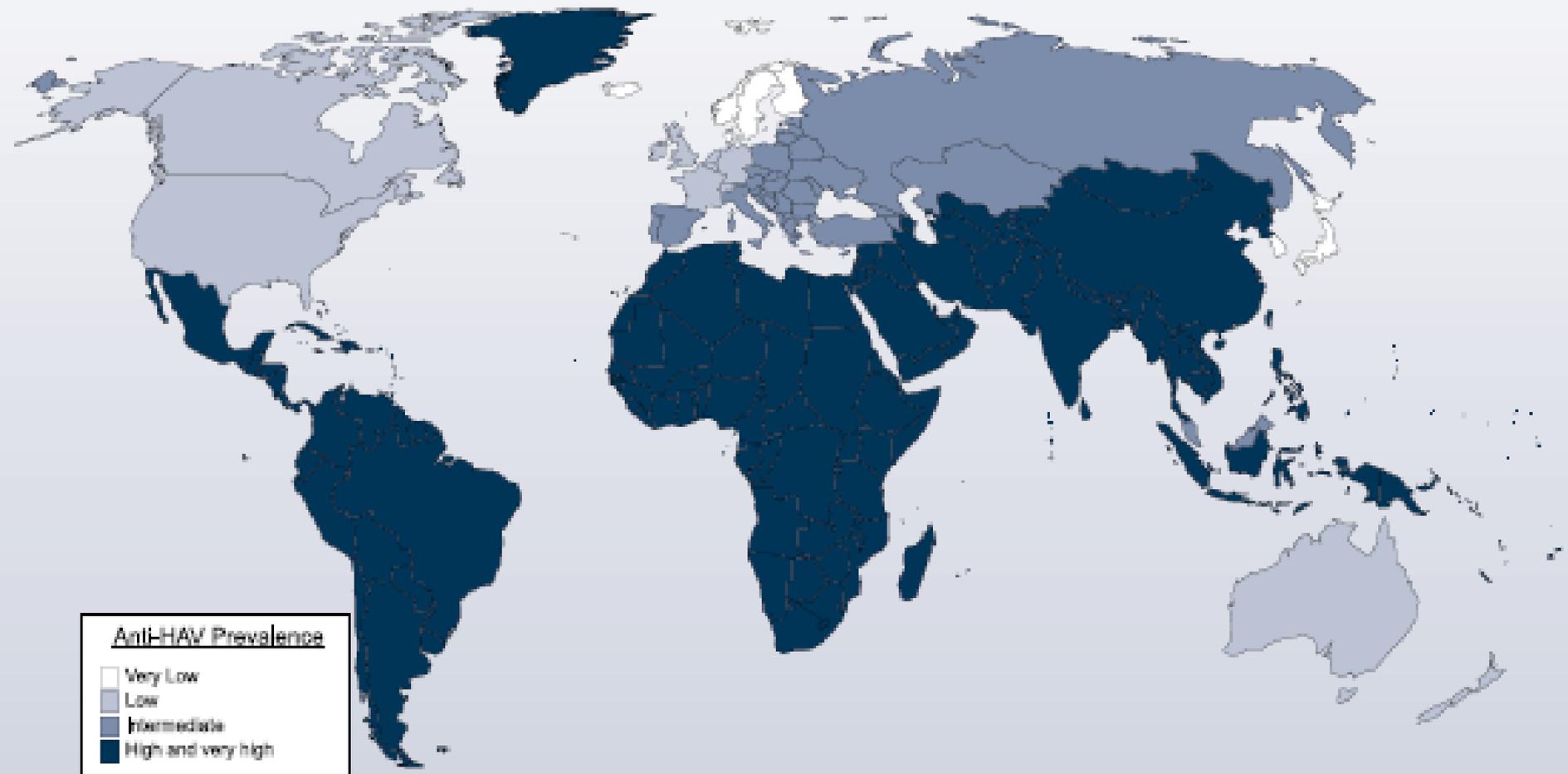
\*\* ALFaleh Ann Saudi Med. 2003

# Prevention Of HCV Transmission

- Avoiding shared use of Razors or brushes and any item that pierces the skin.
- Strict adherence of the universal precautions in health facilities.
- Educating and training of HCW's to the proper use of standard precautions
- Folk medicine?!

