

# Hepatitis B Virus Sero-Prevalence Among Pregnant Females in Saudi Arabia

Mohammed A. Alrowaily, Mostafa A. Abolfotouh<sup>1</sup>, Mazen S. Ferwanah

King Fahad National Guard Hospital, <sup>1</sup>Biobanking Section, Research Center, King Abdulaziz Medical City, NGH, Riyadh, Saudi Arabia

**Address:**

Dr. Mohammed A. Alrowaily, King Fahad National Guard Hospital, NGH, Riyadh, Saudi Arabia.  
E-mail: rowailym1@ngha.med.sa

## ABSTRACT

**Background/Aim:** Since selective screening for Hepatitis B virus (HBV) in pregnant women has failed to identify a high proportion of HBV-infected mothers, pre-natal HBsAg testing of all pregnant women is now recommended. We aimed to determine the prevalence of HBV infection among pregnant women at the ante-natal clinic of a tertiary care center in Saudi Arabia and to identify the target group for postpartum immunization. **Materials and Methods:** A total of 755 pregnant females who attended the antenatal clinic from June 2005 to June 2006 for the first time - before 38 weeks of gestation - constituted the target of the present study. Blood samples 30-39 were drawn from all subjects and sera were tested for HBV serologic markers including Hepatitis B surface antigen, anti-HBs, and anti-HBc using ELISA technique (third generation). **Results:** The overall prevalence of sero-positive HBsAg among pregnant women was 1.6%. As age increased, the prevalence of sero-positive HBsAg significantly increased ( $\chi^2 = 116.43$ ,  $P < 0.001$ ), 30-39 were women aged  $\geq 40$  were five times more likely to be positive for HBsAg as compared to those  $< 30$  years (OR = 4.78). On the other hand, women aged 40 and over were five times more likely to be susceptible to infection with hepatitis as compared to young women aged  $< 20$  (OR = 5.15). Women susceptible to HBV infection constituted about 80% of all pregnant females. **Conclusion:** These findings reflect that the full impact of the Hepatitis B vaccination program that was conducted in 1989 for all Saudi children has not yet reached all pregnant women, with the majority (79.9%) being nonimmune and thus liable to HBV infection. Postpartum HB immunization should be recommended in such cases.

**Key Words:** Antenatal screening, Hepatitis B, postpartum Hepatitis B vaccine

Received 29.09.2007, Accepted: 27.01.2008

The Saudi Journal of Gastroenterology 2008 14(2): 70-2

Hepatitis B virus (HBV) infection is a major public health problem in the Middle East. The majority of the countries in the region have intermediate (2 to  $< 5\%$ ) or high ( $> 5\%$ ) endemicity of HBV infection.<sup>[1]</sup> Studies in Saudi Arabia showed prevalence of Hepatitis B surface antigen (HBsAg) that ranges from 7.4 to 17% denoting high endemicity.<sup>[2]</sup>

The HBV is a major cause of chronic hepatitis, cirrhosis, and hepatocellular carcinoma. It ranks as an important pathogen throughout the world.<sup>[3]</sup> Transmission occurs mainly from a mother to child at time of parturition, as well as person-to-person (horizontal) transmission among children  $< 5$  years of age.<sup>[4]</sup> Even when not infected during the perinatal period, children of HBV-infected mothers remain at a high risk of acquiring HBV infection by horizontal transmission during the first 5 years of life.<sup>[5]</sup>

Since selective screening of pregnant women for HBV has failed to identify a high proportion of HBV-infected mothers,<sup>[6]</sup> pre-natal HBsAg testing of all pregnant women is

now recommended.<sup>[7]</sup> Universal HBsAg screening of pregnant women to prevent perinatal HBV infection has been shown to be cost saving.<sup>[8]</sup> Previous studies showed that the prevalence of HBsAg among Saudi pregnant females ranged from 3.9 to 12.7%.<sup>[9-11]</sup>

A study conducted to investigate the acceptance and efficacy of Hepatitis B immunization in United States women during the postpartum period revealed its feasibility and effectiveness.<sup>[12]</sup> Thus, the aim of the present study was to determine the prevalence of HBV infection among pregnant women at the ante-natal clinic of a tertiary care center in Saudi Arabia and to identify the target group for postpartum immunization.

