

ABO and Rhesus Blood Group System in Tuberculosis Patients

Chukwurah Ejike Felix¹, Obeagu Emmanuel Ifeanyi^{2*}, Agom Daniel Dansy³

¹Department of Haematology and Immunology, Faculty of Clinical Medicine, Ebonyi State University, Abakaliki, Nigeria

²Department of Medical Laboratory Science, Faculty of Health Sciences, Imo State University, Owerri, Nigeria

³Department of Medical Laboratory Science, Faculty of Health Sciences, Ebonyi State University, Abakaliki, Nigeria

*Corresponding author: Obeagu Emmanuel Ifeanyi

| Received: 05.04.2019 | Accepted: 13.04.2019 | Published: 20.04.2019

DOI: [10.21276/sjmps.2019.5.4.1](https://doi.org/10.21276/sjmps.2019.5.4.1)

Abstract

This study on ABO and Rhesus blood group system in tuberculosis patients aimed at the assessment on prevalence of tuberculosis on human ABO and Rhesus blood group system. ABO blood and Rhesus blood group system antigens are hereditary characters that are the most clinically significant in blood transfusion. Tuberculosis is an infectious disease cause by mycobacterium tuberculosis and treated with antibiotics. The samples were collected, diagnosed and analyzed for reliable results using laboratory standard procedures, the results are represented statistically showing no significant effect(s) of tuberculosis (TB) in ABO and Rhesus blood system. This research was carried out among Igbo tribe at location of Ohaozara in Ebonyi State using two hundred (200) Tuberculosis patients coming for treatment at Tuberculosis/ Leprosy Presbyterian Joint Hospital Uburu (Happy Home) and other patient from the same location while three hundred (300) controls (people free from TB). The ABO blood group and Rhesus systems' result present blood group O predominated others and with 48%, followed by A (22%), B (18%) and AB (12%) while the Rhesus blood group system is 90% positive and 10% negative in tuberculosis patients. In comparison, ABO blood groups are; O (44%), A (28%), and AB (4%), while the Rhesus blood group system is 94% positive and 6% negative in controls. From the statistics above, among the patients in control from the same geographical location interprets that blood group O has the highest frequency of about 46 percent (46%) followed by group A with 25%, B (21%) and AB (8%) while Rhesus blood group system is 92% Rhesus positive and 8% Rhesus (D) negative in average among population study of the same location (Ohaozara). With the test (blood grouping) conducted among the population, the results show that no particular blood group is susceptible to Tuberculosis and so, Tuberculosis does not have any phenotypic effect(s) on ABO blood group and Rhesus blood group system.

Keywords: ABO blood group system, Rhesus blood group system, Tuberculosis patients.

Copyright @ 2019: This is an open-access article distributed under the terms of the Creative Commons Attribution license which permits unrestricted use, distribution, and reproduction in any medium for non-commercial use (NonCommercial, or CC-BY-NC) provided the original author and source are credited.

INTRODUCTION

There are many kinds of blood group system existing in different individuals but among all, the ABO and Rh blood group system are the most clinically important blood systems especially the ABO blood system followed by Rh blood group system, and they are the most blood group credited worldwide [1].

The ABO determines the group of individuals and the Rh determines whether the individuals group is positive or negative due to presence or absence of red blood cell D antigens [2].

The ABO blood system was discovered in 1900 at the University of Vienna by Karl Landsteiner who found three different blood types (A, B and O). The Rh blood system was discovered in 1940 by Karl Landsteiner and Wiener and the same great scientist and his student Wiener discovered the Rh blood system in 1940.

Karl Landsteiner during the discovery of ABO system recognized the presence of two separate erythrocyte antigens the A, B and O antigens. Expression of A and B genes in erythrocytes or body fluids results from the independently inheritance of A and B genes. The ABO locus resides on chromosome 9 in human, the H allele produces an enzyme that at the cellular level to construct the H antigen. This contribution to medical sciences was the most significant event in blood group systems research since the discovery of the ABO system. The associated and anti-B antibodies are usually IgM antibodies present in the serum, which are usually produced in the first year of life by sensitization to environmental substances such as bacteria, food, viruses [3].

