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ABO and Rhesus blood group distribution and frequency among blood donors at Kilimanjaro Christian Medical Center, Moshi, Tanzania

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Abstract

Objective: This study aims to determine the distribution of blood groups and the demographic background of blood donors in a referral hospital in Northern Tanzania.

Results: The most common blood group was O (52.3%) and the least common was AB (3.18%). 97.7% of the blood donors were Rh positive and the rest were Rh negative. Most donors were young adults, representing the age group of 19–29. The majority of donors were male (88.1%) and the majority (90.8%) were replacement while the remainder was voluntary donors.

Keywords: ABO, Rhesus factor, Blood donation, Transfusion medicine, Tanzania

Introduction

In 1900, Karl Landsteiner of the University of Vienna identified that red blood cells contain antigens on their surfaces, and that blood plasma contains antibodies targeted to particular antigens [1–3]. This discovery is the basis of modern-day blood grouping and transfusion medicine. Demand for whole blood and blood products is high in Tanzania where current supply does not meet the need. Post-partum hemorrhage is a major cause of maternal mortality in Tanzania and lack of adequate blood supply for transfusion is one of the contributing factors [4]. As well, there is a high rate of road traffic injuries, infectious disease such as HIV, gastrointestinal bleeds, and an increasing rate of elective surgeries [4, 5]. Blood banks require timely information concerning the distribution and frequency of blood groups in order to ensure adequate supply of the most medically useful blood types.

Red blood cells antigens, which are the basis of blood grouping, consist of proteins and carbohydrates attached to lipids or proteins. There are more than 100 blood group systems involving over 500 antigens in which ABO is the most studied group in the human population [6]. These antigens have various functions, such as membrane structural integrity and transportation of molecules through membranes. ABO antigens are highly expressed on human tissues and most epithelial and endothelial cells [7, 8]. Antigen expression can influence the development of particular infections as well as certain malignancies. The Multinational Pancreatic Cancer Consortium successfully identified susceptibility loci in the ABO gene for pancreatic cancer pathogenesis [9]. Other studies have showed an association between gastric cancer and blood group A related to a higher susceptibility of *Helicobacter pylori* infection. Different hypothetical models such as inflammation, immune system surveillance and cell membrane signaling have been developed to explain the mechanism of cancer susceptibility among people with varying blood groups [10–15].

The knowledge of the distribution of Rhesus antigen in a population is critical in managing a transfusion service

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in areas such as antenatal serology, paternity testing as well as selecting compatible blood and blood products. Even after Karl Landsteiner’s discovery in 1900, transfusion reactions were still prevalent [1]. It was not until 1940 when Landsteiner and Weiner discovered the Rh factor that transfusion medicine involved less risk. Immunogenicity of the Rh factor along with A, B antigens made it mandatory for pre-transfusion testing [1, 16]. Currently there are more than 50 antigens in the Rh blood group system but the principal Rh antigens of medical interest are D, C, E, c and e [16]. A person with Rhesus antigen is referred to as Rhesus positive while individuals lacking the antigen are Rhesus negative. When a Rhesus negative person is exposed to Rhesus positive blood, antibodies will be produced, which cause potentially fatal hemolytic reactions. There is a lack of published data in Tanzania describing the distribution of ABO and Rh among blood donors, and this information is scarce in Africa as a whole [17, 18]. This study aims to determine the distribution of blood groups and the demographic background of blood donors in a referral hospital in Northern Tanzania.

Main text

Methods

A cross-sectional study was conducted at KCMC Clinical Laboratory in the blood transfusion unit over a period of 6 months from October 2014 to March 2015. Samples were analyzed from both voluntary and replacement donors at KCMC Clinical Laboratory, Moshi, Tanzania collected from October 2013 to September 2014. Voluntary blood donation involves a donor giving blood, plasma or cellular components of his or her own free will while replacement or family donors are those who give blood when required by a family or community member [19].

