Seroprevalence of cytomegalovirus antibodies amongst normal pregnant women in Nigeria

Objective: Cytomegalovirus (CMV), a ubiquitous virus belonging to the herpes family, is known to be transmitted frequently to developing fetuses in pregnancy. In an immunocompromised state like pregnancy, primary infection through blood transfusion or reactivation of a latent CMV infection can cause severe illness. The study was carried out to determine the seroprevalence of the immunoglobulin G (IgG) antibody to cytomegalovirus amongst pregnant women in correlation with previous exposure to blood transfusion.

Methods: A cross sectional study was carried out amongst 179 HIV negative pregnant women attending the antenatal clinic of Lagos State University Teaching Hospital (LASUTH), Ikeja, Nigeria. Five mL of blood was collected and stored in a plain bottle, centrifuged on the same day and the serum stored at −20°C. All samples were screened for anti-CMV IgG antibodies using the enzyme linked immunosorbent assay (ELISA). Consenting participants were instructed to fill a semi-structured questionnaire to obtain demographic and other related information. Statistical analysis of the results was done using Pearson’s chi squared test for analytical assessment.

Results: A total of 97.2% of the pregnant women recruited for this study were anti-CMV IgG positive. Out of the 179 recruited for the study 174 responded to the question on previous history of blood transfusion, 14.9% of the respondents (26 of 174) had a previous history of blood transfusion and all tested positive to the anti-CMV IgG antibody. However, past history of blood transfusion and educational level were found to be insignificant to the risk of acquiring CMV infection.

Conclusion: The seroprevalence of the CMV antibody amongst pregnant women in this environment is high in relation to findings in other developing countries. There is the need to assess anti-CMV immunoglobulin M antibodies in pregnant women, which is a determinant of active infection.

Keywords: CMV, IgG, pregnant women

Introduction

Cytomegalovirus belongs to the subfamily of herpes known as β herpes and it is found universally in various geographic locations. CMV infection is also known to be frequently transmitted to a developing fetus. This virus remains the leading cause of congenital viral infection and a significant cause of transfusion-acquired infections in patient populations. Its clinical manifestations include asymptomatic forms, severe fetal damage, and death in rare cases due to spontaneous abortion.

For most healthy people who acquire CMV infection after birth or through blood transfusion, there are few symptoms and no long term sequelae. Therefore, for the vast majority of individuals, CMV infection is innocuous. However, CMV
infection is important in pregnant women because of their immunocompromised state and risk of infection to the fetus whose immune system is not fully developed. Furthermore, 10% to 15% of the children who are asymptomatic at birth may develop late sequelae, especially hearing defects, after a period of months or years.\(^3\)

Once a person becomes infected, the virus remains alive but usually dormant within the individual’s body for life. Recurrent disease rarely occurs unless the immune system is compromised. It was reported that the risk of fetal damage is greater if the primary infection occurs during the first trimester of pregnancy.\(^4,6\) The prevalence of congenital infection ranges from 0.2% to 2.5% in different populations.\(^7,11\) with groups at greater risk including Black or Asian women, those from a low socioeconomic background, and those that are born prematurely.\(^10\)

The prevalence of congenital infection varies with the prevalence of the infection in the population.\(^14\) The seroprevalence of CMV among women of childbearing age ranges from 35% to 95% in different countries.\(^5,14-17\) It is more widespread in developing countries and in areas of lower socioeconomic conditions.\(^18\)

The rate of seropositivity of anti-CMV immunoglobulin G (IgG) enzyme linked immunosorbent assay (ELISA) antibodies of pregnant women in Turkey was reported to be 98.5%\(^19\) and 84% in Spain.\(^20\) These rates are much higher than the typical European rate but similar to the rate obtained amongst Black pregnant women.

An overall rate of 87% of anti-CMV IgG ELISA antibodies in pregnant women was reported in Singapore,\(^21\) 100% in Thailand,\(^21\) and 93% in Iranian women of childbearing age.\(^22\)

Since CMV is transmitted through blood transfusion, the strict use of blood during pregnancy and labor cannot be overemphasized as some of the pregnant women could be CMV negative.

Several studies and data support the transmission of CMV from seropositive blood donors to susceptible recipients in a variety of settings.\(^23,24\) Topin et al\(^25\) provided the first biochemical evidence for transfusion associated CMV infection. The study reported that monocyte latently infected with CMV represent the primary vector for Transfusion Associated-CMV (TA-CMV) which can be largely abrogated by transfusing at risk patients with either seronegative units or blood filtered to remove white blood cells. Lamberson et al\(^26\) also determined that a decreased incidence of transfusion associated with CMV (TA-CMV) infection occurred when only blood products that tested negative for CMV immunoglobulin M (IgM) were used.

