Evaluation of the clinical utility of maternal alloantibody screening as a surrogate to antiglobulin crossmatch procedures in resource limited settings

Zaccheaus Awortu Jeremiah, Augustina Mordi

ABSTRACT

Aims: In resource limited settings, cross matching procedures are usually limited to the conventional antiglobulin technique. Pretransfusion screening for red cell alloantibodies are not carried out routinely. The study was aimed at evaluating the usefulness of antibody screening as a surrogate to antiglobulin crossmatch procedure. Methods: A total 250 pregnant women attending the antenatal clinic of the Braithwaite Memorial Specialist Hospital (BMSH), Port Harcourt were screened for the presence of red cell alloantibodies using DiaMed screening and panel cells (DiaCell and DiaPanels). Results: Alloantibodies detected were anti-E (1.2%), anti-K (0.8%), anti-C (0.4%) and anti-Jsb (0.4%). The overall prevalence rate of red cell allo–antibodies was 4.8%. A blind crossmatch performed using the serum of the patients on donor's cells revealed the following results, incompatible 5 (2.0%) and 254 (98.0%) compatible. Taking incompatible results as positive and compatible as negative, the performance indices of the antibody screening procedure was obtained as follows: sensitivity (41.6%), specificity (100%), PPV (100%), NPV (97.1%), efficiency (48.6%). Prevalence (4.8%) and percentage safety (41.6%). The study did not show the type and screen to reach the expected safety level of 99.0%. Its usefulness was however shown through the detection of unexpected antibodies in 4.8% of the subjects. Conclusions: We concluded that with a high specificity obtained, the detection and identification of these antibodies would help select blood in advance for patients undergoing surgery in order to reduce the incidence of haemolytic transfusion reactions.

Keywords: Antibody screening, Antiglobulin crossmatch, Pretransfusion testing, Antibody identification

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Article ID: 100001IBTIZAJ2011.

doi:10.5348/ijbti-2011-1-OA-1

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INTRODUCTION

Antibody detection and identification are fundamental to the practice of immunohaematology. Antibody identification can be a guide to the clinical significance of the antibody and provides information that aids in the selection of suitable blood for transfusion. Prenatal immunohaematologic care of
pregnant women requires the investigation of unexpected RBC antibodies in their sera during pregnancy. When RBC antibody screening is positive, it is necessary to determine specificity of the antibody, its clinical importance and the ability to cross the placenta and cause haemolytic disease of the foetus and newborn (HDFN). In some circumstances, it can be a difficult and time consuming process and thus cause a delay in patient care [1].

In Port Harcourt, as in other developing parts of the world, type and screen procedure is not routinely carried out as part of the pre-transfusion test protocol, hence the incidence of maternal red cell allo-antibodies and the prevalence of these unexpected antibodies in this locality are not known. With the high incidence rate of neonatal jaundice in Port Harcourt [2] and paucity of information on this subject, it becomes necessary for a study like this to be conducted; hence this is the first attempt ever to provide the prevalence of unexpected antibodies in this part of the world.

This study therefore aimed to: 1) provide the frequencies of allo-antibodies among pregnant women attending the Braithwaite Memorial Specialist Hospital, 2) evaluate the clinical utility of the antibody screening procedure as a surrogate to antiglobulin cross matching procedures and, 3) determine the specificity of the antibodies identified in the sera of the pregnant women.

MATERIALS AND METHODS

**Study area and population:** This study was conducted in Braithwaite Memorial Specialist Hospital (BMSH), Port Harcourt, and the capital city of Rivers State of Nigeria. The study population consisted of 250 pregnant women aged 16 – 45 years recruited form antenatal care clinic within the period of six months.

**Study Design:** A prospective cross-sectional design was used in this study. Samples were collected randomly, after obtaining a written or oral informed consent from the patients. Women from different ethnic groups were randomly recruited parameters together with sampling various blood.

**Collection and processing of samples:** Four milliliter of whole blood was drawn with syringe (5 ml) through venepuncture using the antecubital vein. The whole blood was allowed to clot in the plain bottle, centrifuged at 300g for five minutes and the serum separated into a separate plain tube with a cap. The separated serum was stored at – 80°C. Serological parameters were assessed using the achieved sample. All blood samples were collected at the Antenatal unit of the Braithwaite Memorial Specialist Hospital, Port Harcourt.

