SEROPREVALENCE OF TORCH ANTIBODY IN PREGNANT WOMEN
Nabi SN, Wane SMFSA, Haider KMTS, Khan AA, Hique MM

Abstract

Introduction: Infection by Toxoplasma gondii, Rubella, Cytomegalovirus (CMV), Herpes simplex virus-1 (HSV-1) and Herpes simplex virus-2 (HSV-2) TORCH group of organisms in pregnancy are the leading causes of congenital anomalies throughout the world. Most of the TORCH infections cause mild maternal morbidity but have serious fetal consequences and treatment of maternal infection frequently has no impact on fetal outcome. Therefore, recognition of maternal disease and fetal monitoring once the disease is recognized is important.

Objectives: This cross sectional descriptive study was carried out to determine the seroprevalence of TORCH antibodies in pregnant women.

Method: Single blood sample was taken from 111 pregnant women, 46 from 1st trimester, 35 from the 2nd trimester and 30 from the 3rd trimester of pregnancy. Blood samples were collected from out patient departments of Combined Military Hospital (CMH) Dhaka and Armed Forces Institute of Pathology (AFIP) Dhaka Cantonment during the period of October 2006 to July 2007. Blood samples were kept at room temperature to allow clot formation and sera were separated by centrifugation and stored in -2 ml aliquot at -20°C till testing. All the samples were tested by Enzyme Linked Immunosorbent Assay (ELISA) method for TORCH specific immunoglobulin G (IgG) and immunoglobulin M (IgM) classes of antibodies.

Result: The over all rate of seropositivity for TORCH specific antibodies found in this study were as follows: 24.32% for Toxoplasma, 87.38% for Rubella, 96.39% for CMV, 90.90% for HSV-1 and 11.71% for HSV-2. The IgG antibodies against Toxoplasma, Rubella, CMV, HSV-1 and HSV-2 were found positive in 23.42%, 81.08%, 95.49%, 87.39% and 9.91% cases respectively and the IgM antibodies were found positive in 0.90%, 6.30%, 0.90%, 2.70% and 1.80% for Toxoplasma, Rubella, CMV, HSV-1 and HSV-2 respectively.

Conclusion: The finding of this study reveals the seroprevalence of TORCH infections in the country. This study will help to create awareness amongst patients and obstetricians about TORCH infections in pregnancy and their consequences and to take the preventive and other appropriate measures against them.

Keywords: TORCH antibody, seroprevalence, pregnant women

Introduction
TORCH, which includes Toxoplasmosis, Rubella, Cytomegalovirus (CMV), and Herpes simplex infections are some of the most common infections associated with congenital anomalies. Most of the TORCH infections cause mild maternal morbidity but have serious fetal consequences and treatment of maternal infection frequently has no impact on fetal outcome. Maternal TORCH infection during pregnancy is a threat to pregnancy because they can be transmitted to the foetus while in the womb. Therefore, recognition of maternal disease and foetal monitoring once the disease is recognized is important for all clinicians.

The consequences of these infections on feto...
depend upon the type and virulence of the infecting agent and the stage of pregnancy. If a mother is exposed to these infections during the first five months of pregnancy, serious foetal complication may occur. These include miscarriage, small for gestational age, prematurity, failure to thrive, congenital heart disease, cataract, chorioretinitis, microphthalmia, blindness, microcephaly, mental retardation, hearing impairment etc. 4,5

The social and reproductive maladjustment because of repeated pregnancy wastages, cost of treatment and morbidity caused to the TORCH group of infections a major cause of concern. The prevalence of these infections varies from one geographical area to another. Many sensitive and specific tests are available for serological diagnosis of TORCH complex. Enzyme Linked Immunosorbent Assay (ELISA) for immunoglobulin M (IgM) and immunoglobulin G (IgG) antibodies against these infections is highly sensitive and specific. 4,6

This cross sectional study was carried out to determine the seroprevalence of TORCH specific antibodies in pregnant women, in order to assess their immune status and their vulnerability to Toxoplasma, Rubella, Cytomegalovirus and Herpes simplex virus infections.

