Evaluation of PTPN22 polymorphisms and Vogt-Koyanagi-Harada disease in Japanese patients

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Purpose: Vogt-Koyanagi-Harada (VKH) disease is an autoimmune disorder against melanocytes. Polymorphisms of the protein tyrosine phosphatase non-receptor 22 gene (PTPN22) have recently been reported to be associated with susceptibility to several autoimmune diseases. In this study, genetic susceptibility to VKH disease was investigated by screening for single nucleotide polymorphisms (SNPs) of PTPN22.

Methods: A total of 167 Japanese patients with VKH disease and 188 healthy Japanese controls were genotyped by direct sequencing methods for six SNPs (rs3811021, rs1217413, rs1237682, rs3761935, rs3789608, and rs2243471) of PTPN22 including the uncoding exons.

Results: The six SNPs in PTPN22 showed no significant association with susceptibility to VKH disease or its ocular, neurologic, or dermatological manifestation.

Conclusions: Further studies are needed to clarify the genetic mechanisms underlying VKH disease.

Vogt-Koyanagi-Harada (VKH) disease is one of the most frequent forms of uveitis in Japan [1]. It is characterized as bilateral panuveitis accompanied by neurologic and skin lesions [2,3]. This disease is considered to be an autoimmune disease against melanocytes [4,5]. Though the etiology of VKH disease still remains unknown, genetic factors may play an important role in susceptibility as indicated by an established association between VKH disease and specific human leukocyte antigen (HLA)-DRB1 alleles [6,7].

The protein tyrosine phosphatase non-receptor 22 gene (PTPN22) is located on chromosome 1p13.3-p13.1, and it encodes the lymphoid-specific phosphatase (Lyp) that is important in the negative control of T-cell activation and development [8-10]. Recently, it was reported the single nucleotide polymorphism (SNP), R620W (rs2476601), in PTPN22 increased susceptibility to several autoimmune diseases including rheumatoid arthritis (RA), systemic lupus erythematosus (SLE), and insulin dependent diabetes mellitus (IDDM) [11-15]. In the PTPN22 risk variant (rs2476601), this substitution disrupts an interaction between Lyp and the protein tyrosine kinase, Csk, and may translate biologically to a potential for ‘hyperreactive’ pathogenic T-cell responses [8].

This R620W mutation was not observed in the Japanese population [16,17]. Therefore, in this study, we analyzed six SNPs, which belong to the same haplotype block as R620W (rs2476601) in PTPN22. For the efficacy of the linkage analysis, we chose six SNPs of which minor allele frequencies were more than 15% from the database of Japanese Single Nucleotide Polymorphisms [18,19].

METHODS

We recruited 167 VKH (72 males and 95 females) patients and 188 healthy controls for this study. All patients and control subjects were Japanese. Patients were diagnosed according to the “Revised Diagnostic Criteria for VKH Disease” [3] at the Uveitis Survey Clinic of the Hokkaido University Hospital (Sapporo, Japan) and Yokohama City University Hospital (Yokohama, Japan). All patients and control subjects were informed of the study’s purpose, and their consent obtained. The study was approved by the ethics committee from each institute participating in this study.

DNA was prepared from peripheral blood specimens using the QIAamp DNA Blood Mini Kit (Qiagen, Tokyo, Japan). Six SNPs (rs3811021, rs1217413, rs1237682, rs3761935, rs3789608, and rs2243471) from the PTPN22 region were examined (Figure 1). Each of the six SNPs was amplified by standard polymerase chain reactions (PCRs; Table 1). After purification using ExoSAP-IT (USB
Corporation, Cleveland, OH), the PCR products were sequenced with Big Dye Terminator v3.1 (Applied Biosystems, Foster City, CA) using either sense or antisense primers (Table 1). The BigDye XTerminator Purification Kit (Applied Biosystems) was used to purify the DNA from sequencing reactions. The sequencing reactions were analyzed using an ABI3130 sequencer (Applied Biosystems).