## MATERIALS AND METHODS

At the antenatal clinic of King Fahad Hospital at Riyadh National Guard Health Affairs, every pregnant woman was routinely tested in each pregnancy. Serology screening for

HBsAg, Hepatitis B core antigen (HBcAb), and Hepatitis B antibody (HBsAb) is requested in every first booking visit. All pregnant females who attended the antenatal clinic from June 2005 to June 2006 for the first time - before 38 weeks of gestation - constituted the subjects of the present study.

Age and parity status were recorded. A total of 755 women were screened for HBsAg, HBsAb, and HBcAb. Blood samples were drawn from all subjects and sera were tested for HBV serologic markers including HBsAg, and HBsAb, and HBcAb were detected by using chemiluminescent microparticle immunoassay using commercial kits from Abbott Laboratories Diagnostics Division, Abbott Park, USA (ABBOTT).

SPSS software program version 10 was used for all statistical analyses.

## RESULTS

Table 1 shows an overall prevalence of sero-positive HBsAg of 1.6%. However, as age increased, the prevalence of sero-positive HBsAg significantly increased ( $\chi^2 = 116.43$ ,  $P < 0.001$ ). Sero-positivity among those aged 40 years and over was seven times as it was among people  $< 30$  (OR = 7.07). Meanwhile, people aged 30 to  $< 40$  are five times more likely to be positive for HBsAg as compared to those  $< 30$  years (OR = 4.78).

On the other hand, the number of sero-negative cases to Hepatitis B was significantly higher among the elderly compared with the youth. It increased from 48% among those below 20 years to 82.6% among those aged 40 years and more ( $\chi^2 = 59.34$ ,  $P < 0.001$ ). People 30-39 were women aged  $\geq 40$  were five times more likely to be susceptible to infection with hepatitis as compared to young people aged  $< 20$  (OR = 5.15). Moreover, those aged 20-40 are about 4.5 times more vulnerable to hepatitis as compared to their counterparts aged  $< 20$  (OR = 4.5).

## DISCUSSION

The present study shows that the sero-prevalence of HBsAg was 1.6%, which is comparable to the  $< 2.6\%$  reported by Al-Mazrou and colleagues<sup>[13]</sup> and far less than that reported by Al-Shamahy in the study conducted in Yemen<sup>[14]</sup> and by El-Hazmi.<sup>[15]</sup> It also shows that 20.1% had HBsAb titer  $> 10$  (immunity). This finding was significantly higher among mothers below 20 years of age (52%) who perhaps were exposed to the mass HBV vaccination that took place in 1989 for all Saudi children.<sup>[16]</sup>

The study also shows that 79.9% of the pregnant females have a nonimmune status making them liable to HBV infection. This suggests that the full impact of the Hepatitis B vaccination program has not yet reached all women during their maternity period, as also reported by a previous study.<sup>[13]</sup>

Jurema *et al.*<sup>[12]</sup> have previously shown that Hepatitis B immunization in the postpartum period is feasible and effective. The availability of a safe and effective hepatitis vaccine<sup>[16]</sup> encourages us to accelerate viral elimination, and additional intervention such as Hepatitis B immunization in postpartum women can thereon be undertaken. Thus, Hepatitis B immunization can be recommended, giving the first dose immediately on the first postpartum day before the mother gets discharged from the hospital; and the second dose to coincide with her child's first vaccination dose at the age of 2 months, and a third dose to be given to the mother when her child gets vaccinated at the age 6 months. However, further studies to assess the feasibility as well as effectiveness of such a program are necessary.

## ACKNOWLEDGEMENT

The authors would like to extend their gratitude and thanks to the staff members of King Abdullah International Medical Research

**Table 1: Seroprevalence of HBsAg among pregnant women by age**

Age group	HBsAg		OR	Anti-HBs		OR
	Negative	Positive		HBs $> 10$	HBs $< 10$	
$< 20$ years $n = 25$	25 100%	- -	1.0 <sup>@</sup>	13 52%	12 48%	1.0 <sup>@</sup>
20–29 years $n = 213$	212 99.9%	1 0.5%	4.78	41 19.2%	172 80.8%	4.54
30–39 years $n = 455$	446 98%	9 2%	7.07	87 19.1%	368 80.9%	4.58
$> 40$ years $n = 69$	67 97.1%	2 2.9%		12 17.4%	57 82.6%	5.15
Total $n = 762$	750 98.4%	12 1.6%		153 20.1%	609 79.9%	
$\chi^2$	116.43, $P < 0.001$			59.34, $P < 0.001$		

<sup>@</sup> - Reference category, OR - Odds ratio.