The inclusion criteria for this study were donors between the ages of 18–70, with a personal weight above 50 kg, and who met the hemoglobin cut off criteria. All donors were required to have a haemoglobin level of at least 12.0 g/dL for females and 13.0 g/dL for males as per WHO standards. A total of 1845 participants out of 2200 total donors met inclusion criteria. 1815 participants (98.4%) had the complete information required for analysis in this study.

During blood donation approximately 4 mL of blood from each donor was collected in EDTA tubes for analysis. ABO and Rh status were analyzed by tube method using commercially prepared anti-A, anti-B, anti-AB and anti-D antisera blood types. To do so correctly, we followed the specific procedures outlined in the manufacturer’s manual. Prepared 5% suspensions of red blood cells in normal saline were used. Four different tubes

labelled with donor unit numbers were added with one drop of antisera A, B, AB and D. To every tube with specific antisera one drop of 5% cell suspension was added and each sample was macroscopically observed for agglutination.

Descriptive statistics was used whereby data were summarized using frequency and percentages. Fisher’s Exact test was used to test association between exposures and outcome of interest. A *P value* of less than 0.05 (2-tails) was considered as statistically significant.

Interpretation of results

Positive: Agglutination indicates positive reactions to respective group or Rhesus factor.

Negative: No Agglutination indicates negative reactions to respective group or Rhesus factor (Table 3).

Results

A total of 1845 participants met inclusion criteria. Out of the total participants, there was complete information for 1815 (98.4%) blood donors. As shown in Table 1, there were more male participants (88%, n = 1597) as compared with female participants (12%, n = 218). The age distribution of the participants was 43% (n = 773), 29% (n = 530), and 28% (n = 508) for the age groups of 18–29, 30–39 and 40–65 respectively.

The most common blood type among the participants was blood group O (52%, n = 949), followed by blood group A (26%, n = 465), blood group B (19%, n = 342) and blood group AB (3%, n = 59%). As shown in Table 2, 98% (n = 1773) of participants were Rhesus positive while 2% (n = 42) were Rh negative.

In both sexes, blood group O was the commonest being 52% (n = 830) among male and 55% (n = 119) among female participants. The least common blood group was blood group AB representing 3% of male participants and 1.8% of female participants. There was no difference between the blood type and the sex ($\chi^2 = 3.7021$, *P value* 0.895). Except for blood group AB, all of the other blood groups had Rh-negative antigens in the donated blood. The prevalence of Rh-negative antigens were 3%

Table 1 Gender and age distribution of the participants (N = 1815)

Characteristic	Frequency (n)	Percentage (%)
Sex		
Female	218	12
Male	1597	88
Age (years)		
18–29	773	43
30–39	530	29
40–65	508	28

Table 2 Blood type and Rh antigens of the participants (N = 1815)

	Frequency n (%)
Blood type	
A	465 (26)
B	342 (19)
O	949 (52)
AB	59 (3)
Rh antigens	
Rh negative	42 (2)
Rh positive	1773 (98)

(n = 13), 2% (n = 8) and 2% (n = 21) for blood group A, B and O respectively. There were no differences in RH between different blood groups ($\chi^2 = 1.923$, *P value* 0.712) (Table 3).

Discussion

This study determined the distribution of ABO and Rh antigens among blood donors in Northeastern Tanzania, where there is a lack of data on this subject [17, 18]. The current study showed that the majority of donors were male, which is consistent with other studies in Africa and in most regions globally [17–19, 20, 22, 23]. One contributing factor might be that women do not meet donation cut-off values for hemoglobin given normal menses, menorrhagia, prenatal iron deficiency anemia and postnatal blood loss. From a cultural perspective, in various African countries it may be more likely for males to donate blood given long-standing beliefs that women are not as physically strong as men [22, 23]. In Western regions such as Europe, women were found to have higher rates of adverse reactions, primarily vasovagal events, and were also not as likely to meet hemoglobin cut off requirements for donation [23].

