# Prevalence of hepatitis B in pregnancy and vertical transmission rate of HBV in Africa: A systematic review



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## **Outline**

**Background Objectives** Methodology Results **Discussion Conclusion** Recommendations

## Why this research questions

#### To demonstrate the importance of:

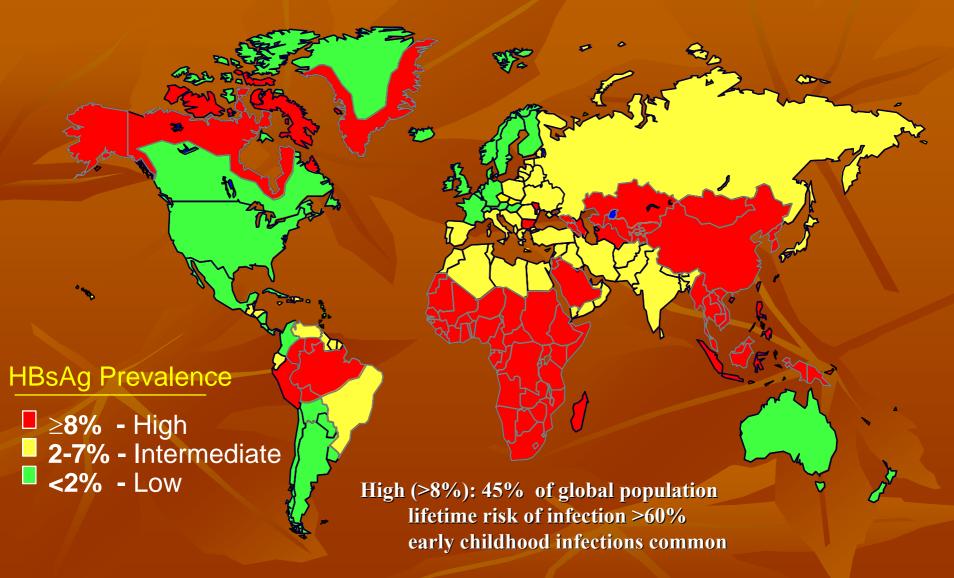
**♦ Systematic** HBV screening during pregnancy in Africa

**◆ Taking <u>special care</u>** of newborns from positive mothers

## Background

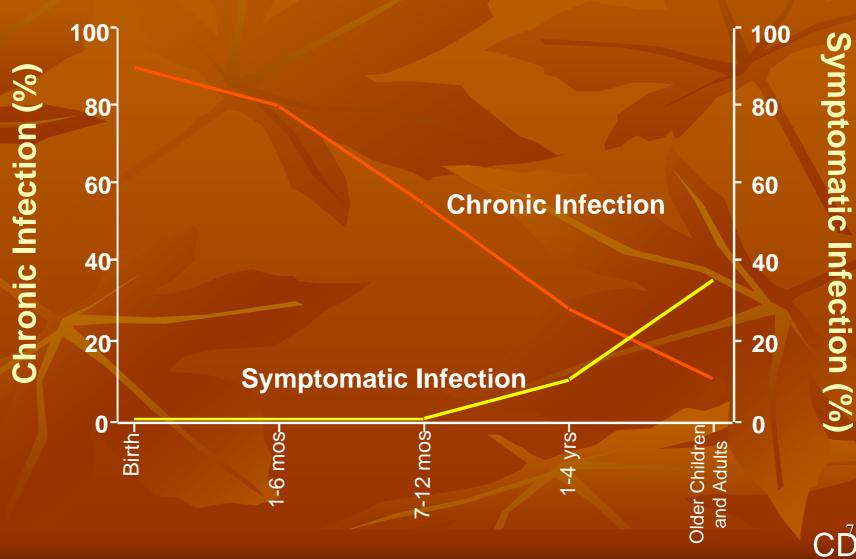
- Hepatitis B is a major public health problem in the developing countries of Africa and Asia (prevalence > 8%)
- 2 billion have markers of current or past infection
- 350 million have chronic infection
  - 15-25% will die from chronic liver disease (liver cancer and cirrhosis) at least 1 million deaths per year
  - Young children who become infected with HBV are the most likely to develop chronic infection
  - 25% mortality in perinatal acquired disease
  - Hepatitis B-associated hepatocellular carcinoma is probably the most common tumour affecting males in sub-Saharan Africa