# HAV INFECTION

# Geographic Distribution of HAV Infection



# HEPATITIS A VIRUS TRANSMISSION

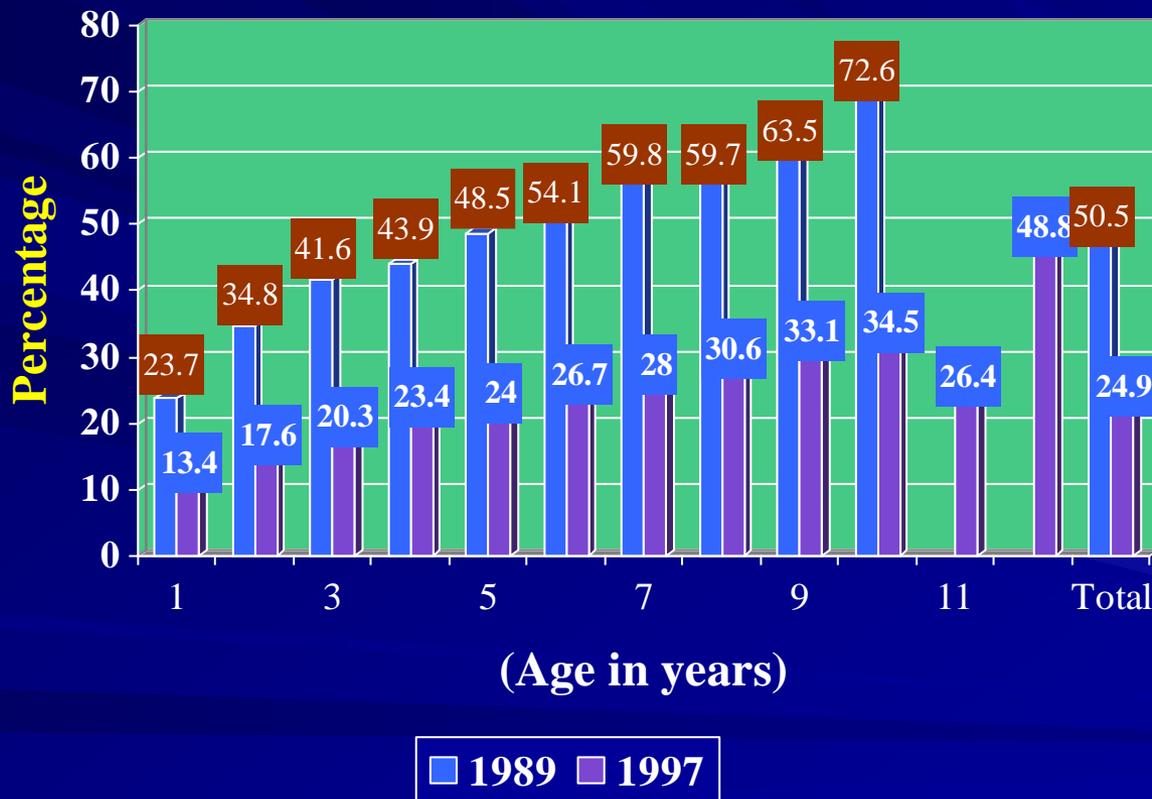
- **Close personal contact**  
(e.g., household contact, sex contact, child day-care centers)
- **Contaminated food, water**  
(e.g., infected food handlers)
- **Blood exposure (rare)**  
(e.g., injection drug use, rarely by transfusion)



