Protein tyrosine phosphatase PTPN22 +1858C/T polymorphism is associated with active vitiligo

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Abstract. Vitiligo is characterized by a skin depigmentation disorder resulting from an autoimmune response targeting melanocytes. Within the genetic factors involved in the development of the vitiligo immune response, various genes in the major histocompatibility complex (MHC) and non-MHC loci have been considered to be risk factors. The PTPN22 gene encodes for a lymphoid protein tyrosine phosphatase, a regulator of the activation and development of T-cells. The +1858C/T polymorphism has been associated to autoimmune disease susceptibility in different populations and could be implicated in the onset of vitiligo. To assess the possible association between the presence of PTPN22 +1858C/T and vitiligo, 187 patients with vitiligo and 223 control subjects were analyzed in the study. Genomic DNA was isolated using the salting-out method and samples were subjected to polymerase chain reaction-restriction fragment length polymorphism in order to detect the PTPN22 +1858C/T polymorphism. Causal associations were determined by χ² test and their respective odds ratio (OR) was assessed in a 2x2 contingency table. The results showed an association between active vitiligo and the allele T load [P=0.0418; OR, 2.5706; 95% confidence interval (CI), 1.0040‑6.5816], and active vitiligo-CT genotype (P=0.0389, OR, 2.6548; 95% CI, 1.0191‑6.9156). In conclusion, the present data indicates a possible association between the PTPN22 +1858C/T genotype and a significant susceptibility of developing an active form of vitiligo.

Introduction

Vitiligo is an acquired multifactorial progressive depigmentation disorder, characterized by the presence of circumscribed white macules on the skin in which the progressive loss of functional melanocytes in the epidermis is observed (1,2). Vitiligo has been reported to affect 0.1-2% of the population worldwide, but the prevalence varies considerably among the populations or ethnic groups analyzed (1,3,4) and can be as high as those reported in Mexico (4%) or India (8.8%), affecting the two genders equally (5). Furthermore, vitiligo is considered an autoimmune disease (AID), as it can be associated with AIDs, including Hashimoto's thyroiditis, diabetes mellitus, Addison's disease, alopecia areata and ophthalmic abnormalities, such as iritis (4,6-8). This association is supported by: i) The presence of lymphocytes in the dermis of early lesions, ii) antibodies against melanocytes and antigens from the melanogenic system in a number of patients with active disease (9,10), and iii) the response to treatment with immunomodulatory agents, including corticosteroids and phototherapy (11,12).

Autoimmunity is responsible for the misdirected attacks by the immune system of the body against itself, as a result of the failure to recognize self-antigens correctly. This phenomenon is usually a harmless process, but has the potential to lead to a broad spectrum of multiple complex AIDs. Genetic and environmental factors are believed to be the triggers of progression from harmless autoimmunity to AID. The major histocompatibility complex (MHC) locus appears to be a major genetic factor that confers susceptibility to multiple AIDs (13). However, a non-MHC gene may contribute to risk in different AIDs (12).
The PTPN22 gene located on chromosome 1p13.2 encodes a lymphoid protein tyrosine phosphatase (LYP) (14) that is important as a negative control of T-cell activation and T-cell development. PTPN22 belongs to a family of protein tyrosine phosphatases that are involved in the prevention of spontaneous T-cell activation by dephosphorylation and inactivation of T-cell receptor-associated kinases and their substrates (15). PTPN22 is specifically expressed in lymphocytes and suppresses the downstream mediators of T-cell receptor signaling through binding with C-terminal Src kinase (CSK) (15).

Polymeric variants of this gene have been previously associated with various AIDs. The +1858C/T single-nucleotide polymorphism (SNP) (rs2476601), in exon 14 of the PTPN22 gene has been associated with susceptibility to several diseases and dermatological pathologies in different populations (15-18). This polymorphism results in an amino acid change from arginine (Arg) to tryptophan (Trp) in codon 620. The functional significance of this SNP has been discussed by a number of studies (19-22). Experimental evidence indicates that the disease-associated LYP variant Trp620 prevents the interaction of LYP with CSK (19). As a result, the T-cell receptor-associated kinases may be able to induce T-cell activation in an uncontrolled manner and this may increase the overall immune system reactivity and predispose an individual to AID (20,21).

