

Identification of Hepatitis A Virus Antibody among Bangladeshi Children and their Correlation with Socioeconomic Condition

S Mahmud¹, A S M B Karim², J Alam³, S K Saha⁴, S S Ahmed⁵, M M Z Islam⁶,
N K Sarker⁷, A S Munshi⁸, S Sarker⁹, M Afroz¹⁰, F Tasneem¹¹

Abstract

Background : Hepatitis A virus infection is endemic in many developing countries including Bangladesh. Children infected with Hepatitis-A virus typically have asymptomatic disease.

Objective : To observe the anti-HAV positivity in children & to determine any relation between anti-HAV positivity & socioeconomic condition.

Materials & Methods : A cross sectional observational study was conducted at Bangabandhu Sheikh Mujib Medical University (BSMMU) Hospital & Dhaka Shishu (Children) Hospital (DSH) from July 2008 to June 2009. Serum samples from 254 children aged between 1-15 years were tested for antibody (IgM & IgG) against hepatitis A virus (HAV) and detailed history was taken to find out the associated risk factors like residence, family income, source of drinking water, sanitation facility etc.

Results : Hepatitis A virus antibody was positive in 141 (55.5%) of 254 children. Among all children anti-HAV positivity was significantly higher ($p=0.004$) in lower income group (64.8%) than higher income group (47.0%). Anti-HAV positivity was significantly higher (93.8%) in non-sanitary latrine users & in those who used unsafe drinking water. ($p=0.006$)

Conclusions : Majority of children were found sero-positive against HAV by 15 year of age. Anti-HAV positivity was significantly higher in lower socioeconomic group.

Key words : Anti-hepatitis A virus (HAV), HAV seroprevalence

Introduction

Hepatitis A is an enterically transmitted viral disease world wide distributed.¹ Acute viral hepatitis caused by HAV is an acute, self-limiting infection.² Hepatitis A virus infection is very common in early childhood and most of the infections are asymptomatic or mildly symptomatic.³ The incidence of HAV is higher in developing countries, the true incidence of hepatitis A is often underestimated because of under-reporting as a result of its widely asymptomatic and milder forms of infection.

Three epidemiologic patterns of endemicity (low, intermediate, and high), are seen worldwide.⁴ The countries with low endemicity include Japan, Singapore, Hongkong and Taiwan whereas those with moderate endemicity include Thailand, Malaysia and Sri Lanka. Countries with high endemicity for HAV infections include India, China, Nepal, Bangladesh, Pakistan, Myanmar and Philippines.⁵ In many

developing countries of Africa, Asia and Latin America, most infections occur by 5 years of age where seroprevalence approach 90-100% by 10-15 years of age.¹ In Africa, Hendricks et al.⁶ showed anti-HAV positivity of >90% among the 5-10 year age group among lower class black children.

In developing countries, low economic status, high crowding and inadequate water treatment contribute to a high endemicity pattern; more than 90% of the population has acquired natural immunity before 10 years of age, and often shows asymptomatic forms. In many developing countries like India, Pakistan, Nepal several sero-prevalence studies have shown high rates of sero-positivity among children with subclinical infection.⁷ The present study was designed to identify HAV antibody (IgG & IgM) among children of different age group attending two tertiary care hospitals of Bangladesh and to observe relation between anti-HAV positivity & socioeconomic condition.

Materials & Methods

A cross sectional observational study was conducted from July 2008 to June 2009 at blood collection centers of Bangabandhu Sheikh Mujib Medical University (BSMMU) Hospital & Dhaka Shishu (Children) Hospital (DSH). A total of 254 children aged 1-15 years no previous history of jaundice or Hepatitis A vaccination were included in this study. The sample size was determined by the prevalence rates of neighboring countries with a similar socioeconomic condition (e.g., India & Pakistan) as there are no previous data on HAV prevalence particularly children of Bangladesh. With prior written consent, clinical history and relevant data like age, sex, date of collection, residence, monthly income of individual family, source of drinking water, food habit & sanitation facility were recorded and 2 ml of blood was collected from the study subjects.

A preformed semi structured data collecting form was used as a data collection instrument. Data were collected by researcher and analyzed by Statistical Package for social Science (SPSS) version 11.5 program. Data analysed by Chi-square (χ^2) test.

