Original Article





Website: www.ajts.org DOI: 10.4103/ajts.AJTS_15_17

Evaluation of serum natural autoantibodies reaction in different hematological disorders with prospective view to their probable utilization in predictive medicine

Maryam Mosaed, Ali Akbar Pourfathollah¹, Mostafa Moghadam², Mir Hadi Jazayeri³, Amir Reza Safdarian^₄

Abstract:

BACKGROUND: There are some antibodies which are present in healthy individuals without any former exposure to foreign antigens; they are known as natural autoantibodies (NAAbs). In recent years, it was shown that they probably contribute to the homeostasis of the whole body and might be present before beginning of some diseases. Thus, as new biomarkers, they are promising factors to diagnose diseases.

MATERIALS AND METHODS: In this study, we drew upon samples of 924 individuals (600 controls and 324 cases) with underlying diseases of anemia, polycythemia, leukocytosis, thrombocytopenia, thrombocytosis, and pancytopenia. For detection of NAAbs against red blood cell, plasma samples were incubated with their own red cell suspension in 4°C for 18 h. Then, positive samples were evaluated for antibody screening and titration.

RESULTS: Fifty-two (8.6%) controls and 58 (17.9%) cases showed positive reaction (Pv < 0.001). The prevalence of positive antibody screens among auto-positive controls was 53% and 100% among cases; moreover, strength of antibody screen reaction had a mean rank of 22.5 in controls and a mean rank of 38.5 in cases (Pv < 0.001). A significant relation was also observed between ABO blood group and prevalence of NAAbs in controls but not in cases (Pv < 0.05).

CONCLUSION: The prevalence and potency of NAAbs increased along with hematological changes; moreover, the antibody reactions' pattern and titration showed significant differences between the two groups and these may be useful as biomarker for monitoring and prediction of some hematological diseases.

Keywords:

Hematological disorders, natural autoantibodies, predictive medicine

Introduction

It is well established that natural antibodies (NAbs) are present in healthy.Individuals without any former deliberate immunization, but generally, are able to react with a widespread range of antigens from selfmolecules (e.g., phospholipids, serum

This is an open access journal, and articles are distributed under the terms of the Creative Commons Attribution-NonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms. proteins, cellular components, insulin, and thyroglobulin) to exogenous antigens.^[1,2] For a long time, NAbs had been considered as an unimportant part of immunity, but recently, more research on NAbs has clarified surprising and even controversial physiologic and pathological aspects. When for the first time, Grabar, in 1975, suggested a physiologic function for NAbs, clearance of cell debris, has

How to cite this article: Mosaed M, Pourfathollah AA, Moghadam M, Jazayeri MH, Safdarian AR. Evaluation of serum natural autoantibodies reaction in different hematological disorders with prospective view to their probable utilization in predictive medicine. Asian J Transfus Sci 2020;14:167-71.

Iran Blood Transfusion Research Center, Tarbiat Modares University, ¹Tarbiat Modares University, Iranian Blood Transfusion Organization, ³Department of Immunology, Iran University of Medical Sciences, ⁴Department of Immunology, Medical Faculty, Tarbiat Modares University, ²Iran Blood Transfusion Research Center, Tehran, Iran

Address for correspondence:

Dr. Ali Akbar Pourfathollah, Tarbiat Modares University, Iranian Blood Transfusion Organization, P. O. Box: 14665-1157, Tehran, Iran. E-mail: pourfa@ibto.ir