Two hypotheses have been investigated as possible mechanisms for CMV transmission by blood. First, since CMV resides in a latent state in the infected host, it has been proposed that all CMV-seropositive blood donors can transmit the latent virus\(^27,28\) following transfusion. In this case latent virus is reactivated and may cause active CMV infection in the recipient. Latency following a primary infection may be punctuated by periodic reactivation that gives rise to recurrent infections, and in utero transmission may occur during either primary or recurrent infections.\(^29\) Yeager and colleagues published an excellent study of TA-CMV infections in newborns to prove this fact.\(^30\) CMV-specific antibody of the IgM class has been recognized as a marker of active or recent primary infection with the virus. Recent reports have shown a positive correlation between post transfusion CMV infection and the receipt of blood from CMV IgM-positive donors.\(^31\)

Such units of blood may be used for CMV negative pregnant women thus increasing the incidence of CMV in the population with its consequent embryopathy such as sensorineural hearing loss, chorioretinitis, mental retardation and fetal death. CMV embryopathy should be a major concern for public health irrespective of the percentage of babies affected. Therefore, the need to determine the seroprevalence of the CMV antibody in pregnant women cannot be overemphasized. This will underscore the significance of CMV screening in blood units for use in pregnancy and labor. This is necessary to avoid the transmission of CMV infected blood to women who are CMV seronegative.

Subjects and method

Study population

A cross sectional study was carried out using the antenatal clinic of Lagos State University Teaching Hospital (LASUTH), Ikeja, Nigeria. The clinic registers about 150 patients daily and all consenting newly registered pregnant women were recruited for the study between July 1 and August 15, 2008. However, approval was obtained from the institution’s research and ethics committee (LASUTH Health Research and Ethics Committee). Participants were instructed to fill out a structured questionnaire including demographic information and previous exposure to blood transfusion. Immunocompromised participants with a history of chronic illness, such as hepatitis, sickle cell disease, renal disorders, gestational diabetes and HIV infection were excluded.
Collection of samples
A blood sample of 5 mL was collected into a sterile, nonan-
ticoagulated bottle from each consenting participant. The
sample was centrifuged and the serum separated into a sterile
bottle and stored at −20°C.

Cytomegalovirus serology
Sera were tested for IgG CMV using the ELISA test. The
CMV-specific IgG antibodies were studied by the commercial
Dia.Pro Diagnostic Bioprobe CMV IgG (Milan, Italy)
according to the manufacturer’s instructions. All specimens
were analyzed using the enzyme immunoassay test. The
cutoff of IgG was set at 0.5 IU/mL (calibrator 2) by the kit’s manufacturer.

Samples with a concentration ≥0.5 WHO IU/mL were
considered positive for CMV IgG, while samples with a
concentration below the cut off were considered as negative
results. The controls and the calibrators passed the validation
check recommended by the manufacturer of the kit.

Statistical analysis
Statistical analysis of results was done using the statistical
package for social science (SPSS 16.0, IBM, USA). The
Pearson chi squared test and multiple comparisons
involving post-hoc analysis such as the Bonferonni correction
were used for the analytic assessment and the differences
were considered to be statistically significant when the
P-value obtained was <0.05.

Results
A total of 179 pregnant women were enrolled into the study.
About 50.8% of them were between the ages of 25–30 years
and 0.60% were under 18 years (Table 1). Almost all of them
(97.20%) had anti-CMV IgG antibodies. The majority of
those (89 of 179) who had anti-CMV IgG antibodies were
between parity 1 and 4 (Table 2). It was observed that most
(70.40%) of the patients had tertiary education, followed
by 24.60% with secondary education. Less than 2% had no
formal education (Table 3). However, levels of education
did not correlate with CMV status (P = 0.14).

The majority (86.03%) of the subjects were married and
most of them had anti-CMV IgG antibodies, 11.17% were
single and 2.7% were separated. (P = 0.11) (Table 4).

One hundred and seventy four of the participants
responded to the question on past history of transfusion while
14.94% of the 174 respondents (26 of 174) had a previous
history of blood transfusion and all tested positive to the
anti-CMV IgG antibody. Equally, 148 of the 174 (85.05%)
subjects had no past history of blood transfusion and 143 of
the 148 (96.62%) tested positive to anti-CMV IgG antibodies,
but only 5 of 148 (3.37%) tested negative to the IgG CMV
P = 0.77 (Table 5). Hence an association cannot be established
between transfusion history and anti-CMV IgG positivity.

Table 1 Age in years with CMV serostatus

<table>
<thead>
<tr>
<th>Age in years</th>
<th>anti-CMV IgG +ve</th>
<th>anti-CMV IgG –ve</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;18</td>
<td>2</td>
<td>0</td>
</tr>
<tr>
<td>19–24</td>
<td>18</td>
<td>2</td>
</tr>
<tr>
<td>25–30</td>
<td>89</td>
<td>2</td>
</tr>
<tr>
<td>31–35</td>
<td>45</td>
<td>1</td>
</tr>
<tr>
<td>36–40</td>
<td>17</td>
<td>0</td>
</tr>
<tr>
<td>41 and above</td>
<td>3</td>
<td>1</td>
</tr>
</tbody>
</table>

Note: P = 0.14.