Other studies (21-23) have demonstrated that the +1858T allele is a hypomorphic variant, which means that the polymorphic variant of the protein has a diminished function and cannot inhibit the T cell receptor signaling in T cells correctly; therefore, LYP-620Trp is considered a loss-of-function protein. This impaired LYP function in T cells may drive hyperresponsive B cells to break the tolerance in the periphery and to the production of autoantibodies, leading to the development of AID (22).

In vitiligo, the SNP rs2476601 has been identified as a risk factor for a specific generalized form in European Caucasian populations of Romania (23). The data collected for this gene in Mexico are obsolete or limited. The results obtained by different studies support the association of the PTPN22 1858T allele with sporadic childhood-onset systemic lupus erythematosus (SLE) in the Mexican population (24), and susceptibility to rheumatoid arthritis in the Western Mexican population (16). However, thus far, no similar studies have been carried out on vitiligo. The aim of the present hospital case-control study was to investigate the possible involvement of the PTPN22 gene 1858C/T polymorphism in Mexican patients with vitiligo from Northeast Mexico.

Materials and methods

Selection of subject and controls. A total of 187 Mexican patients diagnosed with vitiligo and recruited by the Dermatology Department of the ‘Dr Jose E. Gonzalez’ University Hospital of the Universidad Autonoma de Nuevo Leon (UANL; Monterrey, Mexico), were selected in the study conducted between November 2009 and May 2010 (from the states of Coahuila, Nuevo Leon, San Luis Potosi, Tamaulipas and Zacatecas). The University Hospital-UANL Institutional Review Board approved and registered the study under the code DE08-008. Subsequent to providing their informed consent, the patients were evaluated and surveyed by direct interview.

The information gathered from the patients included: Age, gender, birthplace, full clinical history, age of disease onset, previous treatment (if any), vitiligo lesions in previously traumatized regions (Koebner's phenomenon), class and distribution of vitiligo in the patient and their families. The patients were further classified as exhibiting the following types of vitiligo: focal (one or more maculae in an unsegmented pattern); segmental (one or more maculae distributed in a single dermatome or zosteriform pattern), vulgaris (symmetric or asymmetric distribution of maculae in one or more areas) and universal (>80% skin depigmentation). The course of the disease was classified as active (progressive, appearance of new maculae in the last six months) or stable.

The control group consisted of 232 patients from the Mexican Mestizo population who were recruited during an earlier study from a general unbiased healthy population through the Blood Bank of the aforementioned hospital. For study purposes, a Mexican Mestizo was defined as a person born in Mexico, whose last three ascending generations were also born in the country.

Experimental procedure

DNA isolation. Peripheral venous blood samples were obtained from the patients with vitiligo and the controls. The samples were centrifuged, and theuffy-coat was processed for DNA isolation by the salting-out method and finally suspended in Tris-EDTA (pH 7.8), at a final concentration of 0.1-1.0 µg/µl.

PTPN22 +1858C/T genotyping. Genotyping was performed by polymerase chain reaction-restriction fragment length polymorphism (PCR-RFLP) as modified by Harrison et al (25). PCR was carried out using an MJ Research PTC 100 thermal cycler (Bio-Rad, Hercules, CA, USA) as follows: A total amount of 250 ng genomic DNA, 0.5 µM PTPN22 +1858C/T primers (forward, 5'-ATGGTGCTTCAACGGAATT-3' and reverse, 5'-CATGCTGCTATTGCTCTG-3'), 0.2 mM deoxyribo-nucleotide triphosphates, 1.5 mM MgCl2 and 2.5 units of Taq DNA polymerase (Promega Corporation, Madison, WI, USA) at 94°C/1 min, 58°C/1 min and 72°C/1 min (33 cycles). The PCR samples (~1 µg) were subsequently digested overnight with XcmI (New England Biolabs, Ipswich, MA, USA) at 37°C. The digested ampiclons were analyzed by electrophoresis in 2% agarose gels using ethidium bromide.