Materials and Methods

The study was carried out in the Department of Immunology and Virology at Armed Forces Institute of Pathology (AFIP), Dhaka Cantonment during the period of October 2006 to July 2007. A total of 111 pregnant women were investigated. Among them 46 were in the first trimester, 35 in the second trimester and 30 were in the third trimester of pregnancy. The study cases were randomly selected from those who reported to AFIP, different family outpatient departments and indoor cases of Combined Military Hospital (CMH) Dhaka. The cases were referred patients from different clinics and hospitals of Dhaka city. About 4-6 ml of blood was collected aseptically by venipuncture in a sterile dry test tube from each pregnant woman. The blood samples were kept at room temperature to allow clot formation and sera were separated by centrifugation and stored in 0.2 ml aliquot at -70°C until testing. All the sera were tested for TORCH specific IgG and IgM classes of antibodies by Enzyme Linked Immunosorbent Assay (ELISA) method. The tests were performed with commercially available kits and the manufacturer's instructions were followed strictly. The patients were interviewed using a predefined structured questionnaire to collect data and other relevant findings. Finally data were compiled and analyzed manually and then presented in tabular forms.

Out of 111 pregnant women tested for TORCH specific antibodies, 27(24.28%) were seropositive for Toxoplasma gondii specific antibody, 109(98.18%) were positive for Rubella antibody, 107(96.39%) were positive for CMV antibody, 98(89.99%) were positive for HSV-1 antibody and 13(11.71%) were positive for HSV-2 antibody (Table-I). Distribution of specific IgG and IgM antibodies of TORCH agents amongst pregnant women were shown in Table-II. Distribution of pregnant women as per their age group and duration of pregnancy have been shown in Table-III. The age of the youngest one was 17 years and the eldest one was 37 years.

Table III: Distribution of pregnant women as per the age group and trimester of pregnancy (n=111) 7

<table>
<thead>
<tr>
<th>Age group</th>
<th>Total No.</th>
<th>1st Trimester No. (%)</th>
<th>2nd Trimester No. (%)</th>
<th>3rd Trimester No. (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>16-24</td>
<td>40 (36.00)</td>
<td>13 (32.50)</td>
<td>18 (45.00)</td>
<td>9 (22.50)</td>
</tr>
<tr>
<td>25-35</td>
<td>39 (34.90)</td>
<td>12 (30.77)</td>
<td>18 (46.15)</td>
<td>9 (22.87)</td>
</tr>
<tr>
<td>36-45</td>
<td>32 (29.00)</td>
<td>11 (27.00)</td>
<td>11 (28.12)</td>
<td>10 (25.00)</td>
</tr>
<tr>
<td>46-50</td>
<td>10 (9.00)</td>
<td>4 (10.00)</td>
<td>3 (7.89)</td>
<td>3 (7.89)</td>
</tr>
</tbody>
</table>

When IgM antibody is present, it is invariably suggests acute/subacute infection and the presence of IgG antibody suggests past or present active infection. 4,7

In the present study, an attempt was made to assess the seroprevalence of TORCH specific antibodies in pregnant women on the basis of detection of IgG and/or IgM antibodies. The overall prevalence of seropositivity found in this study was 25.42% for toxoplasma, 87.38% for rubella, 98.99% for CMV, 90.09 for HSV-1 and 11.71% for HSV-2 (Table-I). In a similar study carried out in Bangladesh by Rosawun A and Mahamudul K in 2001 where the seropositivity for TORCH specific antibodies found were 48% for toxoplasma, 93% for rubella, 90% for CMV, 88% for HSV-1 and 90% for HSV-2. The distribution of IgM antibodies in their study were 1% for toxoplasma, 5% for rubella, 1% for CMV and 2% for HSV-1. This result well correlates with the present study except the seropositivity of toxoplasma, which was lower in the present study. Another study conducted in Bangladesh by Aif A, Motiur R, Nazimuddin A et al found the seroprevalence of toxoplasma as 20%. A study conducted in India by Turabadi D, Madhar M, Reda M, et al found the seropositivity for toxoplasma, rubella, CMV, HSV-1: 52.62%, 88.10%, 99.47% and 37.18% respectively. Another study conducted in India showed seroprevalence of toxoplasma at 50.52%. Studies in Thailand and Hong Kong showed seroprevalence of toxoplasma to be 13.5% and 9.8% respectively. 8,9

In case of rubella infection, 87.38% seropositivity was found in our study. In a study conducted in Bangladesh by Rababi N, Mondal MIA, and Kawsar NA it was found that 94.33% of pregnant
depend upon the type and virulence of the infecting agent and the stage of pregnancy. If a mother is exposed to these infections during the first five months of pregnancy, serious fetal complications may occur. These include miscarriage, small for gestational age, prematurity, failure to thrive, congenital heart disease, cataract, chorionicvillitis, microphthalmia, blinding, microcephaly, mental retardation, hearing impairment etc. **4**. The social and reproductive maladjustment because of repeated pregnancy wastages, cost of treatment and morbidity caused to women makes the TORCH group of infections a major cause of concern. The prevalence of these infections varies from one geographical area to another. Many sensitive and specific tests are available for serological diagnosis of TORCH complex. Enzyme Linked Immunosensor Assay (ELISA) for immunoglobulin M (IgM) and immunoglobulin G (IgG) antibodies against these infections is highly sensitive and specific. **5**.

