Research DOI: 10.6003/jtad.1591a1

The Relationship Between ABO Blood Groups and Psoriasis Vulgaris

Belma Türsen,¹ MD, Erdinç Terzi,² MD, Bilal Bulut,³ MD, Ümit Türsen,^{3*} MD, Teoman Erdem,⁴ MD

Address: ¹Mersin State Hospital, ²Yenikent State Hospital, Department of Dermatology, ³Mersin University, ⁴Sakarya University, School of Medicine, Department of Dermatology

E-mail: utursen@mersin.edu.tr

* Corresponding Author: Dr. Ümit Türsen, Mersin University, School of Medicine, Department of Dermatology Mersin, Turkey

Published:

J Turk Acad Dermatol 2015; **9 (1)**: 1591a1.

This article is available from: http://www.jtad.org/2015/1/jtad1591a1.pdf

Keywords: ABO, blood groups, psoriasis

Abstract

Background: Studies of associations between various cancers and the ABO blood groups have shown elevated relative risks for some categories of disease. To date, no report has evaluated the relationship between the ABO blood groups and the psoriasis.

Material and Methods: We conducted a retrospective study of psoriasis diagnosed in Turkey. All cases were clinically and histopathologically confirmed. Blood information was obtained for 129 individuals with psoriasis, and the distribution of ABO and Rh blood type for cases was compared with that of 419 healthy blood donors from the same geographic area.

Results: Patient group A and AB blood group was higher than the control group, O and B blood groups lower than the control group. There was no statistically significant differences between the two groups (p = 0.263). The patient group and control group statistically any significant differences were found between the distribution of Rh factor.

Conclusion: Our study shows some association of AB and O blood groups with psoriasis. Further studies in larger series on blood group antigens are needed to elucidate the relationship between these antigens and psoriasis.

Introduction

Psoriasis vulgaris, which is characterized by sharply demarcated erythematous scaling plaques, the reason for the unknown, a chronic inflammatory disease [1,2]. The etiopathogenesis of diseases, despite being one of the most studied to date, is fully unclear. Psoriasis is considered to be a genetically programmed disease of dysregulated inflammation, which is driven and maintained by multiple components of the immune system. The recent literature supports the hypothesis of multifactorial inheritance [1,3,4,5]. In hu-

mans, the major blood group antigens are located on the surface of red blood cells and various epithelial cells.

The relationship with blood groups had been studied in many cancers such as esophagus, cardiac, gastric, lung, laryngeal, hypopharyngeal, salivary gland, gynecologic, colorectal, pancreatic, bone, urinary bladder, ureter, renal, breast, prostate, testicular tumors and uveal melanoma [6,7,8,9,10,11,12,13,14,15,16,17,18,19,20]. There are publications that evaluated the relationship between blood groups and skin

diseases such as vitiligo, pemphigus vulgaris, discoid lupus eritematosus, oral lichen planus and skin tumors [21,22,23,24,25,26,27,28,29,30,31,32]. To date, the literature investigation the relationship between ABO blood groups and psoriasis vulgaris were found in any publication. In this study, a retrospective evaluation was performed to determine the relationship between blood groups with psoriasis vulgaris.

Materials and Methods

In our study group of patients diagnosed with psoriasis vulgaris 66 (51,2%) men and 63 (48,8%) women with a total of 129 patients, in the control group and 303 (72.3%) were male and 116 (27.7%) women were 419 healthy blood donors. Routine blood examination was performed in all patients and controls. The control group, cardiovascular disease, cancer, chronic degenerative neurological disease, chronic obstructive pulmonary disease, hepatitis, allergic diseases, and were selected from among healthy people without a history of alcohol dependence.

Blood samples were obtained into vacuum tubes containing EDTA (vacutainer, Becton Dickinsen, Marseilles, France) from each donor's venous circulation. ABO and Rh blood typing were carried out with tube method and gel method.

Tube method: One drop of anti-A, anti-B, or anti-D (Eryclone, Tulip Diagnostics, Bambolim, India) was added to the appropriately labeled tube. A 5 percent suspension of red blood cells (RBC) was made in isotonic saline. One drop was added to tubes containing anti-A, anti-B, or anti-D. The contests of the tubes were mixed thoroughly, and the tubes were centrifuged for 20 seconds at 3400 rpm. Tubes were read macroscopically for agglutination.

Gel method: A 5 percent RBC suspension was prepared in diluent (modified bromelin solution for red cell suspensions). Gel cards (Diaclon ID, Diamed AG, Cressier, Switzerland) were used for ABO and Rh typing. 10 µL of RBC suspension was added to the gel microtubes containing anti-A, anti-B, anti-D, and control reagents, respectively. 50 µL of donor plasma were added to microtubes for reverse ABO group testing. The ID cards were centrifuged at 895 rpm 10 minutes in the centrifuges (ID-centrifuge). A positive reaction (4+) was determined by the formation of a red line on the gel surface, whereas intermediate reactions were characterized by red agglutinates distributed throughout the gel. With a negative reaction, a compact button of cells formed on the bottom of the microtube.