Statistical analysis. The collected data were organized in a database and statistical calculations were performed with the SPSS 17.0 software for windows (SPSS, Inc. Chicago, IL, USA) and the Epi-INFOTM 7 statistical program (USD Incorporated 2008, Stone Mountain, GA, USA). The Hardy-Weinberg equilibrium was obtained using a goodness-of-fit test. Genotypic dependence between the patients and controls was determined by the χ² test, and the odds ratio (OR) was calculated from 2x2 contingency tables. The Mann-Whitney U test and Spearman's Rho were used to determine the correlation of age, gender and genotype for each studied group. P≤0.05 was considered to indicate a statistically significant difference.

Results

Clinical characteristics of the participants. The present study analyzed the genotype distribution and allele frequency for PTPN22 +1858C/T polymorphism corresponding to...
The study consisted of 187 patients with vitiligo, including 97 (51.9%) females and 90 (48.1%) males. The medium age of the patients at the time of consultation was 41.33±16.194 years, with a median of 44 years. The mean age of vitiligo onset was 26.28±15.752 years, with a median age of 10 years. The patients with vitiligo were classified into four categories according to the type of vitiligo, and into two main groups depending on disease activity (Table I).

The most common illness associated with vitiligo was thyroid alterations, in which hypothyroidism was present in 19 patients (10.2%), subclinical hypothyroidism in 12 patients (6.4%), hyperthyroidism in five patients (2.7%) and subclinical hyperthyroidism in only one patient (0.5%).

In order to assess the Hardy-Weinberg equilibrium, the data were analyzed by means of a $\chi^2$ goodness-of-fit test. Several tests were used (control group, Pearson test $P=0.763198$, likelihood-ratio (Llr) test $P=0.673069$, Fisher’s exact test $P=1.000000$; case group, Pearson test $P=0.541091$, Llr test $P=0.397683$, Fisher’s exact test $P=1.000000$), all of which had a $P$-value >0.05, indicating that the rs2476601 genotype was in Hardy-Weinberg equilibrium in patients with vitiligo and the controls. The statistical results obtained from the study are shown in Table III.

### Table I. General characteristics of patients with vitiligo.

| Type of vitiligo | Gender, n (%) | Vitiligo activity, n (%) | Koëbner phenomenon, n (%) | Age of onset, n (%) |
|------------------|---------------|--------------------------|---------------------------|------------------|
|                  | Female | Male | Active | Stable |  | <30 years, n (%) | ≥30 years, n (%) |
| Vulgaris         | 88 (47.06) | 78 (41.71) | 85 (45.45) | 81 (43.32) | 47 (25.13) | 26.46±15.201 | 93 (39.04) | 73 (35.35) |
| Universal        | 1 (0.53)  | 2 (1.07)  | -      | 3 (1.6)  | -          | 12.33±15.373 | 2 (1.07)   | 1 (0.53)  |
| Focal            | 7 (3.74)   | 10 (5.35) | 6 (3.21) | 11 (5.88) | 2 (1.07)  | 28.41±19.912 | 9 (4.81)   | 8 (4.28)  |
| Segmental        | 1 (0.53)   | -      | 1 (0.53) | -      | -          | 3           | 1 (0.53)   | -      |

*Age is presented as the mean ± standard deviation.

### Table II. Autoimmune pathologies associated with vitiligo in patients.

| Type of vitiligo | Thyroid | AA | DM | HTN | Atopy | SLE | RA | MG | AH |
|------------------|---------|----|----|-----|-------|-----|----|----|----|
| VV, n (%)        | 33 (17.65) | 5 (2.67) | 13 (6.95) | 23 (12.3) | 6 (3.21) | 1 (0.53) | 1 (0.53) | 1 (0.53) | 1 (0.53) |
| UV, n (%)        | 1 (0.50) | - | - | - | - | 1 (0.53) | - | - | - |
| FV, n (%)        | 3 (1.52) | 1 (0.53) | 1 (0.53) | 1 (0.53) | 1 (0.53) | - | - | - | - |
| SV, n (%)        | - | - | - | - | - | 1 (0.53) | - | - | - |
| Total, %         | 19.78 | 3.21 | 7.49 | 12.83 | 4.29 | 1.07 | 0.53 | 0.53 | 0.53 |

AA, alopecia areata; DM, diabetes mellitus; HTN, arterial hypertension; SLE, systemic lupus erythematosus; RA, rheumatoid arthritis; MG, myasthenia gravis; AH, autoimmune hepatitis; VV, vulgaris vitiligo; UV, universal vitiligo; FV, focal vitiligo; SV, segmental vitiligo.