This cross sectional study was carried out to determine the seroprevalence of TORCH specific antibodies in pregnant women, in order to assess their immune status and their vulnerability to Toxoplasma, Rubella, Cytomegalovirus and Herpes simplex virus infections.

**Materials and Methods**

The study was carried out in the Department of Immunology and Virology at Armed Forces Institute of Pathology (AFIP) Dhaka Cantonment during the period of October 2006 to July 2007. A total of 111 pregnant women were investigated. Among them 46 were in the first trimester, 35 in the second trimester and 30 were in the third trimester of pregnancy. The study cases were randomly selected from those who reported to AFIP, different family outpatient departments and indoor cases of Combined Military Hospital (CMH) Dhaka. The cases were referred patients from different clinics and hospitals of Dhaka city. About 4-6 ml of blood was collected aseptically by venipuncture in a sterile dry test tube from each pregnant woman. The blood samples were kept at room temperature to allow clot formation and sera were separated by centrifugation and stored in 0.2 ml aliquot at -20°C till testing. All the sera were tested for TORCH specific IgG and IgM classes of antibodies by Enzyme Linked Immunosensor Assay (ELISA) method. The tests were performed with commercially available kits and the manufacturer’s instructions were followed strictly. The patients were interviewed using a predevised structured questionnaire to collect data and other relevant findings. Finally data were compiled and analyzed manually and then presented in tabular forms.

Out of 111 pregnant women tested for TORCH specific antibodies, 27 (24.32%) were seropositive for Toxoplasma gondii specific antibody, 97 (87.56%) were positive for Rubella antibody, 100 (90.59%) were positive for CMV antibody, 100 (90.59%) were positive for HSV-1 antibody and 13 (11.71%) were positive for HSV-2 antibody (Table-I). Distribution of specific IgG and IgM antibodies of TORCH agents amongst pregnant women were shown in Table-II. Distribution of pregnant women as per their age group and duration of pregnancy have been shown in Table-III. The age of the youngest one was 17 years and the eldest one was 37 years.

**Relationship between TORCH specific antibodies in women with previous normal pregnancy outcome group and women with previous bad obstetric history group** has been shown in Table-IV. Out of 111 cases, 12 (10.81%) were primigravida and 99 (89.19%) were multigravida. Among 59 multigravida cases, 57 (57.57%) bad history of previous normal pregnancy outcome and 42 (44.22%) had previous bad obstetric history. Among the 57 women who had previous normal pregnancy outcome, 22.80%, 84.21%, 94.73%, 85.96% and 12.82% cases were seropositive for Toxoplasma, Rubella, CMV, HSV-1 and HSV-2 antibodies respectively and amongst the 42 women who had previous bad obstetric history, these rates were 26.19%, 92.85%, 97.61%, 97.23% and 11.90% respectively. The seroprevalence rates were slightly higher in women having previous bad obstetric outcome group than the normal pregnancy outcome group but the differences were not statistically significant (p>0.05).

**Discussion**

Any patient infected with the TORCH group of agents, mainly two types of antibodies are produced against the infecting organism*. These are immunoglobulin M (IgM) and immunoglobulin G (IgG) types. By the means of measuring the antibody in mother’s blood, we can identify the type of infection.

**Table III**

<table>
<thead>
<tr>
<th>Age group</th>
<th>1&lt;sup&gt;st&lt;/sup&gt; Trimester (No.)</th>
<th>2&lt;sup&gt;nd&lt;/sup&gt; Trimester (No.)</th>
<th>3&lt;sup&gt;rd&lt;/sup&gt; Trimester (No.)</th>
<th>Total (No.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1-15 yrs</td>
<td>10 (18.52)</td>
<td>20 (36.36)</td>
<td>15 (27.27)</td>
<td>45 (81.82)</td>
</tr>
<tr>
<td>&gt;15 yrs</td>
<td>15 (27.27)</td>
<td>10 (18.52)</td>
<td>15 (27.27)</td>
<td>40 (72.73)</td>
</tr>
</tbody>
</table>

*In the present study, an attempt was made to assess the seroprevalence of TORCH specific antibodies in pregnant women on the basis of detection of IgG and/or IgM antibodies.

