# ABO and Rh Blood Group Incompatibility among Icteric Neonates and their Mothers in Jos, Nigeria

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## Abstract

Context: ABO and Rh blood groups play significant roles in health and diseases, one of which is blood group incompatibility, a common cause of neonatal hyperbilirubinemia. Aims: The aim of the study is to determine the frequency of ABO and Rh blood group incompatibilities among icteric neonates and their mothers for the purpose of instituting intervention for better neonatal outcomes. Settings and Design: This study was conducted at the Special Care Baby Units (SCBUs) of the Jos University Teaching Hospital (JUTH), Bingham University Teaching Hospital (BHUTH), and Plateau State Specialist Hospital (PSSH), Jos, Nigeria, from March 2013 to February 2014. It was a descriptive cross-sectional study that includes all jaundiced neonates admitted into the SCBUs and neonates who developed jaundice on admission in the neonatal wards. Subjects and Methods: A total of 150 icteric neonates admitted into the SCBUs of the JUTH, PSSH, and the BHUTH were recruited for this study with their parent's consent. Blood samples were collected from the neonates and their mothers in ethylene diaminetetra-acetic acid and plain bottles for ABO and Rh blood grouping, direct antihuman globulin test, and serum bilirubin (SB) assays. Statistical Analysis Used: Data obtained were analyzed using Epi Info Version 6 software. The results were reported in tables and frequencies, categorical variables were expressed as proportions, whereas continuous variables were expressed as mean ± standard deviation and were analyzed using Student's "t"-test. The level of statistically significant relationship was set at  $P \le 5\%$  ( $P \le 0.05$ ). Results: Thirty-four (22.7%) mother-neonate pair had ABO incompatibility with 14 mothers (9.3%) with blood group O, having neonates with blood group A and a mean SB of 249.5  $\pm$  131.4  $\mu$ mol/L. Three (2.0%) mothers were Rh D negative while their neonates were Rh D positive and showed a positive direct antihuman globulin test with a mean SB of  $322.1 \pm 246.7 \mu$ mol/L. **Conclusions:** Blood group O and Rh D-positive blood groups predominate while ABO and Rh incompatibilities present a risk for hyperbilirubinemia among icteric neonates in Jos, Nigeria.

Keywords: Blood group, hyperbilirubinemia, incompatibility, neonate

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## INTRODUCTION

Blood groups are based on the existence of antigens made up of molecules such as glycoproteins and glycolipids on the red blood cell membranes.<sup>[1]</sup> Approximately 400 blood group antigens have been described, with the ABO and Rh blood groups being the most frequently studied genetic markers in humans.<sup>[2,3]</sup> The ABO blood group determined by the presence or absence of blood group antigens A or B has a natural distribution among humans with the O blood group predominating in most instances, followed by blood group A.<sup>[4-6]</sup>

The Rh blood group system is the second most frequent blood group system of clinical significance determined by a

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highly immunogenic Rh D antigen.<sup>[6]</sup> Human populations with Rh-negative status are few with slight racial differences, 5.5% in South India, 5% in Nairobi, and 4.8% in Nigeria, while the Caucasians have a 15% rate of Rh-negative blood group.<sup>[7-9]</sup>

In blood group immunology, individuals who lack a particular blood group antigen produce antibodies against that antigen leading to an immunologic reaction, hemolysis, and

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hyperbilirubinemia, as it can occur in pregnancy because of fetomaternal transfusion.<sup>[9]</sup> ABO incompatibility occurs most frequently but rarely causes severe haemolytic disease of the foetus and newborn (HDFN), while the highly immunogenic Rh D antigen can cause immune response with severe HDFN.<sup>[9-11]</sup>

Studies among Nigerian children have shown that neonatal jaundice is a significant cause of morbidity and mortality in the pediatric age group.<sup>[12-17]</sup> This study is to determine the frequency of ABO and Rh blood group incompatibility among icteric neonates and their mothers with a view on raising awareness on the knowledge of blood groups, screening, and prophylaxis against isoimmunization in pregnancy, thereby improving the outcome of hemolytic disease of the fetus and newborn with its attendant complications.