¹Dr. Salahuddin Mahmud
Assistant Professor
Dhaka Shishu Hospital, Dhaka

²Prof. A.S.M. Bazlul Karim
Paediatric Gastroenterology &
Nutrition, BSMMU

³Dr. Jahangir Alam

⁴Prof. S.K. Saha
Dept. of Microbiology,
Bangladesh Institute of Child
Health, Dhaka Shishu Hospital

⁵Dr. Syed Shafi Ahmed

⁶Dr. M.M. Ziaul Islam

⁷Dr. N.K. Sarker

⁸Dr. A.S. Munshi

⁹Dr. Shaoli Sarker

¹⁰Dr. Mahenaz Afroz
Assistant Professor of Gynae,
NICRH

¹¹Dr. Farhana Tasneem
MD (Paediatrics)-Final part
student, BICH

^{3, 5} Professor
Bangladesh Institute of Child
Health, Dhaka Shishu Hospital

^{6, 7, 8, 9} Assistant Professor
Bangladesh Institute of Child
Health, Dhaka Shishu Hospital

Correspondence

Dr. Salahuddin Mahmud
Assistant Professor, Dhaka
Shishu (Children) Hospital, Dhaka
Email: drsmbablu@gmail.com

Results

Hepatitis A virus antibody (total) was found positive in 141 (55.5%) of 254 children

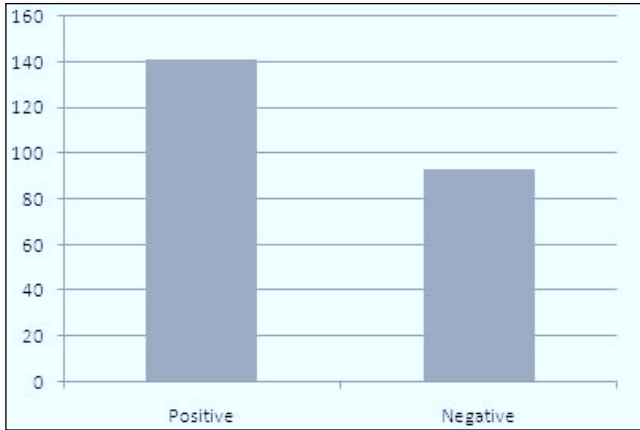


Fig: 1 Anti-HAV positivity among all children

Table I : Anti-HAV positivity with age

Age (yrs)	Total no	HAV antibody		χ^2	p-value
		Positive No (%)	Negative No (%)		
1 – 5	172	77 (44.7)	95 (55.3)	11.397	0.001
5 – 10	55	39 (70.9)	16 (29.1)	4.970	0.026
10 – 15	27	25 (92.6)	2 (7.4)		

Anti-HAV of 1-5 year age group was found to be 44.7%, it gradually increased to 70.9% in 5-10 year age group and finally to 92.6% in 10-15 year age group. Anti-HAV positivity of 5-10 year age group was significantly higher than that of 1-5 year age group (p=0.001) and antibody positivity of 10-15 year age group was significantly higher than that of 5-10 year age group (p=0.026). (Table I).

Table II : Anti-HAV status between urban & rural children

Residence	Total no	HAV antibody		χ^2	p-value
		Positive No (%)	Negative No (%)		
Urban	161	90 (55.9)	71 (44.1)	0.027	0.870
Rural	93	51 (54.8)	42 (45.2)		

Ninety (55.9%) children who live in urban area were found anti-HAV positive. On the other hand 51 (54.8%) children from rural area were found anti-HAV positive. There is no significant difference between urban and rural children regarding HAV antibody positivity (p=0.870). (Table 2)

Table III : Socioeconomic status of children with anti-HAV positivity

Monthly family income (Tk)	Total no	HAV antibody		χ^2	p-value
		Positive No (%)	Negative No (%)		
Lower class (<6000)	122	79 (64.8)	43 (35.2)	8.120	0.004
Higher class (>6000)	132	62 (47.0)	70 (53.0)		

A significantly higher number (64.8%) of children with positive HAV antibody came from lower socioeconomic class compared to that of children from higher class (47%). (p=0.004) (Table 3)