Determination of ABO and Rhesus blood groups: ABO blood grouping was done using anti-A, anti-B, and anti-AB bought from Biotec (Ipswich, UK) with standard tube agglutination technique. All blood group tests were confirmed with known test RBCs. Negative controls were included in all tests. ABO blood group tests were performed only at room temperature. Reverse grouping cells were also supplied by Biotec. Rhesus grouping was done using anti-D monoclonal reagent bought from Biotec. Rhesus controls were supplied in all tests. Tests were done in tubes and all negative results were confirmed using indirect agglutination test technique with 20% bovine albumin and anti-human globulin (AHG) tests at 37°C. After spinning for 20 sec at 100 rpm, the RBC was gently resuspended and immediately observed macroscopically and confirmed microscopically before recording the result as positive or negative.

**Antibody screening and Identification:** Antibody screening panel (3 cells) and identification panel (11 cells) from DiaMed (Switzerland) was used to screen and identify alloantibodies by tube method in low ionic strength solution, albumin and AHG phase according to the manufacturer’s instructions.

**Antiglobulin crossmatch:** Antiglobulin crossmatch using Indirect Antiglobulin Test procedure was carried out on the patients’ serum with pooled donors’ red cells as described by Morgan [3] by a different Medical Laboratory Scientists who was ignorant of the antibody screening results.

**Compatibility testing:** A negative reaction indicates compatibility of the donor blood with the recipient.

A positive reaction indicates incompatibility of the donor blood with the recipient, due to the presence of antibodies directed against antigens on the donor red cells. Further investigation to identify the antibody specificity should be performed.

**Statistics:** Results were analyzed using the SPSS version 16 for windows (Chicago, IL) Frequency table was used to determine the prevalence of allo-antibodies among the pregnant women. Performance indices such as sensitivity, specificity, positive predictive value, negative predictive value, efficiency, prevalence and safety level were calculated using standard formulae.

RESULTS

A total of two hundred and fifty (250) pregnant women attending the Braithwaite Memorial Specialist Hospital (BMSH) were screened for the presence of irregular antibodies.

The demographic characteristics of the pregnant women are shown in table 1. Majority of the study population were in the 26 – 30 years age group (43.9%) and in their second semester (60.4%). Those of the Ibo ethnic group dominated the study population (28%) followed by those of Ikwerre (23.8%) and Ijaw (18.4%). Distribution of ABO and Rh blood groups among the 250 pregnant women revealed that prevalence of group O was 48.0%, A - 41.2%, B - 7.6%, and AB was 3.2%. Rhesus D (Rh D) positive accounted for 91.4%, whereas Rh D negative was 8.6% (table 2). Table 3 shows the specificity of the antibody detected in the study population. The most frequently occurring irregular
antibody detected was anti-E in 3 (1.2%) patients followed by anti-K in 2 (0.8%) patients, then anti-C and anti-Jsb (0.4% respectively). Three of the samples had mixed field reactions while two could not be identified using the panel cell. A total of 238 (95.2%) samples gave negative antibody screen. The antiglobulin crossmatch was performed on all samples and five (2.0%) were found to be incompatible. A total of 245 (98.0%) were compatible as shown in table 4. Table 5 shows the clinical utility of antibody screening as a surrogate to antiglobulin crossmatch. Using performance indices and taking incompatible results as positive, the sensitivity of antibody screening method was found to be 41.6% while specificity was 100%. The positive predictive value (PPV) was 100% while negative predictive value (NPV) was 97.1%. Efficiency was 48.6% while the percentage safety of the

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Number</th>
<th>Percentage (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age group (yrs.)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>16–20</td>
<td>4</td>
<td>16</td>
</tr>
<tr>
<td>21–25</td>
<td>30</td>
<td>12.0</td>
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<tr>
<td>26–30</td>
<td>112</td>
<td>43.9</td>
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<tr>
<td>31–35</td>
<td>70</td>
<td>27.5</td>
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<tr>
<td>36–40</td>
<td>25</td>
<td>9.8</td>
</tr>
<tr>
<td>41–45</td>
<td>9</td>
<td>3.5</td>
</tr>
<tr>
<td>Total</td>
<td>250</td>
<td>100.00</td>
</tr>
<tr>
<td>Trimesters</td>
<td></td>
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</tr>
<tr>
<td>First</td>
<td>23</td>
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</tr>
<tr>
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<tr>
<td>Third</td>
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<tr>
<td>1</td>
<td>71</td>
<td>27.8</td>
</tr>
<tr>
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</tr>
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<td>3</td>
<td>33</td>
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<tr>
<td>0</td>
<td>50</td>
<td>20.0</td>
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<tr>
<td>Ethnic group</td>
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<td>18.4</td>
</tr>
<tr>
<td>Ogoni</td>
<td>19</td>
<td>7.5</td>
</tr>
<tr>
<td>Ekpeye</td>
<td>12</td>
<td>4.7</td>
</tr>
<tr>
<td>Ikwerre</td>
<td>58</td>
<td>23.8</td>
</tr>
<tr>
<td>Ibo</td>
<td>70</td>
<td>28.0</td>
</tr>
<tr>
<td>Yoruba</td>
<td>10</td>
<td>3.9</td>
</tr>
</tbody>
</table>