In studies spanning diverse cultural and geographic groups, the most common age range for blood donation is 18–44 years [24]. Our study results are consistent with this global trend. The improved interest and ability

among younger adults to donate may be related to awareness, better physical health, and greater mobility. Older individuals may suffer from medical conditions such as ischemic heart disease, diabetes mellitus, malignancy and hypertension hence negatively impacting their ability to be well enough to donate blood. There is a need to better understand the potential of adolescents to contribute [25], however, which was not addressed in this study.

The majority of donors in our study were replacement (90%) while the minorities were voluntary (10%). This is consistent with other studies and global trends [19, 21, 26]. The donation of blood by voluntary non-remunerated blood donors is critical for the safety and sustainability of national blood supplies. National blood donation systems in which replacement donors dominate are typically unable to meet clinical demands for blood while paid family members contributing often poses serious threats to the health and safety of the recipients and the donors. WHO recommendations are therefore to create health systems based 100% on voluntary donation [19].

The predominant blood group in our study was type O and the least common was AB. This is consistent with studies in Nigeria, which also showed the predominant group to be O and the least common to be AB [27]. In Kenya, Uganda, Mauritania and Ethiopia similar studies also showed the predominant blood group to be O and the least prevalent to be AB [27–30]. These trends, in keeping with other studies, may suggest that blood group AB is the least dominant while O is the most dominant overall across the continent.

However, there is regional variability; some studies show that in Western and Central Africa, the most predominant group was B while in Eastern and Southern countries, blood group O dominated [31–33]. In Pakistan, one study showed that blood group B is the most dominant while in Nepal it was blood group B. In Britain the most predominant blood group is O and the least is AB [31]. These regional differences may be explained by genetic mapping and the varying origins of diverse ethnic groups [31].

Determination of Rh status is crucial in clinical contexts in order to ensure patient safety. Rh factor is of interest because of its marked immunogenicity. In the case of the D antigen, patients who do not produce the D antigen will produce anti-D if they encounter the D antigen on transfused red blood cells. This process may result in a hemolytic transfusion reaction or, in the case of newborn red blood cells, hemolytic disease of the newborn [31]. For this reason, it is important to determine the Rh status in clinical settings and for research purposes.

Individuals who are Rhesus negative in our study were only 2.3% in contrast to other studies, which showed a range between 5 and 17%. Our study showed a slightly

Table 3 Distribution of Rh antigens per ABO blood group among participants (N = 1815)

	Rh –ve n (%)	Rh +ve n (%)
A	13 (3)	452 (97)
B	8 (2)	334 (98)
O	21 (2)	928 (98)
AB	–	59 (100)

ABO and Rhesus blood groups distribution among the first time blood donors at KCMC referral and teaching hospital in Northern Tanzania

lower prevalence of Rh positive blood donors in comparison to other studies in the African continent as well as in comparison with global trends [34–36]. Given the number of participants in this study, however, this cannot be said to be statistically significant. Globally we share the same blood group types however clearly there are some geographic, regional, and ethnic differences. Ensuring adequate Rh positive blood supply is important in the context of patient safety. As well, the growing literature investigating the association of blood groups with the pathogenesis of cancer requires locally specific information on Rh distribution among other factors [37].

Conclusion

Up-to-date knowledge of the distribution of blood types in a local setting is critical to the functioning of any national health service. To date, there has been a lack of data on this important topic in Tanzania. Our study provides detailed information concerning the blood type and demographic information of blood donors in the North-eastern region of Tanzania. Similar studies are needed across the country and further research and mobilization are required to meet WHO recommendations on voluntary blood donation.

Limitations

This study was conducted in the Northeastern region of Tanzania; the results should not be generalized to Tanzania as a whole.

Abbreviations

Rh: Rhesus; RTI: road traffic injuries; KCMC: Kilimanjaro Christian Medical center; WHO: World Health Organization.

Authors' contributions

JJP, EON, AM and ERS participated in designing the study and performed the data collection. OJ wrote the analysis part of the methodology section, conducted statistical analysis, and wrote the results section of the manuscript. ERS wrote the first draft of the manuscript. All authors read and approved the final manuscript.