The findings of this study evaluated, for statistical analyzes and NCSS (Number Cruncher Statistical System) Statistical Software 2007 & PASS 2008 (Utah, USA) was used. Descriptive statistical methods for evaluating the study data, as well as Student's t test was used to compare quantitative data between groups. Qualitative comparisons of the data in the chi-square test and Fisher exact Chi-square test was used. Statistical significance at p <0.05 level were evaluated.

Results

The mean age of patients was 42.24 ± 14.61 . In the control group mean age was 53 ± 6.2 . The mean age of patients a statistically significantly higher than the control group (p <0.01). Patient group A, B, O and AB blood groups ABO, 55 (42.6%), 19 (14.7%), 47 (36.4%) and 8 (6.3%), respectively. In the control group, ABO blood groups, respectively ,147 (35.1%), 66 (15.7%), 188 (44.9%) and 18 (4.3%) were detected. Patient group A and AB blood group was higher than the control group, O and B blood group lower than the control group. There was no statistically significant differences between the two groups (p = 0.263). The patient group and control group statistically any significant differences were found between the distribution of Rh factor.

Discussion

Psoriasis vulgaris is one of the most common skin diseases. Although the pathogenesis of psoriasis is stil unclear, many studies suggest that immune and hereditary mechanisms may play an important role [1,3,5]. There are publications that evaluated the relationship between blood groups and skin diseases such as vitiligo, pemphigus vulgaris, discoid lupus erythematosus, lichen planus and malignant skin tumors [21,22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32]. Antigenic expression of genes or genetic instability due to the separation from each other and contribute to the production of antibodies because of the different types of diseases associated with blood group [26]. There was no publication in the literature, assessing the relationship between ABO blood groups and psoriasis vulgaris.

In this study, the rate of patient groups A and AB blood group is higher than the control group and the O and B blood group determined that the rate lower than control group. However, we did not observe any sta-

tistically significant differences between the two groups. Antigen A, more patients in this study were found in the erythrocytes of this antigen alone, but also the superficial tissues, such as leather carrying nuclear cells are also available [26]. A blood group B antigen in the form of people isoaglutinin cell membrane protein expression and antibody production may contribute to a hyperproliferative disease such as psoriasis can lead to the development of the phenotype.

ABO blood group genes are map at 9q in which the genetic alteration is common in many skin diseases [26]. Thus, ABO blood group antigen expression may be effected by the genetic change of dermatoses [22]. On the other hand, it is possible the observed associations are not due to the blood group antigens themselves, but to the effects of genes closely associated with them. Additionally it might have nothing to do with molecular mechanisms or genetics. It is merely the result of population history, environment, diet and customs [22].

As a result, with psoriasis vulgaris in our study was observed any statistically significant association between ABO blood groups. Although patients with blood group A and AB were higher than in controls, there was no statistically significant. Some authors observed that A blood type was significantly more frequent in patients with some skin diseases. To explain the relationship between blood group antigens and psoriasis, further studies are needed in larger case series.

References

- Christophers E, Mrowietz U. Psoriasis. In: Fitzpatrick's Dermatology in General Medicine. Eisen AZ, Wolff K, Freedberg IN, Austen KF, editors. Dermatology in General Medicine. 2003, New York: McGraw Hill, 407–427.
- 2. Habif TP. Psoriasis and other papulosquamous diseases. 2004; St. Louis: Mosby, 209-239.
- Gaspari AA. Innate and adaptive immunity and the pathophysiology of psoriasis. J Am Acad Dermatol 2006; 54: 67-80. PMID: 16488332
- 4. Pişkin G. Psoriyazisin patogenezi. T Klin Dermatol 2005;13: 5-12.
- 5. Ertuğrul E, Turgay M. Psoriyazis immünopatogenezi. T Klin Dermatol 2005; 13: 13-15.