**Table III. Association of the genotypes in patients with vitiligo.** When the PTPN22 +1858C/T genotypes and allele frequencies were compared between the whole cohort of cases and controls, evidence of association was observed (Table III). The CT genotype was more frequent in patients with vitiligo than in controls ($P=0.0445$). However, no significant association was observed in the T allele of patients with vitiligo when comparing all the vitiligo types against controls.

Finally, when considering population variables, including gender, type of vitiligo, age of disease onset, immune pathologies and disease activity, an association was only observed between the active vitiligo allele T load ($P=0.0418$; OR, 2.5706; 95% confidence interval (CI), 1.0040-6.5816) and active vitiligo-CT genotype ($P=0.0389$; OR, 2.6548; 95% CI, 1.0191-6.9156) (Table III).

### Discussion

Vitiligo is a complex depigmentation disorder in which there is progressive and selective destruction of the cells that produce the pigment of the skin, hair, and eyes, known as melanocytes (26). Vitiligo is believed to have a non-Mendelian genetic predisposition pattern, with a polygenic and multifactorial inheritance (27). Multiple trigger factors have been associated with the development of vitiligo, including stress, infections, traumatisms, malignancy, sun exposure, neurological abnormalities, and endocrine and autoimmune alterations, which can act independently or in combination (28).
Previous studies associate vitiligo to autoimmune pathologies due to its frequent association with diseases of this nature, in which melanocyte cell death by immune-mediated apoptosis is considered an important susceptibility factor for vitiligo, emphasized by the detection in serum of auto-antibodies and reactive T lymphocytes targeted against this cell type (29). In a number of case-control genetic studies of vitiligo, a genetic association of the MHC was observed by testing different MHC markers in patients with different vitiligo phenotypes versus controls (30). Since MHC loci have been associated with the development of vitiligo, identification of non-associated MHC loci with this pathology and immune diseases have been reported. The PTPN22 locus is one of the strongest risk factors outside of the MHC that associates with AIDS (31).

The product of the PTPN22 gene is a phosphatase known as LYP, which acts as a negative regulator of T cell signaling. Since LYP is also expressed in other cell types, such as B cells, monocytes, neutrophils and dendritic cells (16), the presence of the +1858T allele results in a loss of functional proteins that are unable to regulate T cell activation leading to a hyperresponsive phenotype of T, B and dendritic cells (16,32), thereby having a possible role in the pathogenesis of vitiligo. This observation is supported by a previous study that described the increase of Th17 and activated dendritic cells and cellular immune elements in the active margins of vitiligo lesions, which could potentially drive the inflammatory responses in the skin (33).

In the current study, the presence of the PTPN22 +1858C/T polymorphism in a population affected by vitiligo from northeast Mexico was analyzed. Even though the PTPN22 +1858C/T polymorphism has been reported in Mexico for other diseases, the data remain limited. Previous studies in Mexican populations have described the allele and haplotype frequency distributions of the PTPN22 gene (34) in SLE (24) and susceptibility to rheumatoid arthritis (15). In the case of vitiligo, there are few studies worldwide that make reference to the development of this pathology and its association with this polymorphism. There is evidence that the PTPN22 +1858T allele constitutes a generalized risk for vitiligo in a European Caucasian (Romanian) population, highlighting the importance of a genetically mediated autoimmune mechanism in the pathogenesis of vitiligo (23). By contrast, studies of the segmental form of the disease in Egyptian females failed to find an association with the presence of this polymorphism (35). Some of these positive findings have been confirmed by a meta-analysis performed.
on European and Asian populations by Song et al (36) in 2013. The PTPN22 +1858C/T polymorphism has not been previously assessed for vitiligo in American populations. Although an association was found between all the vitiligo types analyzed in general and the studied polymorphism (P=0.0445; OR, 2.3184; 95% CI, 1.0003-5.3736) this result was not sustainable when focusing on the T allele (P=0.0479; OR, 2.2595; 95% CI, 0.9869-5.1730), however, a notable association was found for the active forms of vitiligo and the T allele load of the polymorphism (P=0.0389), which conferred an increased OR of 2.6548.