### Statistical analysis:
For statistical analyses, the Hardy–Weinberg equilibrium was tested for each SNP among the control subjects. Genotype frequency differences between the case and control genotypes were assessed by the $\chi^2$ test. The calculation of linkage disequilibrium (LD) and pair-wise LD (D' value) between SNPs of the $PTPN22$ region and the haplotypes was performed with Haploview software, version 3.32. The maximum likelihood estimates of haplotype frequencies were estimated by pairs of unphased genotypes using the expectation-maximization (EM) algorithms in the R package ‘haplo.stats’ [20].

### RESULTS

Allele frequencies for the six SNPs covering the gene were in Hardy–Weinberg equilibrium in both the patients and controls. The allelic frequency of each SNP in both groups was nearly equal, and no association was detected when compared independently (odds ratio, OR 1.14–1.35; Table 2). Stratifying the patients by the presence of diffuse choroiditis, sunset glow fundus, nummular chorioretinal depigmented spots, neurologic auditory involvement, meningismus, tinnitus, cerebrospinal fluid pleocytosis, or integumentary findings also revealed no evidence of association in VKH disease (data not shown). We calculated pairwise D’ values for all SNP pairs in $PTPN22$ (Figure 2). The pairwise D’ values in the gene were nearly 1 among almost all SNP pairs, indicating the SNPs were highly associated with each other and the entire $PTPN22$ was contained within a single LD block. Haplotype analysis predicted and revealed that $PTPN22$ was not associated with VKH disease in this Japanese cohort (data not shown).

### DISCUSSION

In the present study, we analyzed polymorphisms of the new candidate gene, $PTPN22$, in Japanese patients with VKH disease.
disease. The gene encodes an important negative regulator of T cell activation [9]. An SNP of PTPN22, R620W (rs2476601) was reported to be associated with several autoimmune diseases such as RA, SLE, and IDDM [11,12,14,15]. However, this SNP, which disrupts an interaction between Lyp and the protein tyrosine kinase, Csk, does not exist as a polymorphism in the Japanese population [9,10,12]. Therefore, in this study, we examined six other SNPs to evaluate the susceptibility locus of PTPN22.

HLA-DRB1 is a common genetic factor in autoimmune diseases (RA and IDDM). Therefore, there may be other common genetic factors in VKH disease [21].

VKH disease is considered to be an autoimmune disease against melanocytes [2-5]. In early studies, activated T lymphocytes were elevated and attacked melanocytes of ocular choroidal tissue in patients in the active phase of VKH disease [22]. Antigen-specific T-cell assay revealed that peptide fragments of the tyrosinase family proteins (tyrosinase, tyrosinase related protein 1 and 2) proliferated in T lymphocytes collected from VKH patients [4,5]. These proteins are found in human melanocytes. These antigen-specific T cell responses were detected in cells collected from HLA-DRB1*04 positive VKH patients only but not from HLA-DRB1*04 negative patients or HLA-DRB1*04 positive healthy people [7,23]. In the Japanese population, 40% of healthy people have HLA-DRB1*04 [7]. However, people having VKH disease represent only 0.01% of the Japanese population [1,7,24,25]. In addition, some patients with VKH disease are HLA-DRB1*04 negative [7]. Thus, it is believed HLA-DRB1*04 is a major susceptible gene in VKH disease. However, other minor genetic factors still remain unclear. To find other susceptible genes, we studied the tyrosinase gene (TYR), tyrosinase related protein 1 gene (TYRP1), tyrosinase related protein 2 gene (TYRP2), and interferon (IFN-γ), but we could not find any association with these genes and VKH disease [7,26]. Genetic influences of VKH were also investigated in other countries, but the etiology of the disease seems to be unresolved [27-29].