The Rhesus blood system is probably the most complex of all erythrocyte blood group system with more than 40 different Rh antigens. Five major antigen

have been observed in the Rh system D, C, c, E, and e. C and c as well as E and e are antithetical refers to two antigens controlled by a pair of allelic genes. The presence of the Rh-antigen on an individual red blood cells means that the person is said to be Rhesus positive, while its absence is designated Rhesus negative [2].

The Rh gene resides on chromosome 1 with the D gene acting as an autosomal discrete unit (Wemer) or as haplotypes (Fisher and Race), (i.e. is HLA system; human leukocyte antigen system; four gene codes A, B, C and D for polymorphic proteins). The ABO and Rh system contain antibodies that can be stimulated as a result of presence of foreign body called antigen and this leads to haemolytic disease of the newborn (HDN). i.e. the human antibody in the subsequently reacts with the fetal antigens. This fetal antigen can induce maternal immunization by the formation of IgG antibodies, which will pass from the maternal circulation into fetal circulation through the placenta, in the fetal reaction with the fetal red blood cell leads to their destruction by the fetal reticuloendothelial system.

Since the discovery of ABO and Rh blood group system blood transfusion has been an easy and successful practice in medicine. And the rate of death as a result of transfusion reaction and rate of HDN occurrence had reduced [4].

Infectious diseases have played a significant role in altering the course of human populations in addition to bringing about change in their genetic constitutions. In Nigeria, tuberculosis remains the single major communicable disease affecting adults of great population suffering, a quarter being openly infectious; some percentage persons die each year with a far fewer number effectively detected and treated. While BCG (bacilli Calmette-Guérin) immunization protects from the most devastating forms of infantile tuberculosis (meningitis and disseminated tuberculosis), it does not appear to offer long term protection against the adult form of disease, which occurs from adult to child in millions of households throughout the country today.

Earlier studies have shown associations between some genetic markers viz; ABO blood group, HLA (human leukocyte antigen), serum proteins and tuberculosis both in India and abroad [5, 6]. However, information on various other red cell markers and disease is negligible in Indian populations. Therefore this pilot study is planned to fill this void by providing phenotype and allele frequency and allele frequency distributions of as many as different red blood cell genetic markers including two blood group systems (A1, A2 BO, Rh D), 4 red cell enzyme polymorphisms (ADA, AKL, ESD, PGM1) and haemoglobin (Hb)

variants in pulmonary tuberculosis patients and controls.

Pulmonary Tuberculosis

Tuberculosis (TB) is an infectious disease caused by bacteria whose scientific name is *Mycobacterium tuberculosis*. It was first isolated in 1882 by a German physician named Robert Koch who received a noble award on the discovery.

Pulmonary tuberculosis commonly affects the lungs but also can involve almost any organ of the body. Many years ago, this disease was referred to as "consumption" because, without effective treatment, these patients often would waste away (die). Today of course, tuberculosis usually can be treated successfully with antibiotics.

There also the group of organisms referred to as atypical tuberculosis there involve other types of bacteria that are in the *Mycobacteria* family. These organisms do not cause diseases and are referred to as "colonizers" because they simply live along side with other bacteria in our bodies without causing damage. At times, these bacteria can cause infections.

Aims and Objectives

- This project study is done to assess the prevalence of tuberculosis patients in ABO and Rhesus blood group system among the Igbo tribe.
- It was conducted to correlate/compare the results to the existing results of other scientists on the similar studies.
- This work was carried out in order to evaluate the causes, multiple risks, preventions and possible treatments for tuberculosis and its effects on ABO and Rhesus blood group system.

Materials and Samples

Data of sample collected from a total of 500 (five hundred) unrelated patients and controls (people free from TB) at random attending PJH (Presbyterian Joint Hospital) Uburu, PJH* Tuberculosis/Leprosy Uburu (Happy Home) and PJH Diagnostic Laboratory. All the subjects for the samples were between 20 and 65 years of age.

- Blood samples were collected at random from 200 (two hundred) patients of pulmonary tuberculosis and 300 (three hundred) persons free from TB (controls). From each subject, about 1-2mls of blood was drawn using 3mls sterilized syringe and collected into E.D.T.A.K2 anticoagulant container then to the hospital laboratory (PJH diagnostic Laboratory).

Procedures

- I sterilized all the containers and materials before used for all the practical.