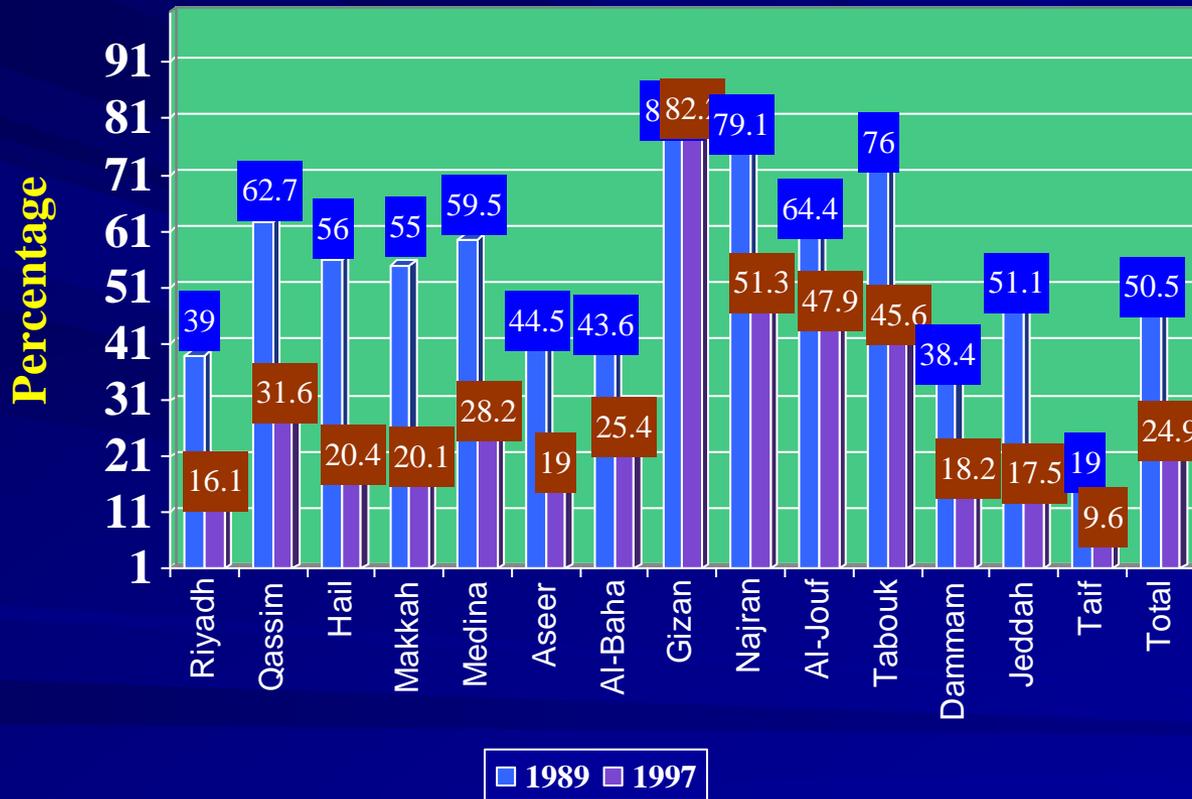
# Modes of HAV transmission

- Faeco-oral route (95%)
  - ==> person-to-person contact
  - ==> contaminated food or water
  - ==> salads and fruits washed in contaminated water
  - ==> contaminated shellfish
- Infected plasma (<5%)
- Sexual route (<5%)

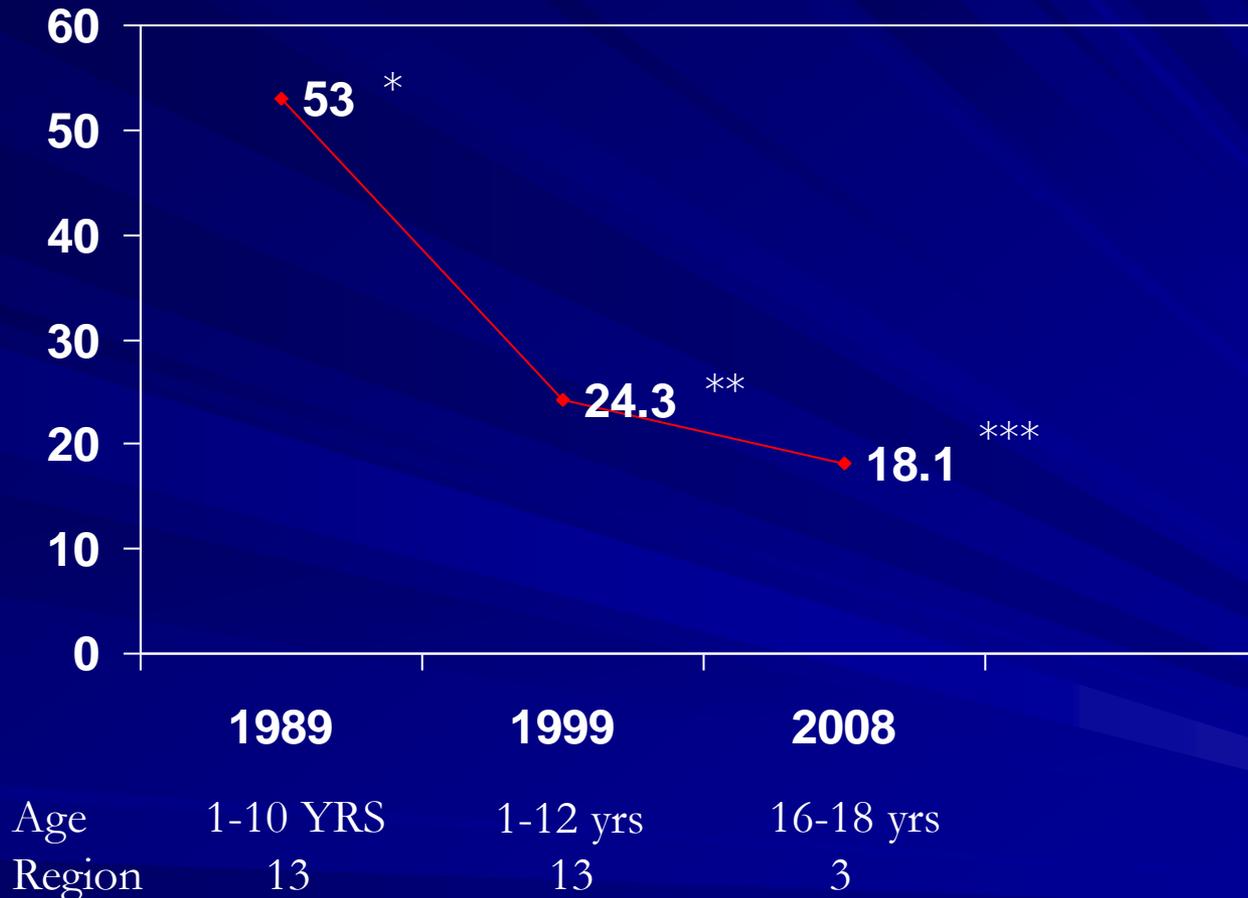
# COMPARISON OF PREVALENCE OF ANTI-HAV AMONG SAUDI CHILDREN IN 1989 (n=4375) AND 1997 (n=5255) – ACCORDING TO AGE