Submission: 02-02-2017 Accepted: 24-11-2017 Published: 19-12-2020 substantially put attention to these multifunctional Abs, and until now broad studies have indicated significant roles in not only protection against exogenous antigens and facilitating apoptosis but also in atherosclerosis, neurodegenerative disorder, and malignancy, that gave rise to present knowledge in their necessary role in maintenance of immune homeostasis.^[3,4] Nowadays, their roles have been clarified including cancer-specific apoptosis, NAbs against phosphorylcholine as a strong protection marker in atherosclerosis, DC differentiation, and inflammation control mediated by antileukocyte NAbs.^[5-9] It has been also suggested that polyreactivity of NAbs probably gives them a beneficial property that leads to formation of a massive, dynamic network controlling whole body homeostasis which in turn actively causes physiologic autoimmunity.^[10] This intellectual system that is permanently affected by internal and external stimulus, purposefully prevents stimulation of pathologic autoimmunity.^[4,11] Accordingly, a hypothesis, termed"Immunological Homunculus." considered natural auto antibodies as the mirror of organism.^[12]This point of view proposed that any excessive changes in physiologic condition of body in terms of quantity and intensity will affect the NAbs content. Then more accomplished results based on more observation like microarray test (antigenchip) have caused some of these NAbs to be considered as "biomarkers". So they can prepare an early diagnosis possibility of "predictive medicine" at the early stage of disease, making an opportunity for identification of pathologic changes in the body.^[13-16] My colleagues have previously shown that the mean immune reactivity of serum antibodies with 24 human autoantigens increased with advancing age, which could be because of the increase in the level of pathological antibodies.^[2] Here, we utilize immune hematology methods for NAbs screening against red blood cells (RBCs) surface antigens under physiological changes of blood in two populations, healthy blood donors (controls) and hematological patients' group (cases).

Materials and Methods

Study design and sample preparation

The study population included the total number of 924 individuals (600 controls and 324 cases) including relative polycythemia, Hb: >18 g/dl, n = 131; anemia, Hb: <10 g/dl, n = 101; leukocytosis, white blood cell >20000 mcl, n = 47; thrombocytopenia, plt: <100,000 mcl, n = 21; thrombocytosis, plt: >450,000, n = 14; and pancytopenia, n = 12. Criteria for selecting this number of samples were based on achievement to an equal number of auto-positive samples in two groups. Blood specimens were obtained from 600 normal individuals and 324 cases into two groups (total 924). Blood samples were collected under the supervision of a physician. Ethylenediaminetetraacetic acid blood

samples of controls were collected from March 2015 to November 2015 in Tehran and Karaj Blood Donor Centers. The cases' samples were collected in Iran Blood Transfusion Organization (IBTO) Hematology Reference Laboratory. The blood donors were informed about the study aim and procedure, and written consent was obtained by the medical interviewer. All the protocols were confirmed by the Ethical Committee of Medical Faculty of Tarbiat Modares University.

Since we were looking for cold reactive natural IgM antibodies with the inability of complement activation, instead of DAT test, all samples with overnight incubation at 4°C underwent a test for natural autoantibodies; then for samples which had shown positive reaction at first stage supplementary tests including antibody screening and specificity test were performed. Standard serologic procedures and manufacturer's directions were followed as described in the AABB Technical Manual.^[17] ABO and Rh(D) blood grouping was performed with reagents purchased from Diagast (251, Av. E. Avinee-BP. 9 59374 Loos France). The antibody screening test was performed with three cells' reagent by tube testing at 4°C in saline media. The tests of antibody specificity were performed using Oladult, Oicord, autologous, and ABO compatible cells.^[17,18] All different cells including O cells, panel cells, cord blood Oi cells, and A1 and B cells were obtained from IBTO Immunohematology Reference Laboratory. To ensure, in the case group, the prewarm indirect antiglobulin test (IAT) was performed with polyspecific anti-immunoglobulin G + Anti-C3d and enhancing agent (LISS), any positive reaction at 37°C led to the removal of the sample. Anti-C3d + Anti IgG (polyspecific) antiglobulin reagents were also obtained from Diagast. Antibody titrations were performed with serial dilutions of the serum in physiologic saline at the same temperature as the antibody screening tests.^[19]

Statistics

Computer software SPSS 16 (IBM, SPSS Inc., Version 16.0. Chicago) and also GraphPad Prism (version: 5.01, GraphPad Software, Inc., USA) were used for data analysis. Analysis of blood groups' prevalence in positive samples was carried out using Chi-Square; moreover, Mann–Whitney and binominal tests respectively were carried out to measure mean rank of antibody screening results and natural autoantibodies' (NAAbs) prevalence rate in different groups. P < 0.05 was interpreted as statistically significant.

Result

Frequency and titration of natural autoantibodies Both prevalence and titration of NAAbs increased among case group; moreover, binomial analysis showed that intended ratio of NAAbs in control group significantly differed from case group (P = 0.001).