Center ( KAIMRC), King Abdulaziz Medical City (KAMC) for their support till the submission of manuscript for publication.

## REFERENCES

1. Qirbi N, Hall AJ. Epidemiology of hepatitis B virus infection in the Middle East. *East Mediterr Health J* 2001;7:1034-45.
2. Toukan A. Control of hepatitis B in the Middle East. *In: Rizzetto M, editor. Proceedings of IX Triennial International Symposium on Viral Hepatitis and Liver Disease. Edizioni Minerva Medica: Turin; 1997. p. 678-9.*
3. Ocamo P, Opio CK, Lee WM. Hepatitis B virus infection: Current status. *Am J Med* 2005;118:1413.
4. Kumar ML, Dawson NV, McCullough AJ, Radivoyevitch M, King KC, Hertz R, *et al.* Should all pregnant women be screened for hepatitis B? *Ann Intern Med* 1987;107:273-7
5. Beasley RP, Hwang LY. Postnatal infectivity of hepatitis B surface antigen-carrier mothers. *J Infect Dis* 1983;147:185-90.
6. Jonas MM, Schill ER, O'Sullivan MJ, de Medina M, Reddy KR, Jeffers LJ, *et al.* Failure of the centers for disease control criteria to identify hepatitis B Infection in a large municipal obstetrical population. *Ann Intern Med* 1987;107:335-7.
7. CDC. Protection against viral hepatitis: Recommendations of the immunization practices advisory committee (ACIP). *MMWR Recomm Rep* 1990;39:1-26.
8. Arevalo JA, Washington E. Cost-effectiveness of prenatal screening and immunization for hepatitis B virus. *JAMA* 1988;259:365-9.
9. Arya SC. Hepatitis B virus in Gizan, Saudi Arabia. *J Med Virol* 1985;17:267-74.
10. Ramia S, Hossain A, Bakir TM, Waller DK, Vivian PA. Prevalence and subtype of hepatitis B surface antigen (HbsAg) in the Saudi population. *Trop Geogr Med* 1986;38:63-9.
11. Ramia S, Abdul-Jabbar F, Bakir TM, Hossain A. Vertical transmission of hepatitis B surface antigen in Saudi Arabia. *Ann Trop Pediatr* 1984;4:213-6.
12. Jurema MW, Polaneczky M, Ledger WJ. Hepatitis B immunization in postpartum women. *Am J Obstet Gynecol* 2001;185:355-8.
13. Al-Mazrou YY, Al-Jeffri M, Khalili MK, Al Ghamdi YS, Mishkhas A, Bakhsh M, *et al.* Screening of pregnant Saudi women for hepatitis B surface antigen. *Ann Saudi Med* 2004;24:265-9.
14. Al-Shamahy H. Prevalence of hepatitis B surface antigen and risk factors of HBV infection in a sample of health mothers and their infants in Sana'a, Yemen. *Ann Saudi Med* 2000;20:5-6.
15. El-Hazmi MA. Hepatitis B virus in Saudi Arabia. *J Trop Med Hyg* 1989;92:65.
16. Al Faleh FZ, Ayoola EA, Arif M, Ramia S, Al-Rashed R, Al-Jeffry M, *et al.* Seroepidemiology of hepatitis B virus infection in Saudi Arabian children: A baseline survey for mass vaccination against hepatitis B. *J Infect* 1992;24:197-206.

**Source of Support:** Nil, **Conflict of Interest:** None declared.

## Author Help: Sending a revised article

- 1) Include the referees' remarks and point to point clarification to those remarks at the beginning in the revised article file itself. In addition, mark the changes as underlined or coloured text in the article. Please include in a single file
  - a. referees' comments
  - b. point to point clarifications on the comments
  - c. revised article with text highlighting the changes done
- 2) Include the original comments of the reviewers/editor with point to point reply at the beginning of the article in the 'Article File'. To ensure that the reviewer can assess the revised paper in timely fashion, please reply to the comments of the referees/editors in the following manner.
  - There is no data on follow-up of these patients.  
**Authors' Reply:** The follow up of patients have been included in the results section [Page 3, para 2]
  - Authors should highlight the relation of complication to duration of diabetes.  
**Authors' Reply:** The complications as seen in our study group has been included in the results section [Page 4, Table]