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Competing interests

The authors declare that they have no competing interests.

Availability of data and materials

An excel sheet with all primary data is available upon request.

Consent to publish

Permission to publish this work was obtained from study participants as well as the KCMC hospital administration.

Ethics approval and consent to participate

The study was conducted following ethical approval from Kilimanjaro Christian Medical University College Review Board and approval from the medical director of KCMC. Written informed consent was obtained from all participants.

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References

- Lefrère J, Berche P. Landsteiner discovers the blood groups. *Transfus Clin Biol*. 2010;17(1):1–8.
- Garraty G, Dzik W, Issitt PD, Lublin DM, Reid ME, Zelinski T. Terminology for blood group antigens and genes—historical origins and guidelines in the new millennium. *Transfusion*. 2000;40(4):477–89.
- Hosei E. Biological and clinical aspects of ABO blood group system. *J Med Investig*. 2008;55(3–4):174–82.
- Pembe A, Paulo C, D'mello B, Roosmalen J. Maternal mortality at Muhimbili national hospital in Dar-es-Salaam, Tanzania in the year 2011. *BMC Pregnancy Childbirth*. 2014;14:320.
- Komba DD. Risk factors and road traffic accidents in Tanzania: a case study of Kibaha District. *NTNU Open*. 2006;23:1–70.
- Dziczkowski JS, Anderson KC. Blood group antigens and therapy in *Harrison's Principles of International Medicine*. 14th ed. New York: McGraw Hill; 1998.
- Franchini M, Favaloro EJ, Targher G, Lippi G. ABO blood group, hypercoagulability, and cardiovascular and cancer risk. *Crit Rev Clin Lab Sci*. 2012;49:137–49.
- Reid ME, Mohandas N. Red blood cell blood group antigens: structure and function. *Semin Hematol*. 2004;41:93–117.
- Amundadottir L, Kraft P, Stolzenberg-Solomon RZ, Fuchs CS, Petersen GM, Arslan AA, et al. Genomewide association study identifies variants in the ABO locus associated with susceptibility to pancreatic cancer. *Nat Genet*. 2009;41:986–90.
- Gong Y, Yang YS, Zhang XM, Su M, Wang J, Han JD, et al. ABO blood type, diabetes and risk of gastrointestinal cancer in northern China. *World J Gastroenterol*. 2012;18:563–9.
- Gates MA, Xu M, Chen WY, Kraft P, Hankinson SE, Wolpin BM. ABO blood group and breast cancer incidence and survival. *Int J Cancer*. 2012;130:2129–37.
- Xie J, Qureshi AA, Li Y, Han J. ABO blood group and incidence of skin cancer. *PLoS ONE*. 2010;5(e11972):16.
- Wang DS, Chen DL, Ren C, Wang ZQ, Qiu MZ, Luo HY, et al. ABO blood group, hepatitis B viral infection and risk of pancreatic cancer. *Int J Cancer*. 2012;131:461–8.
- Sheng L, Sun X, Zhang L, Su D. ABO blood group and nasopharyngeal carcinoma risk in a population of Southeast China. *Int J Cancer*. 2013;133:893–7.