# COMPARISON OF PREVALENCE OF ANTI-HAV AMONG SAUDI CHILDREN IN 1989 (n=4375) AND 1997 (n=5255) – ACCORDING TO REGION



# Changing pattern of Hepatitis A prevalence within the Saudi population over 18 yrs



\*AlRashed R. Ann SM 1997

\*\* AlFaleh et al SMJ 1999

\*\*\* AlFaleh et al WJG 2008

# PREVENTING HEPATITIS A

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

# HEPATITIS A VACCINES

## Recommended Dosages of Hepatitis A Vaccines

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

\* EL.U. – Enzyme-linked immunosorbent assay (ELISA) units

\*\* Units

# has 2-phenoxyethanol as a preservative

## has no preservative

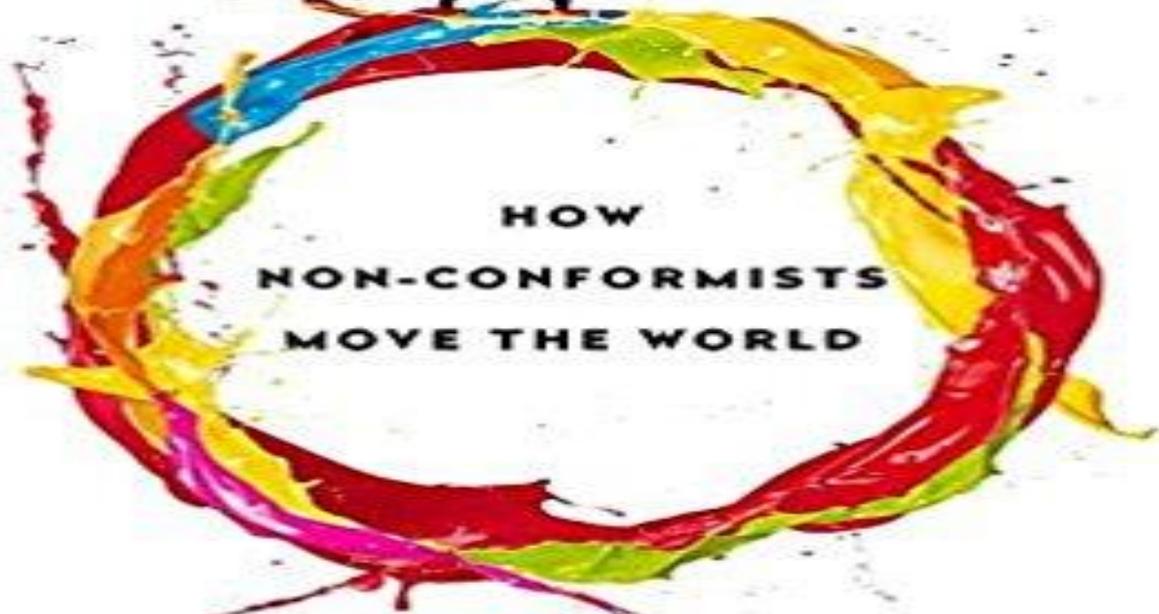


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