Potency of natural autoantibodies reaction

According to antibody screening results, the strength of NAAbs reaction in case group was significantly higher than normal population (P = 0.001) [Figure 1]. Moreover, the number of samples with NAAbs in case group was higher than in the controls [Table 1]. In antibody screening for case and control groups of this study, the potency of antibody in control group was very low and not enough for titration, but in case group, the potency of antibody was very higher even up to 1024-fold titration and also as has been showed in Figure 1. Pan reactivity of the patient samples was significantly increased compare to control group.

Incidence of natural autoantibodies reaction in different hematological disorders

A significant relationship was observed between the incidence of reaction and type of disease in most of the abnormal cases as it has been shown in Table 2; also the highest percent of reactions was observed in pancytopenia both in terms of strength and incidence rate.

Relation between natural autoantibodies and ABO blood group

Among whole positive samples in control group, blood groups were as follows: A = 46.1% (24 cases), O = 23% (12 cases), B = 21% (11 cases), and AB = 9.9% (5 cases). Distribution of ABO blood group in case group compare to control was not significant;^[20] but in control group, the A blood group showed a significant increase in compare the normal population (P = 0.01).

Table 1: The cold reaction of red blood cells with self-serum significantly increased in case group compare to control and subsequently in alloreactivity and titration

Auto check	Control, <i>n</i> (%)	Case, <i>n</i> (%)	
Total	600	324	
Auto-positive	52 (8.6)	58 (17.9)	
Auto-negative	548 (91.4)	266 (82.0)	
Ab screening and titration (n=30)*			
Screen positive	16 (53)	30 (100)	
Screen negative	14 (47)	0	
Titration	All=2	2-1024	

*Only those samples were selected for Ab screening which had at least a minimum + 1 reaction

Discussion

In this study, we investigated sera for NAAbs reaction in 924 individuals in two populations of healthy blood donors as controls and hematologic patients as cases. NAAbs are the capacity of immune system to respond to foreign antigens. It has been shown that the amount of these antibodies increases with age and responsible for the increased susceptibility of the elderly to infection and cancer.^[21] The CD5+ B-1 cells are responsible for the secretion of natural autoantibodies with no stimulation from the early stages of embryogenesis,^[22] In our previous study, we demonstrated that there is association between human B-1 cell frequency and aging.^[23] As well as reactivity of natural autoantibody with auto-antigens changed in different age intervals among healthy individuals and increased in the elderly people. Regarding the crucial roles of natural autoantibodies in maintaining the body homeostasis, these variations might represent the immunological and pathological condition in the elderly.^[2] Concentration of total IgG and IgM was determined in different age intervals. Concentration of IgG has been shown increasing trend along with age. On the contrary, the mean value of total IgM showed an increasing pattern from cord blood to the middle age (40 years) after which it dropped, while, as noted above, IgG concentration followed an increasing trend all the way from infants to the elderly. The decrease in IgM level showed positive

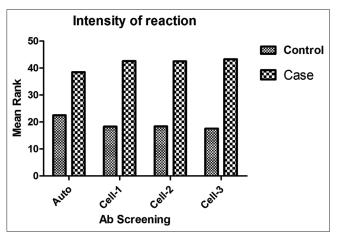


Figure 1: Strength of antibody screen reaction had a mean rank of = 22.5 in controls and a mean rank of = 38.5 in cases (Pv = 0.001)

Table 2: Distribution of positive reaction in different hematopathological conditions, as shown in table, the most association was found with pancytopenia and the least with thrombocytopenia

	Pancytopenia, n (%)	Trombocytopenia, n (%)	Trombocytose, n (%)	Leucocytose, n (%)	Anemia, <i>n</i> (%)	Polycytemia, n (%)
Total	12 (100)	21 (100)	14 (100)	47 (100)	101 (100)	131 (100)
Positive reaction	6 (50)	8 (38)	1 (7)	12 (25.5)	11 (10.8)	20 (15.2)
Negative reaction	6 (50)	13 (62)	13 (93)	35 (74.5)	90 (89.2)	111 (84.8)
Р	<0.001	<0.001	>0.05	<0.001	>0.05	<0.001