In conclusion, the present data indicate the involvement of the PTPN22 +1858T allele as a genetic risk factor for susceptibility to active vitiligo in the Mexican population, supporting the evidence of the possible involvement of the immune response in the pathogenesis of vitiligo.

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References

1. Sehgal V and Srivastava G: Vitiligo: compendium of clinico-epidemiological features. Indian J Dermatol Venereol Leprol 75: 149-156, 2007.
2. Guerra L, Dellambra E, Brescia S and Raskovic D: Vitiligo: pathogenetic hypotheses and targets for current therapies. Curr Drug Metab 11: 451-467, 2010.
3. Steinier D, Bedin V, Brito M, Tadeu R and Steiner T: Vitiligo: An Bras Dermatol 79: 335-351, 2004.
4. Parsad D, Dogra S and Kanwar AJ: Quality of life in patients with vitiligo. Health Qual Life Outcomes 1: 58-63, 2003.
5. Moretti S. Vitiligo: Orphanet Encyclopedia. October 2003. http://www.ophtha.net/data/patho/GB/uk-vitiligo.pdf. Accessed December 13, 2013.
6. Huggins RH, Janusz CA and Schwartz RA: Vitiligo: a sign of systemic disease. Indian J Dermatol Venereol Leprol 72: 68-71, 2006.
7. Jin Y, Mailloix CM, Gowen K, et al: NALP1 in vitiligo-associated multiple autoimmune disease. N Engl J Med 356: 1216-1225, 2007.
8. van Gell N, Speeckaert M, et al: Clinical profile of generalized vitiligo patients with associated autoimmune/autoinflammatory diseases. J Eur Acad Dermatol Venereol 28: 741-746, 2014.
9. Shahil EM, Chatterjee S, et al: Vitiligo: pathomechanisms and genetic polymorphism of susceptible genes. Indian J Exp Biol 44: 526-539, 2006.
10. Kemph EH, Gavastal NG, et al: Autoantibody responses to melanocytes in the depigmenting skin disease vitiligo. Autoimmun Rev 10: 762-765, 2011.
11. Almeida P, Borrego L, Rodríguez-López J, et al: Vitiligo. Treatment of 12 cases with topical tacrolimus. Actas Dermosifiliogr 96: 159-163, 2005 (In Spanish).
12. Xu AE, Zhang DM, Wei XD, Huang B and Lu LJ: Efficacy and safety of tacrolimus cream 0.1% in the treatment of vitiligo. Int J Dermatol 48: 86-90, 2009.
13. Amur S, Parekh A and Mummamani P: Sex differences and genomics in autoimmune diseases. J Autoimmun 38: J254-J165, 2012.
14. Seen S, Dadi H, Shaoul E, Sharfe N and Roifman CM: Cloning and characterization of a lymphoid-specific, inducible human protein tyrosine phosphatase, Lyp. Blood 93: 2013-2024, 1999.
15. Zhebrun D, Kudryashova Y, Babenko A, et al: Association of PTPN22 1858T/G genotype with type 1 diabetes. Graves’ disease but not with rheumatoid arthritis in Russian population. Aging (Albany NY) 3: 368-373, 2011.
16. Torres-Carrillo NM, Ruiz-Noa Y, et al: The +1858C/T PTPN22 gene polymorphism confers genetic susceptibility to rheumatoid arthritis in Mexican population from the Western Mexico. Immunol Lett 147: 41-46, 2012.
17. Ceníz MC, Márquez A, Cordero-Coma M, et al: Lack of association between the protein tyrosine phosphatase non-receptor type 22 R263Q and R620W functional genetic variants and endogenous non-anterior uveitis. Mol Vis 19: 638-643, 2013.
18. Chen YF and Chang JS: PTPN22 C1858T and the risk of psoriasis: a meta-analysis. Mol Biol Rep 39: 7861-7870, 2012.
19. Fjaer JH, Nøssum E, Musumeci L, Alonso A, et al: A functional variant of lymphoid tyrosine phosphatase is associated with type 1 diabetes. Nat Genet 36: 337-338, 2004.