- Following all the procedures from the labeled containers and detailed data, I made the materials ready before collecting the blood samples from the subjects (patients and controls).
- I carefully collected the blood samples from each of the subjects from a total of 500 (five hundred) unrelated TB patients and controls (free from TB) at age bracket of 20-65 years.
- By method of syringe, I collected blood samples randomly of about 2-3 ml with 3ml syringe and via heparinized tube containing EDTA K2 (Ethylene Diamine Tetra Acetic Acid potassium).
- I transported the samples immediately to the Haematology unit in Laboratory.
- I processed the sample for the preparation of haemolyses by freezing and thawing method.
- I confirmed ABO blood groups and Rhesus system of the samples by tube method using antisera. 8.1 worked out mathematically their percentages and frequencies in patients' and controls' samples. 9. I then interpreted my results in tabular form, pie chart and bar chart for understanding.

PROTOCOL

(1) Red Blood cells Grouping and Rhesus systems by Tube method.

Step-1: In three (3) small test tubes labeled A-C was pipetted into each tube as follows;

Tube A: 1 drop of anti-A serum

1 drop of 2-3% patient's or controls red cells.

Tube B: 1 drop of anti-B serum

1 drop of 2-3% patient's or controls red cells.

Tube C: 1 drop of anti-D serum

1 drop of 2-3% patient's or controls red cells.

Step 2: The contents in the above test tubes were gently and thoroughly mixed.

Step 3: The tests with the mixture were stood in test tube rack at room temperature for 3-5 minutes then, observed for agglutination or haemolysis under light.

Step 4: I confirmed the reactions on microscope by making a drop of the mixture on a grease free clean slide and view under microscope with x10 objective lens.

Step-5: Results

Group	Anti- A	Anti-B	Anti AB
A	+	-	+
B	-	+	+
AB	+	+	+
O	—	-	!

Rh (D) +Ve →	Anti- A Plus Red Cells	+
Rh (D) - Ve →	Anti- D Plus Red Cell	-

RESULTS

From the course of my study, I have observed that phenotypically, the distribution of blood groups, (ABO and Rhesus system) in the red cells of tuberculosis patients and controls (persons free from TB) are;

In the ABO blood group system, blood group O predominated in patients and controls with the highest frequency (48%) and (44%) respectively, followed by blood groups A (22% and A (28%), B

(18%) and B (24%), and AB (12%) and AB (4%) in patients and controls respectively as presented in Table 1 and 5 below .

In the rhesus blood group system, the frequency of Rh (D) negative was highest in patients having tuberculosis (10%) and (6%) in controls while their frequencies of Rh (D) positives were 90% and 94% in patients and controls respectively as presented in the Table 2 and 6 below.

Table-1: TB Patients' Result Data of Abo Blood Group and Rhesus System, In Tabular Form