antibody screening method was 41.6%. The overall prevalence of irregular antibodies was 4.8%.

DISCUSSION

Irregular RBC antibodies found in the sera of pregnant women have been studied in many parts of the world where pre-natal immunohaematologic care is given due priority. In this study, the frequency of irregular antibodies in maternal serum was 4.8%. This appears high when compared with values from developed countries like Sweden (0.5%), Netherlands (2.7%), and lower when compared with values from developing countries where higher frequency values of 10.2% in Mexico, and 20% anti D were reported [3-6].

The most frequent and potentially significant non-anti–D antibody in our study was anti-E (1.2%) followed by anti-K (0.8%) then anti-C (0.4%) and anti-Jsb (0.4%). Bowel found an incidence of 14% in D+ pregnant women which was found to be most frequent in his study contrary to the results in this study where anti-E is most frequent [7]. Anti-E can be a naturally occurring IgM antibody, however IgG anti-E can be found in the sera of pregnant women with a history of previous transfusions and pregnancies. This immune form of anti-E is able to cause a mild to moderate HDN [8]. HDN caused by anti-C is usually mild as the C antigen has weak immunogenicity [7-11].
Table 4: Blind compatibility test results using the screened sera of patients (anti globulin cross match).

<table>
<thead>
<tr>
<th>Crossmatch results</th>
<th>Number</th>
<th>Percentage (%)</th>
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</thead>
<tbody>
<tr>
<td>Compatible</td>
<td>245</td>
<td>98.0</td>
</tr>
<tr>
<td>Incompatible</td>
<td>5</td>
<td>2.0</td>
</tr>
</tbody>
</table>

Table 5: Clinical utility of antibody screening as a surrogate to anti globulin cross match.

<table>
<thead>
<tr>
<th>Antiglobulin Crossmatch</th>
<th>Positive</th>
<th>Negative</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Incompatible (+)</td>
<td>TP (5)</td>
<td>FP (0)</td>
<td>5</td>
</tr>
<tr>
<td>Compatible (-)</td>
<td>FN (7)</td>
<td>TN (238)</td>
<td>245</td>
</tr>
<tr>
<td>Total</td>
<td>12</td>
<td>238</td>
<td>250</td>
</tr>
</tbody>
</table>

Positive AB screen includes identified, mixed field and unidentified reaction
TP = True positive, FP = False positive
FN = False Negative, TN = True negative
PPV = Positive predictive value,
NPV = Negative predictive value
Incompatible results = positive results
Compatible results = negative results
Sensitivity = TP/TP + FN = 5/12 x 100 = 41.6%
Specificity = TN/TN + FP = 238/238 x 100 = 100%
PPV = TP/TP + FP = 5/5 x 100 = 100%
NPV = TN/FN + TN = 238/245 x 100 = 97.1%
Efficiency = TP + TN/TP + FP + FN + TN x 100 = 238/245 x 100 = 48.6%
Prevalence 12/250 x 100 = 4.8%
Taking incompatible Crossmatch as 100%
Number of incompatible crossmatch = 100%
Number of incompatible crossmatch with an antibody detected 5/12 x 100% = 41.6%
Percentage safety = 41.6% (safety range 99–99.9%)

Anti-K was seen in this study with a frequency of 0.8%. The frequency of K antigen in this locality is not yet known but it is known that after the D antigen, the K antigen is the most immunogenic. HDN caused by anti-K can be severe [9]. There is evidence that anti-K can recognize K antigens expressed in the early stage of erythroid development in the fetal liver and can cause anaemia by suppressing erythropoiesis [12-14]. Jsb has been reported to be common among people of African jessant. Anti-Jsb was the least frequent of all the four specific cities and has been known to be weakly immunogenic. Anti-D was not seen in this study in contrast to 20% anti-D found among pregnant women in Saudi Arabia [4]. Jeremiah and Buseri [15] reported Rhesus antigen and phenotype frequencies in Port Harcourt as follows. D neg (5.0%), C neg (82.3%), C neg (0.2%), E negative (79.5%) and e neg (1.3%). It is therefore expected that anti-C and anti-E will occur more frequently in this locality while anti-D will be less common. This probably may explain why anti-D was not encountered in this investigation. Contrastingly, the frequency of D-negative has been reported to be approximately 15 percent among Caucasians and 20-30% in Middle East and some West African countries. It is not surprising that variable frequencies of unexpected antibodies were obtained in different region of the globe. However, a more detailed population study with larger sample size may need to be earned out using more sensitive methods to arrive at a more accurate figure.