Abbreviations: CMV, cytomegalovirus; IgG, immunoglobulin G.

Table 2 Parity with CMV seropositivity

<table>
<thead>
<tr>
<th>Parity</th>
<th>anti-CMV IgG +ve</th>
<th>anti-CMV IgG –ve</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>60</td>
<td>3</td>
</tr>
<tr>
<td>1–4</td>
<td>89</td>
<td>2</td>
</tr>
<tr>
<td>5 and above</td>
<td>25</td>
<td>0</td>
</tr>
</tbody>
</table>

Note: P = 0.42.

Abbreviations: CMV, cytomegalovirus; IgG, immunoglobulin G.

Table 3 Education with CMV seropositivity

<table>
<thead>
<tr>
<th>Education</th>
<th>anti-CMV IgG +ve</th>
<th>anti-CMV IgG –ve</th>
</tr>
</thead>
<tbody>
<tr>
<td>None</td>
<td>3</td>
<td>0</td>
</tr>
<tr>
<td>Primary</td>
<td>5</td>
<td>1</td>
</tr>
<tr>
<td>Secondary</td>
<td>42</td>
<td>2</td>
</tr>
<tr>
<td>Tertiary</td>
<td>124</td>
<td>2</td>
</tr>
</tbody>
</table>

Note: P = 0.14.

Abbreviations: CMV, cytomegalovirus; IgG, immunoglobulin G.

Discussion
The 97.20% seropositivity rate of IgG anti-CMV antibodies
observed in this study amongst pregnant women is in keep-
Table 4 Marital status with CMV seropositivity

<table>
<thead>
<tr>
<th>Marital status</th>
<th>anti-CMV IgG +ve</th>
<th>anti-CMV IgG –ve</th>
</tr>
</thead>
<tbody>
<tr>
<td>Single</td>
<td>18</td>
<td>2</td>
</tr>
<tr>
<td>Married</td>
<td>151</td>
<td>3</td>
</tr>
<tr>
<td>Separated</td>
<td>5</td>
<td>0</td>
</tr>
</tbody>
</table>

Note: P = 0.11.

Abbreviations: CMV, cytomegalovirus; IgG, immunoglobulin G.

Table 5 Relationship between blood transfusion and anti-CMV IgG seropositivity

<table>
<thead>
<tr>
<th>Past transfusion history</th>
<th>anti-CMV IgG +ve</th>
<th>anti-CMV IgG –ve</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yes</td>
<td>26</td>
<td>0</td>
</tr>
<tr>
<td>No</td>
<td>143</td>
<td>5</td>
</tr>
</tbody>
</table>

Note: P = 0.77.

Abbreviations: CMV, cytomegalovirus; IgG, immunoglobulin G.

A total of 67.2% of caesarean sections of these cases were emergency.39 The rate of blood transfusion for the various indications were as follows: malpresentation (excluding breech) 66.7%; placenta previa 59.1%; uterine rupture 55.6%; breech delivery 32%; obstructed labor 28.2%; previous baby 25%; previous cesarean section 17.0%; severe preeclampsia 11.1%; fetal distress 10.7%; and others, 10.3%. Akinola et al40 also reported in a similar study a transfusion rate of 13.6% amongst cesarean section patients in southwestern Nigeria.

Despite the reported rate of lower seroprevalence of anti-CMV IgG ELISA antibodies amongst higher socioeconomic classes and in developed nations,18 lower rates of blood transfusion of 4.9% and 5.4% were reported by Dathie et al41 and Rouse et al42 respectively. In Canada43 the overall rate of blood transfusion for postpartum hemorrhage was 0.31%, and 0.49%, 0.28%, 0.23% for emergency caesarean section, vaginal delivery, and elective caesarean section respectively.

In developing countries, the higher rate of anti-CMV IgG seroprevalence associated with higher requirements for blood in pregnant women poses a grave risk of CMV embryopathy. There is the need to assess anti-CMV IgM antibodies in pregnant women in these countries which is a marker of acute infection and more directly related to CMV embryopathy.

Previous history of blood transfusion and socioeconomic status were, however, found in this study not to be significant factors to CMV antibody positivity. This might be related to the high seroprevalence of anti-CMV IgG in the Nigerian population. Future studies in populations with similarly high seroprevalence may need higher recruitment numbers.

Conclusion

The seroprevalence of anti-CMV IgG is very high among pregnant women in Nigeria. Based on previous studies that showed a decrease in the incidence of CMV disease when blood was screened for CMV (IgM), the incidence of the disease can be decreased in Lagos if blood is screened for CMV. Previous blood transfusion exposure was insignificant in the risk of transmission.

Limitations of this study

1. Reliability of information on blood transfusion provided by patients and the possible reasons for finding small numbers of those who have been previously transfused.
2. The use of IgG ELISA assay rather than IgG and IgM in determining seroprevalence of CMV in pregnant women.

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Disclosure
The authors report no conflicts of interest in this work.

References

