Frequency of Wr\textsuperscript{a} antigen and anti-Wr\textsuperscript{a} in Brazilian blood donors

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\textbf{A B S T R A C T}

Background: Wr\textsuperscript{a} is a low-incidence antigen, which is antithetical to the high prevalence red blood cell antigen, Wr\textsuperscript{b}. Anti-Wr\textsuperscript{a} is a naturally occurring antibody that is found in approximately 1–2% of blood donors. The aim of this study was to determine the frequency of Wr\textsuperscript{a} and anti-Wr\textsuperscript{a} in Brazilian blood donors.

Methods: A total of 1662 Brazilian blood donors were molecularly analyzed using the SNaPshot methodology to determine the WR\textsuperscript{A/B} alleles and to predict the frequency of the Wr\textsuperscript{a} antigen. To detect the anti-Wr\textsuperscript{a}, samples from 1049 blood donors were analyzed using a gel test with Wr(a+) red blood cells. The serum was treated with dithiothreitol (DTT) to determine the immunoglobulin classes. Immunoglobulin (Ig)-G isotype classification was performed in a gel test using the IgG1/IgG3 card. A monocytic monolayer assay was employed to predict the clinical significance of IgG anti-Wr\textsuperscript{a}.

Results: Of the 1662 donors, only one sample had the Di'02.03 allele in heterozygous predicting the Wr(a+b+) phenotype. Anti-Wr\textsuperscript{a} was detected in 34 (3.24%) samples, 64.7% in females and 35.3% in males. Regarding the immunoglobulin class, eight (23.5%) cases of anti-Wr\textsuperscript{a} were classified as IgG and 26 (76.5%) as IgM. Of the eight cases of IgG anti-Wr\textsuperscript{a}, four were IgG1, two were IgG3 and three anti-Wr\textsuperscript{a} were not IgG3 or IgG1, and thus probably IgG2 or IgG4. The results of the monocytic monolayer assay showed that IgG anti-Wr\textsuperscript{a} might be of clinical significance.

Conclusion: This study shows a very low frequency (0.06%) of the Wr\textsuperscript{a} antigen in Brazilian blood donors. Additionally, it shows that the frequency of anti-Wr\textsuperscript{a} in this population is higher than previously reported.

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**Introduction**

The Diego blood group system is carried on band 3, a multi-pass membrane glycoprotein, which is encoded by the SLC4A1 gene. The Diego system is composed of 22 antigens: three pairs of antithetical antigens, Di\(^a\) and Di\(^b\), Wr\(^a\) and Wr\(^b\), Wu and DISK, and 16 very low frequency antigens.\(^1\) Wr\(^a\) and Wr\(^b\) antigens are related to a SNP in exon 16 (1972G>A) that encodes a lysine in Wr\(^a\) or a glutamic acid in Wr\(^b\) at amino acid position 658.\(^2\)

The Wr\(^a\) antigen, first described by Holman in 1953, has an incidence of around 1 in 1000 in Caucasian populations, but it is not reported in other ethnic groups.\(^3\) Although the Wr\(^a\) antigen has a very low incidence, anti-Wr\(^a\) is a relatively common antibody since it is often a naturally occurring antibody.\(^4\) The described incidence of anti-Wr\(^a\) in the sera of normal donors varies in different studies; it has been estimated at 1 of 100 in healthy volunteer blood donors.\(^5\) The immunoglobulin (Ig) class of anti-Wr\(^a\) can be IgM, IgG or IgM plus IgG. Alloanti-Wr\(^a\) is rarely involved in hemolytic transfusion reactions, however there are some cases reporting hemolytic disease of the fetus and newborn (HDFN) caused by anti-Wr\(^a\).\(^1\)

Antibodies against low-incidence antigens, including anti-Wr\(^a\), are difficult to identify, because the screening and panel cells rarely express these antigens.\(^6,7\) Hence, little is known about the frequency of anti-Wr\(^a\) in many populations. The knowledge of the molecular basis of the Diego blood group system and the development of molecular assays to identify the Di alleles has allowed the frequency of these alleles to be assessed in different populations. The aim of this study was to determine the frequency of the Wr\(^a\) antigen and anti-Wr\(^a\) in a Brazilian population of blood donors.

**Methods**

A total of 1662 blood samples were obtained from healthy volunteer Brazilian blood donors at the Associação Beneficente de Coleta de Sangue (Colsan), São Paulo, Brazil. The population studied was from Southeast of Brazil and it is composed of a highly admixed population.

**Molecular analysis**

DNA was extracted using the QIAamp DNA Mini Kit (Qiagen\(^\text{®}\) Inc. Valencia, CA, USA) according to the manufacturer’s instructions. To determine the WR\(^A\) and WR\(^B\) alleles and predict the frequency of the Wr\(^a\) antigen, genotyping was performed using a previous described SNaPshot\(^\text{®}\) protocol (Latini et al.\(^8\)). Fragment analyses were performed in a 3500xl Genetic Analyzer (Applied Biosystem, Foster City, CA, USA) as shown in Figure 1.