Table IV : Antibody status in children living in different housing conditions

House	Total no	HAV antibody		χ^2	p-value
		Positive No (%)	Negative No (%)		
Pucka	161	87 (54.0)	74 (46.0)	0.387	0.534
Semi-pucka & kacha	93	54 (58.0)	39 (42.0)		

Eighty seven (54.0%) children who lived in pukka house were found anti-HAV positive. On the other hand 54 (58.0%) children who lived in semi-pucka & kacha house were found anti-HAV positive. Children who were living in the semi-pucka & kacha house were found to be anti-HAV positive more than that of children living in pukka house but result is not statistically significant (p=0.534). (Table 4).

Table V : Antibody status of children with their sanitation facility

Sanitation facilities	Total no	HAV antibody		χ^2	p-value
		Positive No (%)	Negative No (%)		
Sanitary latrine users	238	126 (53.0)	112 (47.0)	10.110	0.001
Non-sanitary latrine users	16	15 (93.8)	1 (6.2)	10.110	0.001

Anti-HAV was found significantly (p=0.001) higher among non-sanitary latrine users (93.8%) than sanitary latrine users (53.0%). (Table 5)

Table VI : Antibody status of children with their source of drinking water

Source of drinking water	Total no	HAV antibody		χ^2	p-value
		Positive No (%)	Negative No (%)		
Safe water users	214	111 (51.6)	104 (48.4)	5.354	0.006
Unsafe water	40	30 (76.9)	9 (23.1)		

* Safe water (Boiled WASA, tube well & filter water)

* Unsafe water (Unboiled WASA & ponds water)

Anti-HAV was found more (76.9%) among unsafe drinking water user group than those of safe water users (51.6%) and this difference is statistically significant (p=0.006). (Table 6)

Discussion

In the present study among 254 children the anti-HAV positive was 55.5%. (141). Children at the age group of 1-5 years were found 44.7% positive Anti-HAV. It was found that sero-positivity increased with age, significant age groups were 5-10 year 70.9% ($p=0.001$) and 10-15 year 92.6% ($p=0.026$). Similar results were also observed in other studies in Bangladesh by Ahmed et al.⁸ A study by Kamath et al.⁹ reported anti-HAV positivity of 61.6% in 5-10 year age group and 97% in 11-15 year age group in Chennai, India. Agboatwalla et al.¹⁰ & Sawayama et al.¹¹ also reported similar results from Pakistan (94.1% seropositive by the age of 5 years) and Nepal (91.1% seropositive) respectively. Anti-HAV positivity was found 94.1% in 1-5 year age group at Rawalpindi and 99% in two rural villages in Nepal.

In the present study no significant difference was observed between urban (56%) and rural (55%) population regarding anti-HAV positivity. Similar result was also reported by Raharimanga et al.¹² from Madagascar, Africa where no difference was observed between urban and rural population regarding anti-HAV positivity. But opposite picture was found b Ahmed et al.⁸ showed that seropositivity of anti-HAV of rural subjects (82%) were significantly higher than those of urban (73%) population.

Significantly ($p<0.05$) higher proportion of children with positive HAV antibody came from lower income group (64.8%) compared to that of children with higher income group (47%) in urban population. Saha et al.¹³ reported that seroprevalence of anti-HAV was lower (49.8%) in higher income group than that of lower income group (96.5%). Ahmed et al.⁸ showed that anti-HAV was significantly higher in the lower income group (79.2%) compared to that with higher income group (68.8%). Socio-economic condition has some effect on anti-HAV positivity. Children living in the semi-pukka & kacha house were found to be anti-HAV positive in 58.0% cases which was nearly similar to children living in pukka house 54% ($p>0.05$). Ahmed et al.⁸ also reported the similar results. Overcrowding, poor hygienic and poor living conditions in kacha house may be the reasons of these differences.