In this study, we found no association between PTPN22 and VKH disease in the individuals studied. Our results suggest that further molecular genetic studies are needed to detect novel genetic loci and predisposing genes and to elucidate the true genetic mechanisms underlying VKH disease.

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REFERENCES
1. Goto H, Mochizuki M, Yamaki K, Kotake S, Usui M, Ohno S. Epidemiological survey of intraocular inflammation in Japan. Jpn J Ophthalmol 2007; 51:41-4. [PMID: 17295139]
2. Kitamura M, Takami K, Kitaichi N, Namba K, Kitamei H, Kotake S, Ohno S. Comparative study of two sets of criteria for the diagnosis of Vogt-Koyanagi-Harada's disease. Am J Ophthalmol 2005; 139:1080-5. [PMID: 15953440]
3. Read RW, Holland GN, Rao NA, Tabbara KD, Ohno S, Arellanes-Garcia L, Pivetti-Pezzi P, Tessler HH, Usui M. Revised diagnostic criteria for Vogt-Koyanagi-Harada disease: report of an international committee on nomenclature. Am J Ophthalmol 2001; 131:647-52. [PMID: 11336942]

4. Yamaki K, Gocho K, Hayakawa K, Kondo I, Sakuragi S. Tyrosinase family proteins are antigens specific to Vogt-Koyanagi-Harada disease. J Immunol 2000; 165:7323-9. [PMID: 11120868]

5. Yamaki K, Takiyama N, Itoh N, Mizuki N, Seiya M, Sinsuke W, Hayakawa K, Kotani T. Experimentally induced Vogt-Koyanagi-Harada disease in two Akita dogs. Exp Eye Res 2005; 80:273-80. [PMID: 15670805]

6. Islam SM, Numaga J, Fujino Y, Hirata R, Matsuki M, Maeda H, Masuda K. HLA class II genes in Vogt-Koyanagi-Harada disease. Invest Ophthalmol Vis Sci 1994; 35:3890-6. [PMID: 7928186]

7. Horie Y, Takemoto Y, Miyazaki A, Namba K, Kase S, Yoshida K, Ota M, Hasumi Y, Inoko H, Mizuki N, Ohno S. Tyrosinase gene family and Vogt-Koyanagi-Harada disease in Japanese patients. Mol Vis 2006; 12:1601-5. [PMID: 17200659]

8. Bottini N, Musumeci L, Alonso A, Rahmouni S, Nika K, Rostamkhani M, MacMurray J, Meloni GF, Lucarelli P, Pellecchia M, Eisenbarth GS, Comings D, Mustelin T. A functional variant of lymphoid tyrosine phosphatase is associated with type I diabetes. Nat Genet 2004; 36:337-8. [PMID: 15004560]

9. Siminovitch KA. PTPN22 and autoimmune disease. Nat Genet 2005; 37:1300-2. [PMID: 16314859]

10. Gregersen PK. Gaining insight into PTPN22 and autoimmunity. Nat Genet 2004; 36:1248-9. [PMID: 15565104]

11. Gregersen PK. Gaining insight into PTPN22 and autoimmunity. Nat Genet 2005; 37:1300-2. [PMID: 16314859]

12. Begovich AB, Carlton VE, Honigberg LA, Schrødi SJ, Chokkalingam AP, Alexander HC, Ardlie KG, Huang Q, Smith AM, Spoerke JM, Conn MT, Chang M, Chang SY, Saiki RK, Catanese JJ, Leong DU, Garcia VE, McCallister LB, Jeffery DA, Lee AT, Batiwalla F, Remmers E, Criswell LA, Seldin MF, Kastner DL, Amos CI, Sninsky JJ, Gregersen PK. A missense single-nucleotide polymorphism in a gene encoding a protein tyrosine phosphatase (PTPN22) is associated with rheumatoid arthritis. Am J Hum Genet 2004; 75:330-7. [PMID: 15208781]