## SUBJECTS AND METHODS

This study was conducted at the Special Care Baby Units (SCBUs) of the Jos University Teaching Hospital, Bingham University Teaching Hospital, and Plateau State Specialist Hospital, Jos, Nigeria, from March 2013 to February 2014. It was a descriptive cross-sectional study that includes all jaundiced neonates admitted into the SCBUs and neonates who developed jaundice on admission in the neonatal wards. Neonates whose parents did not consent, those with cephalhematomas, bleeding tendencies, birth asphyxia, and those recently transfused were excluded from the study. Ethical approval was obtained from the Ethics Committee of the participating institutions, while written informed consent was obtained from the parents or guardians of the neonates. The neonates were recruited using the nonprobability convenience sampling technique, and the questionnaire was self-administered to obtain relevant information. The neonates were examined, and 5 ml of the blood sample was collected for ABO and Rh blood group, direct antihuman globulin test, and serum bilirubin (SB) assays. Three milliliters of mothers' blood sample was also collected for ABO and Rh blood grouping. SB concentration >104 µmol/L was considered pathologic. Data obtained were analyzed using Epi Info Version 6 software (CDC 2000, Atlanta, Georgia, USA). The results were reported in tables and frequencies, categorical variables were expressed as proportions, whereas continuous variables were expressed as mean  $\pm$  standard deviation and were analyzed using Student's "t"-test. The level of statistically significant relationship was set at  $P \le 5\%$  ( $P \le 0.05$ ).

## RESULTS

Sixty-three (42.0%) of 150 neonates and 85 (56.7%) of 150 mothers were ABO blood group O. One hundred and forty-six (97.3%) neonates were Rh D positive, while 4 (2.7%) mothers were Rh D negative [Table 1].

Mother-neonates ABO and Rh blood group incompatibility

Thirty-four (22.7%) mother-neonate pairs were ABO incompatible. Twenty (13.3%) blood group O mothers had 20 (13.3%) neonates with blood group B and a mean SB of  $249.5 \pm 131.4$ , while 14 (9.3%) mothers had neonates with blood

group A and a mean SB of  $229.5 \pm 110.0 \,\mu$ mol/L. Two (1.3%) of those with ABO incompatibility had a positive direct antihuman globulin test. Three (2.0%) mother–neonate pairs were Rh D negative-positive with positive direct antihuman globulin test and SB of  $322.1 \pm 246.7 \,\mu$ mol/L, whereas three others pairs were Rh D positive–negative and with negative direct antihuman globulin test. No statistically significant difference was found in the severity of hyperbilirubinemia between the groups with ABO or Rh incompatibility, *P* values of 0.63 and 0.99, respectively [Table 2].

## DISCUSSION

This study demonstrated the predominance of blood group O and Rh positivity in both the neonates and their mothers. These findings are in accordance with other studies in Nigeria as documented by Bakare et al. in Ogbomosho, South West, Nigeria, and Pennap et al. in Keffi, North Central Nigeria.<sup>[18,19]</sup> Racial differences may exist in the distribution of some of the ABO blood group antigens, but no significant differences were found comparing the prevalence of 45.2%, 39.7%, 10.9%, and 4.1% for blood groups O, A, B, and AB among the Caucasians in the United States with the frequencies for the neonates and their mothers in our study.<sup>[20]</sup> ABO incompatibility is the most commonly reported serologic cause of neonatal jaundice but clinically mild as a cause of hemolytic disease of the newborn (HDFN) with hyperbilirubinemia than the Rh incompatibility.<sup>[20,21]</sup> This could be attributed to the fact that Rh incompatibility which was once an extremely common cause of severe hemolysis has now been reduced by the prophylactic