20. Cantón I, Akhtar S, Gavalas NG, Kawokrder BJ, Blomhoff A, Watson PF, Westman AP and Kemp EH: A single-nucleotide polymorphism in the gene encoding lymphoid protein tyrosine phosphatase (PTPN22) confers susceptibility to generalised vitiligo. Genes Imm 6: 584-587, 2005.
21. Yang T, Miletic AV, Bottini N and Mustelin T: Protein tyrosine phosphatase PTPN22 in human autoimmunity. Autoimmunity 40: 452-461, 2007.
22. Zakhman J, Hermiston M, Steiner D, Hasegawa K, Chan A and Weiss A: PTPN22 deficiency cooperates with the CD45 E613R allele to break tolerance on a nonautoimmune background. J Immunol 182: 4093-4106, 2009.
23. Laberge GS, Bislea SA, Pain PR and Spritz RA: The PTPN22 +1858C/T (R620W) functional polymorphism is associated with generalized vitiligo in the Romanian population. Pigment Cell Melanoma Res 21: 206-208, 2008.
24. Baca V, Velázquez-Cruz R, Salas-Martínez G, Espinosa-Rosas F, Saldaña-Alvarez Y and Orozco L: Association analysis of the PTPN22 gene in childhood-onset systemic lupus erythematosus in Mexican population. Genes Immun 7: 693-695, 2006.
25. Harrison P, Pointon JJ, Farrar C, Brown MA and Wordsworth BP: Effects of PTPN22 C1858T polymorphism on susceptibility and clinical characteristics of British Caucasian rheumatoid arthritis patients. Rheumatology (Oxford) 45: 1009-1011, 2006.
26. Aris M: Origin of normal and malignant melanocytes. Acta Bioquim Clin Latinoam 43: 333-337, 2009.
27. Zhang XJ, Chen JJ and Liu JH: The genetic concept of vitiligo. J Dermatol Sci 39: 137-146, 2005.
28. Sandoval-Cruz M, García-Carrasco M, Sánchez-Porras R, Mendoza-Pinto C, Jiménez-Hernández M, Munguía-Realpozo P and Ruiz-Aruguelles A: Immunopathogenesis of vitiligo. Autoimmun Rev 10: 762-765, 2011.
29. Sabín K and Parsad E: Stability in vitiligo: Is there a perfect way to predict it? J Cutan Aesthetic Surg 6: 75-82, 2013.
30. Spritz RA: The genes of generalized vitiligo and associated autoimmune diseases. Pigment Cell Melanoma Res 20: 271-278, 2007.
31. Burn GL, Svensson L, Sanchez-Blanco C, Saini M and Cope AP: Why is PTPN22 a good candidate susceptibility gene for autoimmune disease? FEBS Lett. 585: 3689-3698, 2011.
32. Zhang J, Zahir N, Jiang Q, Miliotis H, Heyraud S, Meng X, Dong B, Xie G, Qiu F, Hao Z, McCulloch CA, Keystone EC, Peterson AC and Siminovitch KA: The autoimmune disease-susceptible variant PTPN22 CT60 predicts caplan-mediated Lyp/Pep degradation associated with lymphocyte and dendritic cell hyperresponsiveness. Nat Genet 43: 902-907, 2011.
33. Wang CQ, Cruz-Inigo AE, Fuentes-Duculan J, Moussai D, Gulati N, Sullivan-Whalen M, Gilleaudeau P, Cohen JA and Krueger JG: TH17 cells and activated dendritic cells are increased in vitiligo lesions. PLoS One 6: e18907, 2011.
34. Lins TC, Vieira RG, Grattapaglia D and Pereira RW: Allele and haplotype frequency distribution in PTPN22 gene across variable ethnic groups: Implications for genetic association studies for autoimmune diseases. Autoimmunity 43: 308-316, 2010.
35. Elmongy NN and Abu Khalil RE: PTPN22 gene polymorphism in Egyptian females with non-segmental vitiligo. Comp Clin Pathol 22: 961-964, 2013.
36. Xu AE, Kim JH and lee YH: The CTLA-4 +49 A/G, CT60 A/G and PTPN22 1858 C/T polymorphisms and susceptibility to vitiligo: a meta-analysis. Mol Biol Rep 40: 2985-2993, 2013.