#### Geographic Distribution of Chronic HBV Infection



- The vaccine will not cure chronic hepatitis, but it is 95% effective in preventing chronic infections from developing, and is the first vaccine against a major human cancer.
- In 1991, the WHO called for all children to receive the hepatitis B vaccine.
- Children in the poorest countries, who need the vaccine the most, have not been receiving it because of many reasons.
- Nowadays the vaccine is available in most african countries, and children are getting vaccinated, from six weeks after birth.

### **Outcome of Hepatitis B Virus Infection** by Age at Infection



#### **Hepatitis B – Clinical Features**

Incubation period:

Average 60-90 days Range 45-180 days

Clinical illness (jaundice): <5 yrs, <10% >5 yrs, 30%-50%

Acute case-fatality rate:

0.5%-1%

Chronic infection:

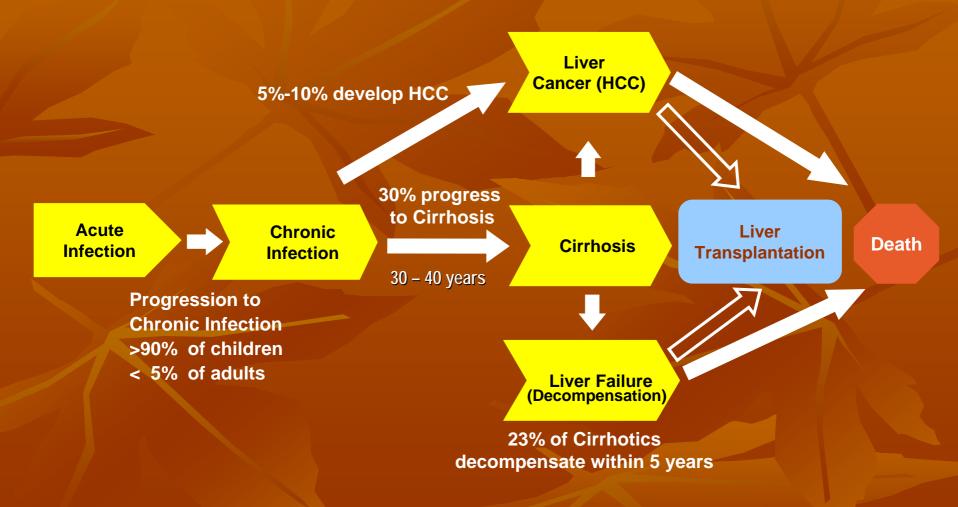
<5 yrs, 30%-90% >5 yrs, 2%-10%

 Premature mortality from chronic liver disease:

15%-25%

Pregnancy is well tolerated by women who are chronic carriers of hepatitis B

## Hepatitis B Disease Progression



Torresi, J, Locarnini, S. Gastroenterology. 2000. Fattovich, G, Giustina, G, Schalm, SW, et al. Hepatology. 1995.; Moyer, LA, Mast, EE. Am J Prev Med. 1994.

#### Vertical transmission of HBV

- ♣ Rate of transmission: HBeAg-positive ~85% HBeAg-negative ~10%
- ♣ Transmission at birth is more likely if the mother is: HBeAg positive B or has high circulating levels of HBV-DNA
- ♣ The placenta forms an excellent barrier against transmission of this large virus (DNA) and intrauterine infection is rare

Thus vertical transmission is effective during delivery

#### Prevention of vertical transmission

- ♣ Active (vaccine) and passive (HBIG) immunisation interrupts transmission in over 90%
- ♣ What about Lamivudine during the last trimester of pregnancy?
- ♣ The best protocol seems to be:
  - ◆ HBV vaccine at birth and every 4 weeks (3 doses)
  - ◆ HBIG at birth and 4 weeks later

## **Objectives**

- To provide comprehensive and reliable information on available data on global prevalence of hepatitis B in pregnant women in Africa
- To assess the vertical transmission rate of HBV to newborns

### Methods of review

- Electronic : Pubmed, WHO regional databases
- Manual search of references from original articles
- Keywords:
  - **♣** Hepatitis B AND pregnancy AND Africa
- Inclusion criteria :
  - **♠** All studies done in Africa emphasizing on prevalence and vertical transmission
- Exclusion criteria : Case report, brief communications
- Total of 144 articles were retrieved, 10 were eligible for prevalence, and 4 for vertical transmission.