15. Tavasolian F, Abdollahi E, Vakili M, Amini A. Relationship between ABO blood group and Acute Lymphoblastic Leukemia. *Iran J Pediatr Hematol Oncol.* 2014;4:1–4.
16. Gundrajukuppam DK, Vijaya SB, Rajendran A, Sarella JD. Prevalence of principal Rh blood group antigens in blood donors at the blood bank of a Tertiary Care Hospital in Southern India. *J Clin Diagn Res.* 2016;10(5):EC07–10.
17. Bloch EM, Vermeulen M, Murphy E. Blood transfusion safety in Africa: a literature review of infectious disease and organizational challenges. *Transfus Med Rev.* 2012;26(2):164–80.
18. Metee M, Magesa PM, Lyamuya EF. Seroprevalence of human immunodeficiency virus, hepatitis B and C viruses and syphilis infections among blood donors at the Muhimbili National Hospital in Dar es Salaam, Tanzania. *Biomed Cent.* 2016;6(21):419–24.
19. World Health Organization. Towards 100% voluntary blood donation: a global framework for action. Geneva: WHO Publishing Library; 2010.
20. Tessema B, Yismaw G, Kassu A, et al. Seroprevalence of HIV, HBV, HCV and syphilis infections among blood donors at Gondar University Teaching Hospital, Northwest Ethiopia: declining trends over a period of five years. *Biomed Cent Infect Dis.* 2010;10:111.
21. Messih IYA, Ismail MA, Saad AA, Azer MR. The degree of safety of family replacement donors versus voluntary non-remunerated donors in an Egyptian population: a comparative study. *Blood Transfus.* 2014;12(2):159–65.
22. Rushton DH, Dover R, Sainsbury AW, et al. Why should women have lower reference limits for haemoglobin and ferritin concentrations than men. *BMJ.* 2011;322(7298):1355–7.
23. Bani M, Giussani B. Gender differences in giving blood: a review of the literature. *Blood Transfus.* 2010;8:278–87.
24. World Health Organization. Global database on blood safety. Age distribution of blood donors by country. Geneva: WHO Publishing Library; 2011.
25. Zito E, Alfrieri S, Maurizio M, Vincenzo S, Giovanna C. Adolescents and blood donation: motivations, hurdles and possible recruitment strategies. *Blood Transfus.* 2012;10(1):45–58.
26. Nuinon M, Kruachan K, Sengking K, Horpet D, Sungyuan U. Thalassemia and hemoglobin E in southern Thai blood donors. *Adv Hematol.* 2014;2014:932306.
27. Akanmu AS, Oyedeji OA, Adeyemo TA, Ogbenna AA. Estimating the risk of ABO hemolytic disease of the newborn in Lagos. *J Blood Transfus.* 2015;2015:560738.
28. Lyko J, Gaertner H, Kaviti JN, Kariithi MW, Akoto B. Blood-group systems ABO and RH in the Kenyan population. *Folia Med Cracov.* 1992;33(1–4):85–92.
29. Sebabi EC, Nzaro E. Distribution of ABO and Rh (D) phenotypes in Uganda. *Vox Sang.* 1974;26(1):74–82.
30. Alemu G, Mama M. Assessing ABO/Rh blood group frequency and association with asymptomatic malaria among blood donors attending Arba Minch blood bank, South Ethiopia. *Malar Res Treat.* 2016;2016:8043768.
31. Bodmer W. Genetic characterization of human populations: from ABO to a genetic map of the British people. *Genetics.* 2015;199(2):267–79.
32. Hamed CT, Bollahi MA, Abdelhamid I, Med Mahmoud MA, Ba B, Ghaber S, Habti N, Houmeida A. Frequencies and ethnic distribution of ABO and Rh(D) blood groups in Mauritania: results of first nationwide study. *Int J Immunogenet.* 2012;39(2):151–4.
33. Dean L. Blood groups and red cell antigens. Bethesda: National Center for Biotechnology Information (US); 2005.
34. Gundrajukuppam DK, Vijaya SB, Rajendran A, Sarella JD. Prevalence of ABO and Rhesus blood groups in blood donors. *J Clin Diagn Res.* 2014;8(12):7–10.
35. McLachlan S, Giambartolomei C, White J, Charoen P, Wong A, Finan C, et al. Replication and characterization of association between ABO SNPs and red blood cell traits by meta-analysis in Europeans. *PLoS ONE.* 2016;11(6):e0156914.
36. Osaro E, Charles AT. The challenges of meeting the blood transfusion requirements in Sub-Saharan Africa: the need for the development of alternatives to allogenic blood. *J Blood Med.* 2011;2:7–21.
37. Jin T, Li PJ, Chen XZ, Hu WH. ABO blood group is a predictor of survival in patients with laryngeal cancer. *Chin J Cancer.* 2016;35(1):90.

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