Maternal serum is screened to make sure pregnant mother has no antibodies to react with fetal cells. Haemolytic disease of the foetus and newborn is caused by the mother’s IgG antibodies crossing the placenta and attaching to the baby’s red blood cells. It is therefore necessary to know as early as possible in the pregnancy whether HDN can be a possibility.

Determining the specificity of an unexpected alloantibody is important in prenatal testing. If the antibody specificity is known, it is possible to test donor blood for the presence of the corresponding antigen. In prenatal testing, knowledge of the specificity of the antibody helps predict the likelihood of the haemolytic disease of the newborn.

In most settings, antibody screening is done as part of pre-transfusion tests but the question as to its use as a surrogate to cross-matching procedure in emergency situation still remain not clear. In this study, a full blind crossmatch was performed in order to assess the safety level of antibody screening procedures as a surrogate to crossmatch. This study did not show the antibody screening procedure to reach the expected safety level of 99%. The safety was low (41.6%) but with a high specificity of 100%, it can be used to select blood in advance for patients undergoing surgery in order to reduce the incidence of haemolytic transfusion reaction. Apart from having a low safety level of 41.6%, it has earlier been reported that the “type and screen” porocedure has the weakness that antibodies to rare non-polymorphic antigens will not be detected (approximately 0.06% of cases), but the antibodies missed are rarely of clinical importance [16,17].

Many of the red blood cell (RBC) alloantibodies of the Rhesus system have been associated with HDFN; however, the severity of the disease is usually the greatest with anti-D [18]. Prevention of RhD HDFN became feasible in the late 1960s after pioneering research by Finn Clarks and Freda; there was a dramatic decline in Rh HDFN [19, 20]. Since then other Rh and non-Rh red cell alloantibodies have become relatively more important and are now responsible for the greater proportion of the HDFN cases. Anti-c and anti-E are the most common implicated non-D Rh antibodies in the pathogenesis of HDFN [21]. In this study, anti-E was found to be most frequently and this supports the earlier report to a large extent. Another study assessed the outcomes of anti-E in pregnancy and they concluded that a substantial proportion of infants are sufficiently affected by anti-E and suffer from clinically significant HDFN. In that study, 21% of the affected infants required exchange transfusion and 10% had severe or very severe disease and they concluded that anti-E titres and poor predictor of HDFN severity. Another study also showed that anti-E was a caused of moderately severe HDFN [7].

Most (95.2%) of the pregnant women who participated in this study showed no antibodies i.e. they
were non-immunized. Most of the immunized women (n = 12, 4.8%) developed allantibodies of the non-D Rhesus and Kell blood groups.

The study shows that immunization due to antibodies belonging to the Rhesus system is 1.6% (i.e. non-D Rhesus antibodies) of all examined Port Harcourt pregnant women. Anti-D antibodies were not encountered in this study. It is possible the use of rhesus immunoglobulin anti-D to prevent sensitization due to anti-D could have led to the decrease in the incidence of anti-D antibodies. Besides the percentage of D-negative women in our locality is quite small when compared with E-negative and C-negative (15).

Anti-C was found in 0.4% of the immunized patients. Baker et al., [22] reported one case on a group A Rh positive, C negative woman in whom anti-C was developed as a result of blood transfusion in childhood.

In summary, the number of antibodies against antigen of the rhesus system formed the highest percentage (1.6%) of the allantibodies detected. The Kell system antibodies were found only in 1.2% of the total antibodies detected (anti-K and anti-Jsb).

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Author Contributions
Zaccheaus Awortu Jeremiah – Substantial contributions to conception and design, Acquisition of data, Analysis and interpretation of data, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published
Augustina Mordi – Substantial contributions to conception and design, Acquisition of data, Analysis and interpretation of data, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published

Guarantor
The corresponding author is the guarantor of submission.

Conflict of Interest
Authors declare no conflict of interest.

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REFERENCES