Table 1: Mothers' and neonates' blood groups						
Blood group	Mother ( <i>n</i> =150), <i>n</i> (%)	Baby ( <i>n</i> =150), <i>n</i> (%)				
ABO						
0	85 (56.7)	63 (42.0)				
А	24 (16.0)	34 (22.7)				
В	33 (22.0)	47 (31.3)				
AB	8 (5.3)	6 (4.0)				
Total	150 (100)	150 (100)				
Rh						
Positive	146 (97.3)	146 (97.3)				
Negative	4 (2.7)	4 (2.7)				
Total	150 (100)	150 (100)				

## Table 2: Mother-neonates ABO and Rh blood groupincompatibility

Blood group	Mother-neonate incompatibility	п	DAT		SB	Р
			Positive	Negative	(µmol/L)	
ABO	O-A	14	0	14	249.5±131.4	0.63
	O-B	20	2	18	229.5±110.0	
Total		34	2	32		
Rh	Positive-negative	3	0	3	324.2±198.8	0.99
	Negative-positive	3	3	0	322.1±246.7	
Total		6	3	3		

DAT: Direct antihuman globulin test, SB: Serum bilirubin

administration of anti-D globulin to Rh-negative mothers.<sup>[21]</sup> Frequency of ABO incompatibilities in our study was higher than Rh incompatibility, but the direct antihuman globulin test did not confirm the isoimmunization in most of them, except for Rh-negative mothers with Rh-positive neonates where the evidence of isoimmunization was demonstrated in all. The low antigenicity expressed at birth, naturally occurring in anti-A and anti-B which are predominantly immunoglobulin (Ig) M, and anecdotal reports suggesting that direct antihuman globulin test has low sensitivity may be responsible for the inability to confirm isoimmunization in the neonates at risk of blood group incompatibilities in this study.<sup>[22]</sup> Hyperbilirubinemia of varying degrees was demonstrated where these incompatibilities were found; however, no statistical difference was noticed among those with ABO or Rh incompatibilities. This finding does not support variation in immunogenicity, especially between anti-A and anti-B, where HDFN in blacks due to anti-B is said to be more severe than with anti-A.<sup>[23]</sup> Other risk factors for neonatal hyperbilirubinemia were not considered in this studies such as neonatal sepsis and enzymopathies could coexist with the blood group incompatibility; however, Kaplan et al. reported no difference in the degree of hyperbilirubinemia even in the presence of these risk factors.[24]

Frequency of ABO incompatibility, 22.7% in this study is in contrast to 7.6% reported by Israel-Aina and Omoigberale in Benin.<sup>[25]</sup> This marked difference may not be unconnected to the size of our study population. On direct antihuman globulin test, our finding was close to the rate of 5% reported by Fadairo *et al.* in Ile-Ife, but in contrast to a previous report that about 36% of Nigerian babies with ABO incompatibility may have evidence of isoimmunization.<sup>[26,27]</sup> Contrary to the frequency found in our study, Manning *et al.* study in the United Kingdom and several other researchers in Canada and Turkey, reported ABO incompatibility as the most common cause of neonatal jaundice followed by Rh incompatibility and G6PD deficiency.<sup>[28-30]</sup> This report may be as a result of the few Rh-negative individuals as well as the low prevalence of G6PD deficiency in their environment.<sup>[28-30]</sup>

Introduction of anti-D and increase awareness has reduced the spectrum of HDFN over the past few decades in Europe and America shifting attention to other allo-antibodies which could be the cause of the emerging HDFN being encountered in developed societies. The low prevalence of Rh incompatibility observed in this study is not surprising due to increase awareness and antenatal clinic attendance as well as the few Rh-negative women in the African and Asian populations.<sup>[31,32]</sup> However, isoimmunization was confirmed in all the Rh-positive neonates, whose mothers were negative and this call for early monitoring and administration of anti-D to Rh-negative mothers for better neonatal outcomes.

Neonatal sepsis, G6PD deficiency, and other causes of neonatal jaundice may be responsible for the hyperbilirubinemia seen in the other neonatal population not accounted for in this study. Collaboration for further studies that will include detailed red blood cell-specific antigen and antibody screening, G6PD enzyme assay, as well as microbiological search for pathogens that could cause neonatal hyperbilirubinemia in our environment will be pursued in due course.