In the current study, antibody was found significantly higher in non sanitary latrine users 93.7% than that of sanitary latrine users 53% ($p=0.001$). Ahmed et al.⁸ reported that anti-HAV positivity depend on sanitation facility. Person to person transmission of HAV generally occurs by fecal-oral route. Common source outbreaks can occur from faecally contaminated food and water. General hygiene, especially in relation to sanitation, water supply and food preparation reflects the living standards and has major influence on HAV endemicity. For the same reason, anti-HAV positivity was found higher among unsafe (unboiled, filter & ponds) water users (75%) than that of safe (boiled & tube well) water users (51.4%). Occurrence of anti HAV antibody was significantly lower among users of tube well water than those who used unboiled supply water or common tap water.

Conclusion

In the studied children anti-HAV positivity was more than 45% after 5 years of age and finally increased to more than 90% after 10 years of age. So, high proportion of children in the present study acquired HAV antibody early childhood and anti HAV positivity increased with increase in age. There is strong relation between anti-HAV positivity & lower socioeconomic condition. Anti-HAV positivity was significantly higher in low income group, non-sanitary latrine users and in those who use unsafe drinking water.

Acknowledgement

Prof. C. A. Kawser, PhD, Chairman of Pediatrics, BSMMU.

Prof. Samir K. Saha, PhD, Head, Dept. of Microbiology, BICH, DSH.

References

1. M. Z. Amina L. N. Siddiqueb, M. A. Satterc* and K. K. Biswasd; Increasing incidence of hepatitis A in Bangladesh, Bangladesh J. Sci. Ind. Res. 47(3), 309-312, 2012, Available online at www.banglajol.info
2. Feinstone SM, Gust ID. Hepatitis A virus. In: Richman DD, Whitley RJ, Hayden FG, editors. *Clinical Virology*. 2nd ed. ASM press, Washington D.C; 2002. P 1019-32
3. Arora NK & Mathur P. 'Epidemiological transition of hepatitis A in India: Issues for vaccination in developing countries'. *Indian Journal of Medical Research* 2008; 128: 699-704
4. Jacobson KH, Koopman JS, 2004. Declining hepatitis A seroprevalence: a global review and analysis. *Epidemiol Infect* 132: 1005-1022
5. Kar P. 'Is there a change in seroepidemiology of hepatitis A infection in India?' *Indian Journal of Medical Research* 2006; 123: 727-29.
6. Hendrickx G, Herck KV, Vorsters A, Wiersma S, Shapiro C, Andrus JK et al. 'Has the time come to control hepatitis A globally? Matching prevention to the changing epidemiology'. *Journal of Viral Hepatitis* 2008; 15: 1-15
7. Chadha MS, Lole KS, Bora MH & Arankalle, VA. 'Outbreaks of hepatitis A among children in Western India'. *Transactions of the Royal Society of Tropical Medicine and Hygiene* 2009; 1088: 1-6.
8. Ahmed M, Munshi SU, Nessa A, Ullah MS, Tabassum S & Islam MN. 'High prevalence of hepatitis A virus antibody among Bangladeshi children and young adults warrants pre-immunization screening of antibody in HAV vaccination strategy'. *Indian Journal of Medical Microbiology* 2009; 27(1): 48-50.
9. Kamath SR, Sathiyasekaran M, Raja TE & Sudha. 'Profile of viral hepatitis A in Chennai'. *Indian Pediatrics* 2009; 46: 642-3
10. Agboatwalla M, Isomura S, Miyake K, Yamashita T, Morishita T & Akram DS. 'Hepatitis A, B and C seroprevalence in Pakistan'. *The Indian Journal of Pediatrics* 1994; 61: 545-9
11. Sawayama Y, Hayashi J, Ariyama I, Furusyo N, Kawasaki T, Kawasaki M et al. 'A ten year serological survey of hepatitis A, B and C viruses infections in Nepal'. *Journal of Epidemiology* 1999; 9(5): 350-54.
12. Raharimanga V, Carod JF, Ramarokoto CE, Chertien JB, Rakotomanana F, Talarmin A et al. 'Age specific seroprevalence of hepatitis A in Antananarivo (Madagascar)'. *BMC Infectious Diseases* 2008; 8(78): 1-6
13. Saha SK, Setarunnahar S, Shakur S, Hanif M, Habib MA, Dutta SK et al. 'Seroprevalence of hepatitis A infection by age group and socioeconomic status of Bangladesh'. 13th International Congress on Infectious Diseases Abstracts, Poster Presentations 2008; 16(43): 101-02