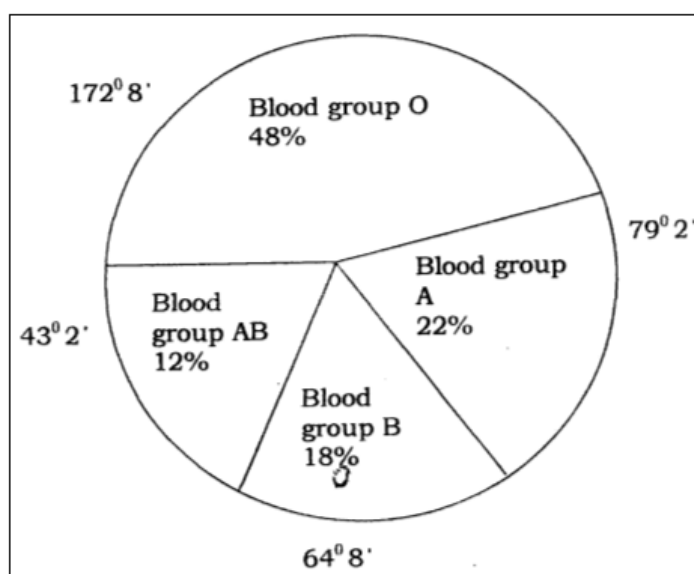
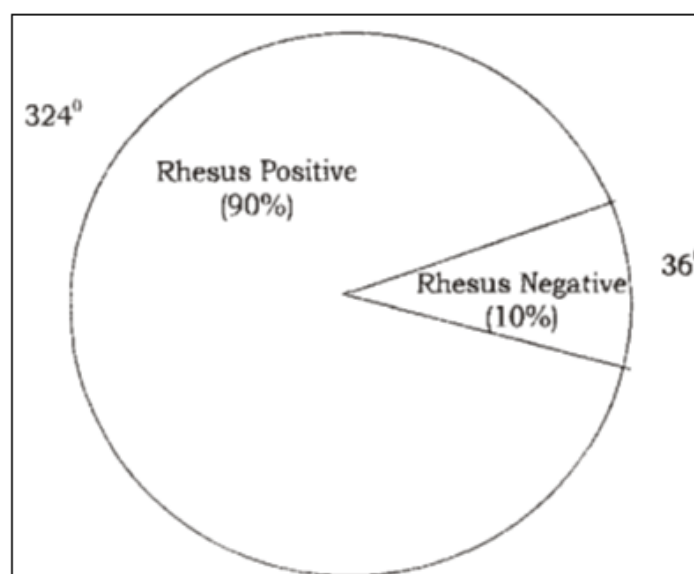
Blood Group	Rhesus (D) Factor	Frequency of TB Patients	Percentage
A	Positive	40	20%
A	Negative	4	2%
B	Positive	32	16%
B	Negative	4	2%
AB	Positive	20	10%
AB	Negative	4	2%
O	Positive	88	44%
O	Negative	8	4%
TOTAL		200	100%

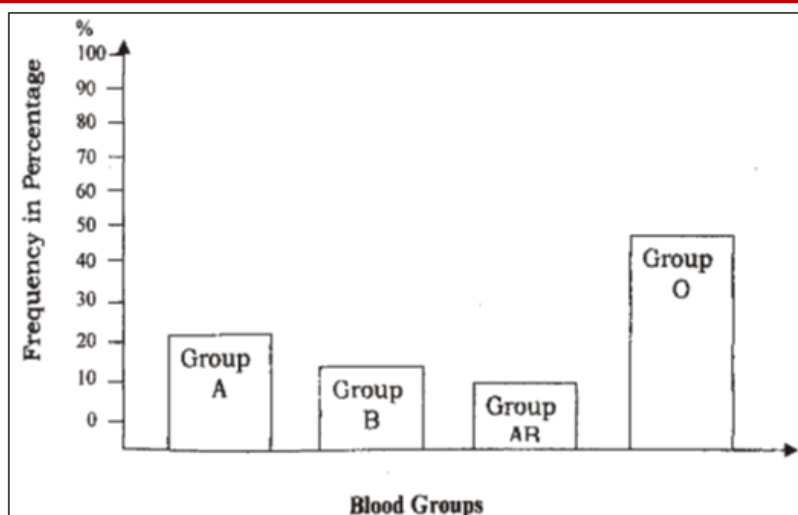
Table-2: ABO Blood Group of TB Patients

Blood Group	Frequency of TB Patients	Percentage Values
A	44	22%
B	36	18%
AB	24	12%
O	96	48%
TOTAL	200	100%

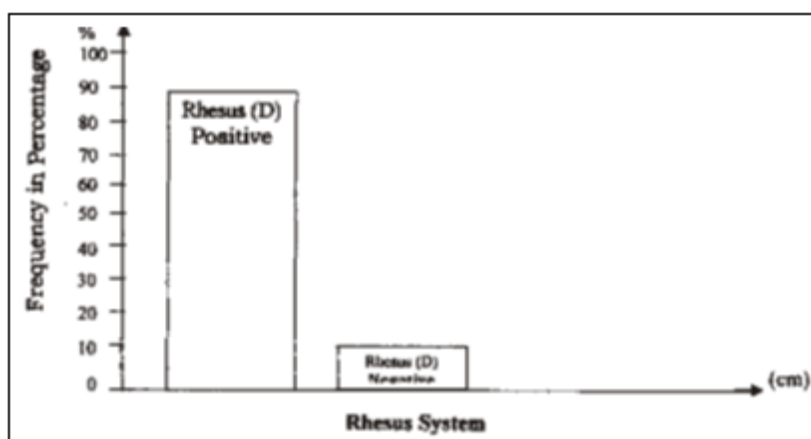
Table-3: Rhesus (D) Blood System of TB Patients