- Su M, Lu SM, Tian DP, Zhao H, Li XY, Li DR, Zheng ZC. Relationship between ABO blood groups and carcinoma of esophagus and cardia in Chaosan inhabitants of China. World J Gastroenterol 2001; 7: 657-661. PMID: 11819849
- Nakagoe T, Fukushima K, Nanashima A, Sawai T, Tsuji T, Jibiki MA, et al. Comparison of the expression of ABH/Lewis-related antigens in polypoid and non-polypoid growth types of colorectal carcinoma. J Gastroenterol Hepatol 2001; 16: 176-183. PMID: 11207898
- 8. You WC, Ma JL, Liu W, Gail MH, Chang YS, Zhang L, et al. Blood type and family cancer history in relation to precancerous gastic lesions. Int J Epidemiol 2000; 29: 405-407. PMID: 10869310
- Graziano SL, Tatum AH, Gonchoroff NJ, Newman NB, Kohman LJ. Blood group antigen A, and flow cytometric analysis in resected early-stage nonsmall cell lung cancer. Clin Cancer Res 1997; 3: 87-93. PMID: 9815542
- Pyd M, Rzewnicki I, Suwayach U. ABO blood groups in patients with laryngeal and hypopharyngeal cancer. Otolaryngol Pol 1995; 49: 396-398.
 PMID: 9454190
- Pinkston JA, Cole P. ABO blood groups and salivary gland tumors (Alabama, United States). Cancer Causes Control 1996; 7: 572-574. PMID: 8932916
- 12. Marinaccio M, Traversa A, Carioggia E, Valentino L, Coviella M, Salamanna S, et al. Blood groups of the ABO system and survival rate in gynecologic tumors. Minerva Ginecol 1995; 47: 69-76. PMID: 7630512
- Juhl BR. Blood group antigens in transitional cell tumors of the urinary bladder. An immunohistochemical study. Dan Med Bull 1994; 41: 1-11. PMID: 8187557
- 14. Vioque J, Walker AM. Pancreatic cancer and ABO blood types: a study of cases and controls. Med Clin (Barc) 1991; 96: 761-764. PMID: 1875761
- 15. Jia DX. Bone tumor and ABO blood type. 1991; 13: 220-222. Zhonghua Zhong Liu Za Zhi PMID: 1664798
- 16. Cordon-Cardo C, Reuter VE, Finstad CL, Sheinfeld J, Lloyd KO, et al. Blood group related antigens in human kidney: modulation of Lewis determinants in renal cell carcinoma. Cancer Res 1989; 49: 212-218. PMID: 2461798
- 17. Anderson DE, Haas C. Blood type A and familial breast cancer. Cancer 1984; 54: 1845-1849. PMID: 6478419
- Walker PD, Karnik S, de Kerion JB, Pramberg JC.
 Cell surface blood group antigens in prostatic carcinoma. Am J Clin Pathol 1984; 81:503-506. PMID: 6702752
- Jordan GH, Lynch DF. Relationship of blood group to testicular carcinoma. Urology 1983; 22: 265-267.
 PMID: 6623772
- 20. Jager MJ, Všlker-Dieben HJ, De Wolff-Roundaal D, Kakebeeke-Kemme H, DÕAmaro J. Possible relation between HLA and ABO type and prognosis of

- uveal melanoma. Doc Ophthalmol 1992; 82: 43-47. PMID: 1305026
- Olasode OA. Is ABO blood grouping a gene marker for vitiligo? Niger J Med 2002; 11: 193. PMID: 12956000
- 22. Mohan L, Singh G, Kaur P, Pandey SS, Mohan R, Niyogi AK. "ABO blood groups and vitiligo". Indian J Dermatol 1982; 27: 60-62. PMID: 7129532
- 23. Wasfi AI, Saha N, El Munshid HA, El Sheikh FS, Ahmed MA. Genetic association in vitiligo: ABO, MNSs, Rhesus, Kell and Duffy blood groups. Clin Genet 1980; 17: 415-417. PMID: 6772363
- 24. Tirado-Sánchez A, Ponce-Olivera RM. ABO and Rhesus blood groups and their non-existent relationship with pemphigus vulgaris. Acta Dermatovenerol Alp Pannonica Adriat 2010; 19: 47-48. PMID: 20976424
- Shahkar H, Fallahzadeh MK, Namazi MR. ABO blood groups and pemphigus vulgaris:no relationship. Acta Dermatovenerol Alp Pannonica Adriat 2010; 19: 49-51. PMID: 20372776
- 26. Tamega Ade A, Bezerra LV, Pereira Fde P, Miot HA. [Blood groups and discoid lupus erythematosus]. An Bras Dermatol 2009; 84: 477-481. PMID: 20098849

- 27. Choudhury SD, Gupta LC. ABO blood groups in lichen planus. Indian J Dermatol 1979; 24: 27-29. PMID: 540973
- 28. Vaish RP, Jena DC, Panigrahi RK. Blood roups in oral lichen planus in southern Orissa. J Indian Dent Assoc 1985; 57: 183-186. PMID: 3864866
- 29. Thomopoulou- Doukoudakis A, Squier CA, Hill MW. Distribution of ABO blood group substances in various types of oral lichen planus. J Oral Pathol 1983; 12: 47-56. PMID: 6403685
- 30. De Giorgi V, Grazzini M, Gori A, Alfaioli B, Rossari S, Crocetti E, Vocioni F, Lotti T. ABO blood group and risk of cutaneous malignant melanoma. Eur J Cancer Prev 2011; 20: 121-122. PMID: 21332097
- 31. Xie J, Qureshi AA, Li Y, Han J. ABO blood group and incidence of skin cancer. PLoS One 2010 4; 5: e11972. PMID: 20694147
- 32. Tursen U,Tiftik EN, Unal S, Gunduz O, Kaya TI, Camdeviren H, Ikizoglu G. Relationship between ABO blood groups and skin cancers. Dermatol Online J 2005; 11: 44. PMID: 16409940