## CONCLUSIONS

ABO and Rh blood group distribution in this study closely reflect the global distribution of blood groups, while ABO incompatibility has been shown to be a relatively common occurrence among mothers and neonates in our environment compared to Rh incompatibility. In view of the neonatal morbidity and mortality associated with hyperbilirubinemia, every possible cause of hyperbilirubinemia including all blood group incompatibilities in our environment must be investigated using the quality standard screening test during pregnancy for early diagnosis and institution of prophylactic measures for better neonatal care.

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### **Conflicts of interest**

There are no conflicts of interest.

## REFERENCES

- Hoffbrand AV. Blood transfusion. In: Hoffbrand AV, Moss PA, Pettit JE, editors. Essential Haematology. 6<sup>th</sup> ed. Oxford: Blackwell Publishers Ltd.; 2011. p. 379-412.
- Enosolease ME, Bazuaye GN. Distribution of ABO and Rh-D blood groups in the Benin area of Niger-Delta: Implication for regional blood transfusion. Asian J Transfus Sci 2008;2:3-5.
- Conteras M, Lubenko S. Immunohaematology: Introduction. In: Hoffbrand AV, Lewis SM, Tuddenhan EG, editors. Postgraduate Haematology. 4<sup>th</sup> ed. London: Arnold Publishers; 2005.p. 207-24.
- Egesie UG, Egesie OJ, Usar I, Johnbull TO. Distribution of ABO, Rhesus blood and haemoglobin electrophoresis among the undergraduate students of Niger Delta State University, Nigeria. Niger J Physiol Sci 2008;23:5-8.
- Knowles S, Poole G. Human blood group systems. In: Practical Transfusion Medicine. 1<sup>st</sup> ed. Oxford: Blackwell Publishers Ltd.; 2002. p. 72-9.
- Durosinmi M. Blood transfusion medicine. In: Olusegun O, editor. Essential Pathology for Clinical Students in the Tropics. 1<sup>st</sup> ed. Ibadan: Caltop Publications; 2004. p. 444-69.
- Adeyemo OA, Soboyemo OB. Frequency distribution of ABO, Rh blood groups and Blood Genotypes among the cell biology genetics students of University of Lagos, Nigeria. Afr Biotech 2006;5:2062-5.
- Khan MN, Khaliq I, Bakhsh A, Akhtar MS, Amin-ud-Din M. Distribution of ABO and Rh D blood groups in the population of Poonch District, Azad Jammu and Kashmir. East Mediterr Health J 2009;15:717-21.
- Reid ME, Francis CL. Erythrocyte antigens and antibodies. In: Beutler E, Coller B S, Litchman MA, Kipps TJ, Seligsohn U, editors. Williams Hematology. 9th ed. New York: McGraw Hill Education; 2000. p. 2329-51.