Rhesus (D) Blood System	Frequency of TB Patients	Percentage Value
Positive	180	90%
Negative	20	10%
TOTAL	200	100%

**TB Patients' Result Data of ABO Blood Group Presented In Pie Chart****TB Patients' Result Data of Rhesus (D) System Presented In Pie Chart**



TB Patients' Result Data of ABO Blood Group Presented in Bar Chart



TB Patients' Result Data of Rhesus System Presented In Bar Chart

Table-4: Controls' Result Data of ABO Blood Group and Rhesus System in Tablar Form

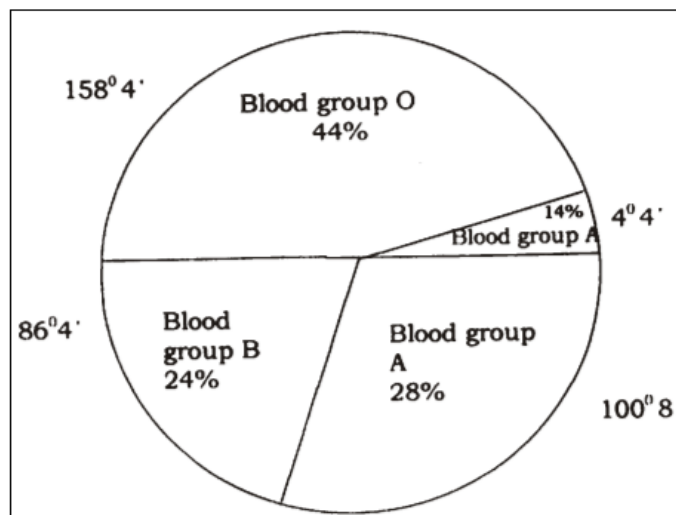
Blood Group	Rhesus (D) Factor	Frequency of TB Patients	Percentage
A	Positive	70	24%
A	Negative	0	0%
B	Positive	84	28%
B	Negative	0	0%
AB	Positive	12	4%
AB	Negative	0	0%
O	Positive	114	38%
O	Negative	18	6%
TOTAL		300	100%

Table-5: ABO Blood Droup of Controls

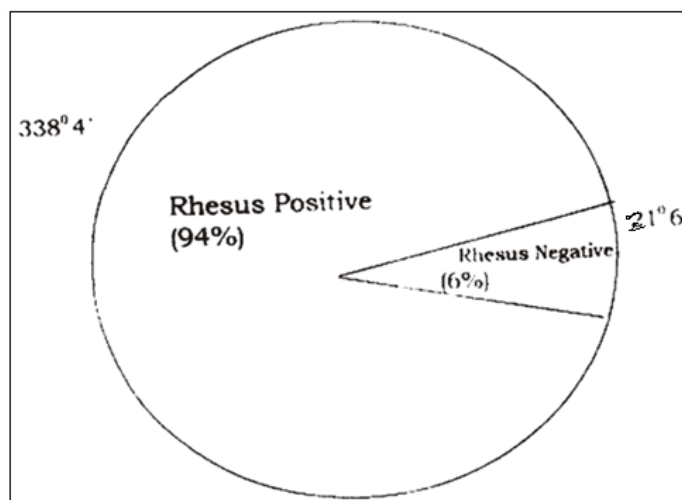
ABO Blood Group	Frequency of control	Percentage value
A	84	28%
B	72	24%
AB	12	4%
O	132	44%
TOTAL	300	100%