**Table IV**

<table>
<thead>
<tr>
<th>Type of previous pregnancy outcome</th>
<th>Total No. (N=111)</th>
<th>Total No. (N=57)</th>
<th>Total No. (N=42)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal pregnancy outcome</td>
<td>57 (57.57)</td>
<td>13 (22.85)</td>
<td>44 (81.82)</td>
</tr>
<tr>
<td>Bad obstetric history</td>
<td>54 (49.73)</td>
<td>44 (80.36)</td>
<td>60 (72.73)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Hemoglobin (g/dL)</th>
<th>Total No. (N=111)</th>
<th>Total No. (N=57)</th>
<th>Total No. (N=42)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal pregnancy outcome</td>
<td>10 (18.52)</td>
<td>15 (27.27)</td>
<td>3 (7.14)</td>
</tr>
<tr>
<td>Bad obstetric history</td>
<td>15 (27.27)</td>
<td>15 (27.27)</td>
<td>15 (35.71)</td>
</tr>
</tbody>
</table>

**Table V**

<table>
<thead>
<tr>
<th>Type of previous pregnancy outcome</th>
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<th>Bad obstetric history (%)</th>
</tr>
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The overall prevalence of seropositivity found in this study was 26.32% for toxoplasma, 87.38% for rubella, 99.39% for CMV, 90.09 for HSV-1 and 11.71% for HSV-2 (Table-I). In a similar study carried out in Bangladesh by Rowshan A and Mahmud K in 2001 where the seropositivity for TORCH specific antibodies were found 48% for toxoplasma, 93% for rubella, 95% for CMV, 88% for HSV. The distribution of IgM antibodies in their study were 1% for toxoplasma, 8% for rubella, 1% for CMV and 2% for HSV-1. This result well corresponds with the present study except the seroprevalence of toxoplasma, which was lower in the present study. Another study conducted in Bangladesh by Afi A, Motive R, Nazirml F A et al found the seroprevalence of toxoplasma as 20%. A study conducted in India by Turabadi D, Mathur M, Rani S, Das AK showed 36.2% seropositivity for toxoplasma, rubella, CMV, HSV-1 and HSV-2 antibodies respectively. Another study conducted in India showed seroprevalence of toxoplasma at 50.52%. Studies in Thailand and Hong Kong showed seroprevalence of toxoplasma as 13.5% and 9.8% respectively. In case of rubella infection, 87.38% seropositivity was found in our study. A study conducted in Bangladesh by Jabeet N, Mordall MIEA, and Kawser NA I It was found that 94.33% of pregnant

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women were seropositive for rubella specific IgG antibody and 0.75% for IgM antibody. In another study carried by Nahar N in Bangladesh observed that the seroprevalence of rubella specific IgG antibody was 69.25%. Studies conducted in Pakistan and Malaysia showed the seroprevalence of rubella as 94% and 92.3% respectively. The reason for this difference in the immunity in various countries is difficult to explain, however factors such as birth rate, population density, level of health immunity, ethnicity etc may play a role.

Seroprevalence of CMV was 96.99% in the present study. Samad, Dey and Khan found 84% seropositivity in a study in Bangladesh. Studies from India and Turkey reported CMV seropositivity as 84.5% and 96.4% respectively. In this study HSV-1 antibody was found in 90.09% cases and HSV-2 antibody in 11.71% cases. In USA the seropositivity reported 90% for HSV-1 and 20-25% for HSV-2 and the IgG antibody seropositivity reported 90.90% for HSV-1 and 37.10% for HSV-2 from Saudi Arabia.

Conclusion
The present study demonstrates a high seroprevalence rate of TORCH specific antibodies in pregnant women in our country. Recognition of these infections will help the clinician to appropriately counsel mothers on preventive measures to avoid these infections and aid in counseling parents on the potential for adverse fetal outcomes when these infections are present. Early diagnosis will help in proper management of the cases. It is therefore recommended that all pregnant women should be routinely screened for TORCH agents at the first antenatal visit and the patient should be managed accordingly. Awareness amongst clinicians and patients about TORCH infections and their consequences should be developed.

References
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