13. Kyogoku C, Langefeld CD, Ortmann WA, Lee A, Selby S, Carlton VE, Chang M, Ramos P, Baechler EC, Batiwalla FM, Novitzke J, Williams AH, Gillett C, Rodine P, Graham RR, Ardlie KG, Gaffney PM, Moser KL, Petri M, Begovich AB, Gregersen PK, Behrens TW. Genetic association of the R620W polymorphism of protein tyrosine phosphatase PTPN22 with human SLE. Am J Hum Genet 2004; 75:504-7. [PMID: 15273934]

14. Michou L, Lasbleiz S, Rat AC, Migliorini P, Balsa A, Westhovens R, Barrera P, Alves H, Pierlot C, Glikmans E, Garnier S, Dausset J, Vaz C, Fernandes M, Petit-Texeira E, Lemaire I, Pascual-Salcedo D, Bombardieri S, Dequeker J, Radstake TR, Van Riel P, van de Putte L, Lopes-Vaz A, Prum B, Bardin T, Dieude P, Cornelis F. Linkage proof for PTPN22, a rheumatoid arthritis susceptibility gene and a human autoimmune gene. Proc Natl Acad Sci USA 2007; 104:1649-54. [PMID: 17237219]

15. Criswell LA, Pfeiffer KA, Lum RF, Gonzales B, Novitzke J, Kern M, Moser KL, Begovich AB, Carlton VE, Li W, Lee AT, Ortmann W, Behrens TW, Gregersen PK. Analysis of families in the multiple autoimmune disease genetics consortium (MADGC) collection: the PTPN22 R620W allele associates with multiple autoimmune phenotypes. Am J Hum Genet 2005; 76:561-71. [PMID: 15719322]

16. Wu H, Cantor RM, Graham DS, Lingren CM, Farwell L, Jager PL, Bottini N, Grossman JM, Wallace DJ, Hahn BH, Julkunen H, Hebert LA, Rovin BH, Birmingham DJ, Rioux JD, Yu CY, Kere J, Vyse TJ, Tsao BP. Association analysis of the R620W polymorphism of protein tyrosine phosphatase PTPN22 in systemic lupus erythematosus families: increased T allele frequency in systemic lupus erythematosus patients with autoimmune thyroid disease. Arthritis Rheum 2005; 52:2396-402. [PMID: 16052563]

17. Ichimura M, Kaku H, Fukutani T, Koga H, Mukai T, Miyake I, Yamada K, Koda Y, Hirosumi Y. Associations of protein tyrosine phosphatase nonreceptor 22 (PTPN22) gene polymorphisms with susceptibility to Graves' disease in a Japanese population. Thyroid 2008; 18:625-30. [PMID: 18578611]
26. Horie Y, Kitaichi N, Takemoto Y, Namba K, Yoshida K, Hirose S, Hasumi Y, Ota M, Inoko H, Mizuki N, Ohno S. Polymorphism of IFN-gamma gene and Vogt-Koyanagi-Harada disease. Mol Vis 2007; 13:2334-8. [PMID: 18199975]

27. Du L, Yang P, Hou S, Lin X, Zhou H, Huang X, Wang L, Kijlstra A. Association of the CTLA-4 gene with Vogt-Koyanagi-Harada syndrome. Clin Immunol 2008; 127:43-8. [PMID: 18282809]

28. Hou S, Yang P, Du L, Zhou H, Lin X, Liu X, Kijlstra A. Small ubiquitin-like modifier 4 (SUMO4) polymorphisms and Vogt-Koyanagi-Harada (VKH) syndrome in the Chinese Han population. Mol Vis 2008; 14:2597-603. [PMID: 19122825]

29. Meng Q, Liu X, Yang P, Hou S, Du L, Zhou H, Kijlstra A. PDCD1 genes may protect against extraocular manifestations in Chinese Han patients with Vogt-Koyanagi-Harada syndrome. Mol Vis 2009; 15:386-92. [PMID: 19234630]