Table-6: Rhesus (D) Blood System of Controls

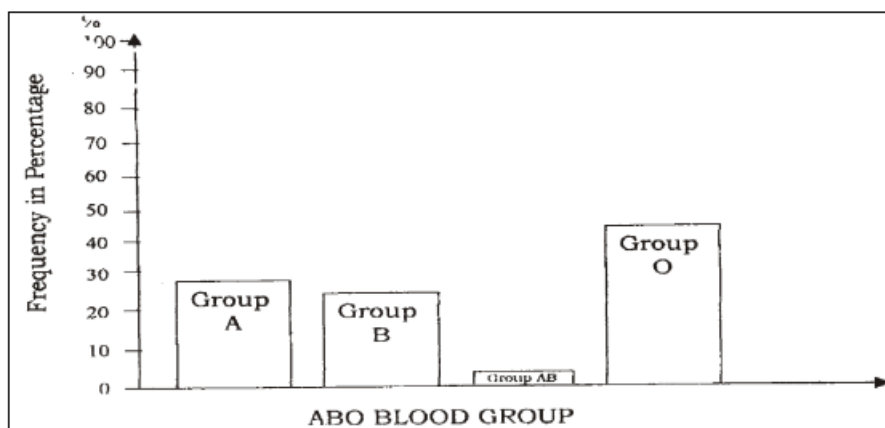
Rhesus (D) Blood System	Frequency of control	Percentage Value
Positive	282	94%
Negative	18	6%
TOTAL	300	100%

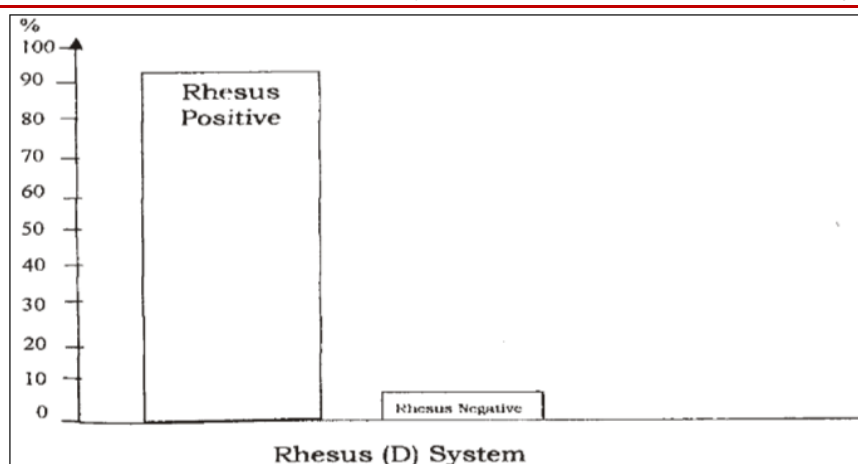


Controls' Result Data of ABO Blood Group Presented In Pie Chart



Controls' Result Data of Rhesus (D) System Presented In Pie Chart





Controls' Result Data of Rhesus (D) System Presented in Pie Chart

DISCUSSION

The research on ABO blood group and Rhesus system in TB (tuberculosis) patients at Eastern region of Nigeria and to be (individuals free of TB) was carried out in Tuberculosis/ Leprosy Presbyterian Joint Hospital (Happy Home, Uburu in Ohaozara L.G.A Ebonyi State of Nigeria. We observed that in the ABO blood group system, blood group O predominated in the Tb patients with the highest frequency (48%), followed by A (22%) B(18%) and AB (12%), while in controls (people free from TB) from the same Igbo tribe and location follow the same pattern of results from that of Tb patients. Blood group O was highest in frequency (44%), while A (28%, B (24%) and AB (4%).

The rhesus blood group system, the frequency of Rh (D) negative in patients was 10% and 6% in controls, while the frequencies of Rh (D) positive were 90% and 94% in patients and controls respectively.

This study was carried out with two hundred (200) Tb patients and three hundred (300) controls and there was no significant difference between the Tb patient and controls in the ABO and Rhesus blood groups.

Chessbrough reported his finding on the same study that blood group O (49%); A (26%), region are the average occurrence which is similar to the controls I obtained in my study outcome. Cheesbrough also discovered that rhesus (D) positive is of 94-95% [7].

Laha and Duta [8] found a very high frequency of blood group O in pulmonary tuberculosis patients while Bhosale and kulkani [9] observed that group A showed the highest frequency (36%) in the case of TB patients of Bombay.

Reddy *et al.*, [10] considering the proportions of ascariasis, cataract, conical ulcer and tuberculosis occurring in Badaga patients found that 47.65% of them

belonged to the O blood group as compared to 31.25% of controls.

Kshatriya and Kapoor [11], observed appreciably high frequency of blood group B and Rhesus factor D negative in tuberculosis patients.

Maror and Bhanwer [12] observed higher incidence of pulmonary tuberculosis among blood group AB and B individuals.