association with the frequency susceptibility of the elderly to autoimmune disease and cancer.^[24] Thus, as expected, we apparently detected the increased levels of NAAbs at least in a part of healthy population that had exceeded micromolar level. Therefore, on the one hand, it can be said that one of the reasons for success of immune system largely depends on proper repertoires of NAAbs against carbohydrates that stand for their various and essential function in all tissues,^[25] but on the other hand, affecting internal and external stimuli and regulation mechanisms should be considered although they are little known.^[11] Based on prior research, a blood group shows more heterogeneity, subgroups even show more genes than B blood groups. Therefore, it likely represents which more anti-A reflects variety of heterogeneous antigens.^[25,26] It should be noted that the term "polyreactivity" of NAAbs does not mean lack of specificity rather a tendency to a set of particular epitopes that give them so many essentially biological roles.^[27] According to observation following immunoculus theory, NAAbs are reflectors of body inner picture in a molecular scale, and if any alteration for any reasons occurs in body condition, it will be reflected in the NAAbs content.[12-16] RBCs transfusion in patients can lead to the remodeling of NAAbs autoreactivity.^[28] NAAbs are able to identify Damage-associated molecular patterns (DAMPs) and in fact as sensors play a protective role in immunity against oxidative epitopes.^[29] Furthermore, it has been shown that NAAbs regulate immune system and a large number of NAbs have a key role in tissue homeostasis. Immunosurveillance mechanism has a role in cancer prevention and the presence of NAbs in normal range/activity in healthy individuals is essential for body to maintain hemostasis.[30]

In the present study, we were able to demonstrate that the reaction of sera NAAbs both in terms of incidence and intensity or potency that increases along with alteration in physiologic and pathologic changes depends on the related tissue (blood) and the most alterations were related to the group with all blood cells disorder. Exploring the fact hidden behind NAAbs as a boundary among physiology and pathology serves us to understand a part of this complex network.

Acknowledgments

This study was supported by IBTO. The work is part of the master thesis of Maryam Mosaed.

Financial support and sponsorship Nil.

Conflicts of interest

There are no conflicts of interest.

References

- 1. Elkon K, Casali P. Nature and functions of autoantibodies. Nat Clin Pract Rheumatol 2008;4:491-8.
- 2. Jazayeri MH, Pourfathollah AA, Rasaee MJ, Farhadi M, Zarei N, Jafari ME. The reactivity of human serum natural autoantibodies with certain autoantigens increases along with aging. Biomed Aging Pathol 2013;3:115-8.
- 3. Rahyab AS, Alam A, Kapoor A, Zhang M. Natural antibody Biochemistry and functions. Glob J Biochem 2011;2:283-8.
- 4. Lacroix-Desmazes S, Kaveri SV, Mouthon L, Ayouba A, Malanchère E, Coutinho A, *et al.* Self-reactive antibodies (natural autoantibodies) in healthy individuals. J Immunol Methods 1998;216:117-37.
- 5. Vollmers HP, Brändlein S. Natural antibodies and cancer. J Autoimmun 2007;29:295-302.
- Kyaw T, Tipping P, Bobik A, Toh BH. Protective role of natural IgM-producing B1a cells in atherosclerosis. Trends Cardiovasc Med 2012;22:48-53.
- Fiskesund R, Su J, Bulatovic I, Vikström M, de Faire U, Frostegård J, et al. IgM phosphorylcholine antibodies inhibit cell death and constitute a strong protection marker for atherosclerosis development, particularly in combination with other auto-antibodies against modified LDL. Results Immunol 2012;2:13-8.
- Bayry J, Lacroix-Desmazes S, Donkova-Petrini V, Carbonneil C, Misra N, Lepelletier Y, *et al.* Natural antibodies sustain differentiation and maturation of human dendritic cells. Proc Natl Acad Sci U S A 2004;101:14210-5.
- Lobo PI, Bajwa A, Schlegel KH, Vengal J, Lee SJ, Huang L, et al. Natural IgM anti-leukocyte autoantibodies attenuate excess inflammation mediated by innate and adaptive immune mechanisms involving Th-17. J Immunol 2012;188:1675-85.
- Lobo PI, Schlegal KH, Vengal J, Okusa MD, Pei H. Naturally occurring IgM anti-leukocyte autoantibodies inhibit T-cell activation and chemotaxis. J Clin Immunol 2010;30 Suppl 1:S31-6.
- 11. Avrameas S, Ternynck T, Tsonis IA, Lymberi P. Naturally occurring B-cell autoreactivity: A critical overview. J Autoimmun 2007;29:213-8.
- Poletaev A, Osipenko L. General network of natural autoantibodies as immunological homunculus (Immunculus). Autoimmun Rev 2003;2:264-71.
- Poletaev AB. The immunological homunculus (immunculus) in normal state and pathology. Biochemistry (Mosc) 2002;67:600-8.
- 14. Cohen IR. Biomarkers, self-antigens and the immunological homunculus. J Autoimmun 2007;29:246-9.
- Quintana FJ, Merbl Y, Sahar E, Domany E, Cohen IR. Antigen-chip technology for accessing global information about the state of the body. Lupus 2006;15:428-30.
- Poletaev AB, Churilov LP, Stroev YI, Agapov MM. Immunophysiology versus immunopathology: Natural autoimmunity in human health and disease. Pathophysiology 2012;19:221-31.
- Blaney KD, Howard PR. Basic & Applied Concepts of Immunohematology. Missouri, USA: Mosby Elsevier; 2009.
- Roback JD, Combs R, Grossman J, Hillyer C. AABB Technical Manual. Bethesda, MD: American Association of Blood Banks (AABB); 2008.
- Campbell SA, Shirey RS, King KE, Ness PM. An acute hemolytic transfusion reaction due to anti-IH in a patient with sickle cell disease. Transfusion 2000;40:828-31.
- 20. Pourfathollah A, Oody A, Honarkaran N. Geographical distribution of ABO and Rh (D) blood groups among Iranian blood donors in the year 1361 (1982) as compared with that of the year 1380 (2001). Sci J Iran Blood Transfus Organ 2004;1:11-7.