## Prevalence rates of HBV in pregnant women

| S.<br>N° | Author<br>Year | Country      | Study<br>design | Setting<br>sampling<br>frame | Sample<br>size | Prevalence<br>HBsAg | Prev<br>HBeAg |
|----------|----------------|--------------|-----------------|------------------------------|----------------|---------------------|---------------|
| 1        | Madzime 1997   | Zimbabwe     | C S             | Hospital                     | 984            | 25%                 | 3.3%          |
| 2        | Nacro 2000     | Burkina Faso | C S             | Hospital                     | 917            | 10.7%               | 18.2%         |
| 3        | Rouet 2004     | Ivory Coast  | C S             | Hospital                     | 1002           | 9%                  | -             |
| 4        | Sidibe 2001    | Mali         | C S             | Hospital                     | 829            | 15.5%               | -             |
| 5        | Ahmed 1995     | Malawi       | C S             | Hospital                     | 253            | 13%                 | -             |
| 6        | Oshitani 1995  | Zambia       | C S             | Hospital                     | 2098           | 6.5%                | 16.1%         |
| 7        | Itoua 1995     | Congo brazza | C S             | Hospital                     | 292            | 6.5%                | 2.7%          |
| 8        | Ndumbe 1992    | Cameroon     | C S             | Hospital                     | 1014           | 25%                 | 5.2%          |
| 9        | Acquaye 1994   | Ghana        | C S             | Hospital                     | 692            | 6.4%                | 15%           |
| 10       | Marinier 1985  | Senegal      | C S             | Hospital                     | 1442           | 9.8%                | 19.8%         |

## Perinatal transmission rates

|   | S.<br>N° | Author<br>Year    | Country         | Study design | Setting sampling frame | Sample size | Prevalence<br>HBsAg | M-C<br>trans.<br>rate |
|---|----------|-------------------|-----------------|--------------|------------------------|-------------|---------------------|-----------------------|
|   | 1        | Roingeard<br>1993 | Senegal         | Cohort       | Hospital               | 152         | 13.8%               | 7%                    |
|   | 2        | Kew 1975          | South<br>Africa | Cohort       | Hospital               | 630         | 0.16%               | 12.5%                 |
| 7 | 3        | Badawy<br>2000    | Egypt           | Cohort       | Hospital               | 352         | 15.3%               | 51.8%                 |
|   | 4        | Menendez<br>1999  | Tanzania        | Cohort       | Hospital               | 980         | 6.3%                | 8%                    |

#### Results

Total number of pregnant women tested: 9523

Prevalence of HBsAg: 6.5 to 25%

Prevalence of HBeAg: 2.7 to 19.8% of HBsAg pos.

Vertical transmission: 7 and 57.8% of HBsAg pos.

#### **Discussion**

- All the studies dealing with prevalence and vertical transmission rate are from Africa
- We could not have any study from Africa, talking about care for newborns from infected women
- Prevalence of HBsAg between 6.5 and 25%
- Proportion of infections acquired perinatally in Africa varies between 7 to 51.8%, probably because of low prevalence of HBeAg or low circulating levels of HBV-DNA

#### **Conclusions**

Africa is an hyperendemic region for HBV

• The prevalence within pregnant women is almost the same as in the general population

• Low proportion of chronic infections acquired perinatally in Africa

#### Recommendations

- Systematic screening for hepatitis B during antenatal care (from 28 weeks of pregnancy)
- Early passive/active immunisation of babies born from all HBsAg-positive mothers is advocated. For that, HBIG should be available
- Of course, the national programme of vaccination should be continued, trying to reach all the children.



Merci pour votre aimable attention Thank you for lending me your ears