Jain [13] found a significantly higher incidence of group AB among pulmonary tuberculosis patients.

In other hands several studies reported in literature [14-18] observed no significant difference between the blood groups of TB patients and person free from TB (Controls).

CONCLUSION

The haematological, Serological and biochemical studies on the effects tuberculosis in ABO blood groups and rhesus system of the patients have apparently not given a much significant evidence for phenotypic association with tuberculosis.

Tuberculosis (TB) to adopt the preventing measures of drug abuse and individuals free from Tuberculosis to adopt the preventive measures from contacting mycobacterium organisms that causes Tuberculosis and still maintain good body immunity from balance die.

REFERENCES

1. Obeagu, E. I. (2019). An update on susceptibility of individuals to diseases based on. *International Journal of Current Research in Medical Sciences*, 5(3): 1-8.
2. Moore, S., & Gahmberg, C. G. (1997). Identification of Rh polypeptide and Rh

- polypeptide/Rh glycoprotein complexes. *Biotest Bull*, 5, 409-413.
3. Obeagu, E. I., Ogbodo, O. R., Onyenweaku, F., Emelike, C. U., & Udochukwu, A. I. (2013). Frequency distribution of ABO, Rh blood groups and blood genotypes among the students and staff of Michael Okpara University of Agriculture, Umudike Abia State, Nigeria. *Int J Res Rev Pharm Appl Sci*, 3(4), 561-565.
4. Tricot, F., Crozat, Y., & Pellerin, S. (1997). Root system growth and nodule establishment on pea (*Pisum sativum* L.). *Journal of Experimental Botany*, 48(11), 1935-1941.
5. Mourant, A. E., Kopec, A. C., & Domaniewska-Sobczak, K. (1978). *Blood groups and diseases. A study of associations of diseases with blood groups and other polymorphisms*. Oxford University Press, Walton Street, Oxford OX2 6DP.
6. Bhasin, M. K., & Chahal, S. M. S. (1996). *A laboratory manual for human blood analysis*. Delhi: Kamla-Raj Enterprises, Delhi, 21: 51-59.
7. Cheesbrough, M. (2001). District laboratory practice in tropical countries part 2 published by press syndicate of the University of Cambridge, 362-369.
8. Laha, P. N., & Dutta, M. (1963). Association between blood group and pulmonary tuberculosis. *Journal Ass. Phs India*, 11: 287-288.
9. Bhosale, N., & Kulkarni, V. S. (2003). ABO blood group and some diseases -A correlation study of Maharashtrais man in India, 57: 164-169.
10. Reddy, V. R., Ramamohan, K., & Reddy, P. G. (1978). ABO blood groups and diseases among the Badages and Tadas of the Nitgiris, Douth India. Vth Ann. Conf. Ind. Soc. Hum. Genet.
11. Kshatriya, G. K., & Kapoor, A. K. (1991). Some observations on blood group in relation to pulmonary tuberculosis. *Indian Anthropologist*, 21: 17-21.
12. Marok, G. K., & Bhanwer, A. J. S. (1989). Excess of blood group AB and B in pulmonary tuberculosis. *Ind. J. Phys. Anthropol. Hum. Genet*, 15, 93-96.
13. Jain, R. C. (1970). ABO blood groups and pulmonary tuberculosis. *Tubercle*, 51(3), 322-323.
14. Kothare, S. N. (1959). ABO blood group in relation to pulmonary tuberculosis. A preliminary report. *Journal of postgrad Med*, 5: 94-98.
15. Navani, H., & Narang, R. K. (1962). A study of ABO blood groups in pulmonary tuberculosis. *Ind. Jour. Chest. Dis.*, 4, 109-113.
16. Nath, K., Jolly, J. G., & Parasha, S. K., (1963). Blood groups and susceptibility in disease. *Journal Assoc physician Indian*, 11; 567-671.
17. Saha, N., & Banerjee, B. (1968). Incidence of ABO and RH blood groups in pulmonary tuberculosis in different ethnic groups. *Journal of medical genetics*, 5(4), 306-307.
18. Sidhu, L. S., Singh, J., Bhatnagar, D. P., & Pahuja, J. K. (1974). Association of pulmonary tuberculosis with ABO and Rh (D) blood groups. *Human Population Genetics In India*, 135-140.