**Antibody screening**

In order to investigate the occurrence of anti-Wr\(^a\), serum samples from 1049 blood donors (638 male and 411 female donors) were initially cross-matched with a Wr\(^a\)+ red blood cell (RBC) from our collection in a gel test by an automated system (WADiana\(^\text{®}\) EXT; Grifols, Barcelona, Spain). The presence of anti-Wr\(^a\) in positive cross-matches was confirmed with two sources of Wr\(^a\)+ RBCs from commercial panels (BioRad\(^\text{®}\), Lagoa Santa, Brazil).

**Immunoglobulin classes**

To determine the Ig classes (IgG or IgM), the serum was treated with dithiothreitol (DTT, Sigma-Aldrich, Brazil). The IgG isotype classification was performed in a gel test using the IgG1/IgG3 card (BioRad\(^\text{®}\), Lagoa Santa, Brazil).

**Monocyte monolayer assay**

To predict the clinical significance of anti-Wr\(^a\), the monocyte monolayer assay (MMA) was performed as previously described\(^9\) in two samples with anti-Wr\(^a\) classified as IgG1 and one sample classified as IgG3. Using an optical microscopy, 600.
monocytes were counted to determine the percentage of reactive monocytes (RBC adhered and phagocytosed). MMA results <4% were considered negative while results ≥4% were considered positive.

Results

Wr® antigen

Of the 1662 genotyped blood samples, only one sample presented the WR’A allele in heterozygous. It was genotyped as WR’A/WR’B predicting the Wr(a+b+) phenotype.

Anti-Wr®

Anti-Wr® was detected in 34 samples from 1049 screened blood donors representing a frequency of 3.24%. Regarding the Ig classes, 8/34 (23.5%) were IgG and 26/34 (76.5%) were IgM. Of the eight IgG anti-Wr®, four were classified as IgG1 and one was isotypes as IgG3. Three samples were not classified as IgG1 or IgG3; these are probably IgG2 or IgG4, Ig classes that are not involved in severe transfusion reactions. As shown in Table 1, a higher frequency of anti-Wr® was observed in female donors (p = 0.0036, Fisher’s exact test).

MMA results (Figure 2) show that Wr® antibodies classified as IgG can potentially be clinically significant, as IgG1 antibodies presented 7–7.5% of reactive monocytes and 12.7% of IgG3 had reactive monocytes.

Discussion

This study shows novel information regarding the presence of anti-Wr® in a Brazilian population of blood donors. Although the frequency of the Wr® antigen (1:1662) is lower than that previously reported in Europeans (1:1000),6 the occurrence of anti-Wr® was higher (1:31) when compared to other studies where it ranged from 1 in 80 to 1 in 200. The frequency of anti-Wr® found in this study is similar to that found in Spain (1:37), however, the presence of the antigen in Spanish population is around 2-times (1:785)6 the frequency found in Brazilians.

The mechanisms involved in anti-Wr® production are still unclear. Some authors believe that, besides the alloimmunization in response to antigen exposure, certain proteins that can cross-react with the Wr® antigen are formed when the immune system becomes more active.7 Situations described to be involved in anti-Wr® alloimmunization are also related to immune system activation, including pregnancy, autoimmune hemolytic anemia and patients with other RBC antibodies.7 Therefore, our hypothesis is that the difference in anti-Wr® distribution between genders could be associated to pregnancy, as anti-Wr® was found in 5.2% of women and 1.8% of men.

The nature of alloimmunization might determine the antibody behavior. Our results comprising Ig class showed that IgM anti-Wr® was the predominant class, corroborating with the hypothesis of it being a naturally occurring antibody. On the other hand, four IgG1 and one IgG3 anti-Wr® with possible clinical significance were identified. Even though anti-Wr® is described to rarely cause HDFN or hemolytic transfusion reactions, probably due to the fact that anti-Wr are usually nonimmune antibodies10 1.4% of anti-Wr® found in this study

Figure 2 – Monocyte monolayer assay slides (1000×). (A) Negative control and (B) Monocyte monolayer assay with anti-Wr®.
can be of clinical significance. Due to the low incidence of the Wr antigen and the low risk of hemolytic transfusion reaction, the use of screening panels containing Wr(a+) RBCs is not required. Thereby Wr(a) incompatible transfusion can occur, but few cases of hemolytic transfusion reaction were described, been estimated to be 1 in 500,000.11

In summary, the Wr(a) antigen has a very low frequency in Brazilian blood donors and anti-Wr(a) has a higher frequency than reported in other populations. Considering the low frequency of the antigen and the few cases of mild HDFN related to anti-Wr(a), clinical impact is discussable as well the requirement of RBC reagent to identify them.

Conflicts of interest

The authors declare no conflicts of interest.

REFERENCES