- Attanasio R, Brasky KM, Robbins SH, Jayashankar L, Nash RJ, Butler TM, et al. Age-related autoantibody production in a nonhuman primate model. Clin Exp Immunol 2001;123:361-5.
- 22. Baumgarth N, Tung JW, Herzenberg LA. Inherent specificities in natural antibodies: A key to immune defense against pathogen invasion. Springer Semin Immunopathol 2005;26:347-62.
- 23. Jazayeri MH, Pourfathollah AA, Jafari ME, Rasaee MJ, Dargahi ZV. The association between human B-1 cell frequency and aging: From cord blood to the elderly. Biomed Aging Pathol 2013;3:20-2.
- 24. Jazayeri MH, Pourfathollah AA, Rasaee MJ, Porpak Z, Jafari ME. The concentration of total serum IgG and IgM in sera of healthy individuals varies at different age intervals. Biomed Aging Pathol 2013;3:241-5.
- Bovin N, Obukhova P, Shilova N, Rapoport E, Popova I, Navakouski M, *et al.* Repertoire of human natural anti-glycan immunoglobulins. Do we have auto-antibodies? Biochim Biophys Acta 2012;1820:1373-82.

- Huflejt ME, Vuskovic M, Vasiliu D, Xu H, Obukhova P, Shilova N, *et al.* Anti-carbohydrate antibodies of normal sera: Findings, surprises and challenges. Mol Immunol 2009;46:3037-49.
- 27. Avrameas S, Ternynck T. The natural autoantibodies system: Between hypotheses and facts. Mol Immunol 1993;30:1133-42.
- Stahl D, Lacroix-Desmazes S, Sibrowski W, Kazatchkine MD, Kaveri SV. Red blood cell transfusions are associated with alterations in self-reactive antibody repertoires of plasma IgM and IgG, independent of the presence of a specific immune response toward RBC antigens. Clin Immunol 2002;105:25-35.
- 29. Uchida K. Natural antibodies as a sensor of electronegative damage-associated molecular patterns (DAMPs). Free Radic Biol Med 2014;72:156-61.
- Ebrahimnezhad S, Jazayeri M, Hassanian SM, Avan A. Current status and prospective regarding the therapeutic potential of natural autoantibodies in cancer therapy. J Cell Physiol 2017;232:2649-52.