- Regan F. Blood cell antigens and antibodies. In: Lewis SM, Bain BJ, Bates I, Laffan MA, editors. Practical Haematology. 11<sup>th</sup> ed. London: Churchill Livingstone; 2012. p. 25-58.
- Marwaha N, Chaudhary RK. Blood groups: In: Saxena R, Pati PH, Mahapatra M, editors. De Gruchy's Clinical Haematology in Medical Practice. 6<sup>th</sup> ed. New Delhi: Wiley India; 2013: 432-453.
- Maisels MJ, Gifford K, Antle CE, Leib GR. Jaundice in the healthy newborn infant: A new approach to an old problem. Pediatrics 1988;81:505-11.
- Airede AI. Relation of peak total serum bilirubin concentrations to neurodevelopmental outcome at 2 years of age in premature African neonates. Ann Trop Paediatr 1992;12:249-54.
- Slusher TM, Angyo IA, Bode-Thomas F, Akor F, Pam SD, Adetunji AA, et al. Transcutaneous bilirubin measurements and serum total bilirubin levels in indigenous African infants. Pediatrics 2004;113:1636-41.
- Lagunju IA, Okafor OO. An analysis of disorders seen at the Paediatric Neurology Clinic, University College Hospital, Ibadan, Nigeria. West Afr J Med 2009;28:38-42.
- Jones L, Wilson D. The blood and Haematologic system. In: Martin RJ, Fanarroff A, Walsh M, editors. Neonatal-Perinatal Medicine Diseases of Fetus and Infant. 9<sup>th</sup> ed. Missouri: Saunders-Elservier; 2011, p. 1303-60.
- Toma BO, Ige OO, Abok II, Onwuanaku C, Abah RO, Donli A. Pattern of neonatal admissions and outcome in a tertiary institution in North central Nigeria. J Med Trop 2013;15:121-5.
- Bakare AA, Azeez MA, Agbolade JO. Gene frequencies of ABO and Rhesus blood groups and haemoglobin variants in Ogbomosho, South West, Nigeria. Afr J Biotech 2006;5:224-9.
- Pennap GR, Envoh E, Igbawua I. Frequency distribution of haemoglobin variants ABO and Rhesus blood groups among students of African descent. Br Microbiol Res J 2011;1:33-40.
- Garratty G, Glynn SA, McEntire R; Retrovirus Epidemiology Donor Study. ABO and Rh (D) phenotype frequencies of different racial/ethnic groups in the United States. Transfusion 2004;44:703-6.
- 21. Reddy VV. Intracorpuscular defects leading to increased erythrocyte

destruction. In: Rodak BF, Fristma GA, Doig K, editors. Haematology: Clinical Principles and Applications. 3<sup>rd</sup> ed. Philadelphia: Saunders-Elsevier; 2007. p. 286-310.

- Wainer S, Rabi J, Lyon M. Coombs' testing and neonatal hyperbilirubinemia. CMAJ 2007;176:972-3.
- Dutta AB. Transfusion practice: Clinical aspects and Applications. In: Jain VK, Jain SK, editors. Blood Banking and Transfusion. 1<sup>st</sup> ed. New-Delhi: CBS Publishers; 2006. p. 213-321.
- 24. Kaplan M, Vreman HJ, Hammerman C, Leiter C, Rudensky B, MacDonald MG, *et al.* Combination of ABO blood group incompatibility and glucose-6-phosphate dehydrogenase deficiency: effect on hemolysis and neonatal hyperbilirubinemia. Acta Paediatr 1998;87:455-7.
- Israel-Aina II, Omoigberale AI. Risk factors for neonatal jaundice in babies presenting at the University of Benin Teaching Hospital Benin City. Niger J Paed 2012;39:159-63.
- Fadairo JK, Aladenika ST, Osaiyuwa C, Olaniyan MF, Agatishe K. Evaluation of some aetiological factors of haemolytic disease of the New born in Ile-Ife. Open J Clin Diag 2014;4:5-11.
- Owa JA, Durosinmi MA, Alabi AO. Determinants of severity of neonatal hyperbilirubinaemia in ABO incompatibility in Nigeria. Trop Doct 1991;21:19-22.
- Manning D, Todd P, Maxwell M, Jane Platt M. Prospective surveillance study of severe hyperbilirubinaemia in the newborn in the UK and Ireland. Arch Dis Child Fetal Neonatal Ed 2007;92:F342-6.
- Sgro M, Campbell D, Shah V. Incidence and causes of severe neonatal hyperbilirubinemia in Canada. CMAJ 2006;175:587-90.
- Atay E, Bozaykut A, Ileak I. Glucose-6-phosphate dehydrogenase deficiency in neonates and indirect hyperbilirubinaemia. J Trop Paediatr 2006;52:56-8.
- World Health Organization. Neonatal and Perinatal Mortality: Country, Regional and Global Estimates. Geneva, Switzerland: World Health Organization; 2006. p. 1-75.
- Chandra T, Gupta A. Frequencies of ABO and Rh Blood groups in Blood Donors. Asian J Trans Sci 2012;6:52-3.