Investigation of the Correlation between Graves’ Ophthalmopathy and CTLA4 Gene Polymorphism

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Abstract: Graves’ disease (GD) is an autoimmune inflammatory disease, and Graves’ ophthalmopathy (GO) occurs in 25–50% of patients with GD. Several susceptible genes were identified to be associated with GO in some genetic analysis studies, including the immune regulatory gene CTLA4. We aimed to find out the correlation of CTLA4 gene polymorphism and GO. A total of 42 participants were enrolled in this study, consisting of 22 patients with GO and 20 healthy controls. Chi-square or Fisher’s exact test were used to appraise the association between Graves’ ophthalmopathy and CTLA4 single nucleotide polymorphisms (SNPs). All regions of CTLA4 including promoter, exon and 3’UTR were investigated. There was no nucleotide substitution in exon 2 and exon 3 of CTLA4 region, and the allele frequencies of CTLA4 polymorphisms had no significant difference between patients with GO and controls. However, the genotype frequency of “TT” genotype in rs733618 significantly differed between patients with GO and healthy controls (OR = 0.421, 95%CI: 0.290–0.611, p = 0.043), and the “CC” and “CT” genotype in rs16840252 were nearly significantly differed in genotype frequency (p = 0.052). Haplotype analysis showed that CTLA4 Crs733618Crs16840252 might increase the risk of GO (OR = 2.375, 95%CI: 1.636–3.448, p = 0.043). In conclusion, CTLA4 Crs733618Crs16840252 was found to be a potential marker for GO, and these haplotypes would be ethnicity-specific. Clinical application of CTLA4 Crs733618Crs16840252 in predicting GO in GD patients may be beneficial.

Keywords: Graves’ disease; Graves’ ophthalmopathy; CTLA4; gene polymorphism

1. Introduction

Graves’ disease (GD) is an autoimmune inflammatory disease. The annual incidence of Graves’ disease is 20 to 50 cases per 100,000 persons [1]. Graves’ ophthalmopathy (GO) also called thyroid eye disease occurs in 25–50% of patients suffered from GD [2]. The reported prevalence rates of Graves’ ophthalmopathy among different ethnic populations are limited. An early study demonstrated 6.4 times higher risk of developing GO in Caucasians than that of Asians [3] but has been criticized for a small sample size. Recent studies reported similar prevalence rate in Malaysians (Malay, Chinese, and Indian) [4] and Indians [5] with GD as compared with Caucasian GD patients. The clinical
presentation of GO includes eyelid retraction, restrictive extraocular myopathy, proptosis, glaucoma, exposure keratitis and compressive optic neuropathy [6]. Patients may experience frequent diplopia, dry eye, chemosis, eyelid swelling and even vision loss in severe cases [7]. Additionally, patients with Graves’ disease tend to have concomitant hyperthyroidism.

Genetic analysis has identified several susceptible genes, including genes encoding thyroglobulin, thyrotropin receptor, HLA-DRβ-Arg74, protein tyrosine phosphatase nonreceptor type 22 (PTPN22), cytotoxic T-lymphocyte–associated antigen 4 (CTLA4), CD25, and CD40 [8]. Recent studies have identified circulating autoantibodies which target the thyroid-stimulating hormone receptor and insulin-like growth factor-1 receptor on the orbital fibroblasts [9,10]. Further activation of inflammatory genes leads to overproduction of cytokines, chemoattractants, hyaluronan and glycosaminoglycan, which results in orbital inflammation in the early phase and tissue hypertrophy in the late phase [11]. The expression level of these gene associated with GD and inflammation would differ according to their genotype, and further study on genetic polymorphism in these genes would predict prognosis or complications in GD patients.

Treatment of GO varies according to the disease severity [12]. Establishing a euthyroid state and quitting smoking are helpful [13]. Active inflammatory phase often requires corticosteroids or other immunosuppressants to reduce orbital congestion [14]. Early intervention and close monitoring may hamper the progression and the severity of the disease [15]. It would be beneficial to find biomarkers early predicting severe GO in GD patients. Herein, all regions of CTLA4 including promoter, exon and 3’UTR were investigated to find the correlation of CTLA4 gene polymorphism and GO.

2. Experimental Section

2.1. Study Subjects

Twenty-two patients with GO (46.2 ± 16.5 years old, eight males and fourteen females) and 20 healthy controls (28.3 years old, 5 males and 15 females) were included in Linkou Chang Gung Memorial Hospital during the period from June 2017 to May 2019.

2.2. DNA Extraction

The blood samples were collected in EDTA-coated vacuum tubes. The genomic DNA was extracted using QIAamp® DNA Mini kit (Qiagen GmbH, Hilden, Germany) according to the manufacturer’s instructions. DNA concentration and purity were evaluated by measuring the optical density at 260 and 280 nm through UV spectrometer.

2.3. PCR Amplification

The PCR mixture contained 1 µL DNA, 10 µL HotStarTaq DNA Polymerase (Qiagen GmbH, Hilden, Germany), 1 µL forward primer (10 µM), 1 µL reverse primer (10 µM) and 12 µL ddH2O. Table 1 displayed the primers used in the present study. The PCR program for promoter and exon 1 was 1 cycle 95 °C for 10 min, 35 cycles of 94 °C for 30 sec, 65.5 °C for 30 sec, and 72 °C for 1 min. The PCR program for exon 2, exon 3, exon 4, and 3’UTR was 1 cycle 95 °C for 10 min, 35 cycles of 94 °C for 30 s, 59 °C for 30 s, and 72 °C for 1 min. The final elongation step was 3 min at 72 °C and then soaking at 10 °C. For gel electrophoresis visualization, 5 µL of the PCR products was pipetted onto a 1.5% agarose gel and run at 100 V for 20 min. Correct PCR products were visualized under UV illumination.

2.4. Purifying and Sequencing

The PCR production was purified by mixture containing 2.5 µL shrimp alkaline phosphatase and 0.05 µL exonuclease I (New England Biolabs, UK), and the purified PCR products were sequenced using ABI PRISM 3730 DNA Analyzer (Applied Biosystems, Foster City, CA, USA). Single nucleotide
polymorphisms (SNPs) analysis were performed on CTLA4 gene, and rs11571315, rs733618, rs4553808, rs11571316, rs62182595, rs16840252, rs5742909, rs231775, and rs3087243 were selected for genotyping.

### Table 1. Primers used for CTLA4 in this study.

| Primer | GC Content | Tm (°C) | Base Pair |
|--------|------------|---------|-----------|
| promotor and exon1 | | | |
| F: 5’ GGC AAC AGA GAC CCC ACC ACC GTT 3’ | 21/13 (62%) | 65.3 | 1234 |
| R: 5’ GAG GAC CTT CCT TAA ATC TGG AGA G 3’ | 25/12 (48%) | 65.8 | |
| F: 5’ CTC TCC AGA TTT AAG GAA GGT CCT C 3’ | 25/12 (48%) | 65.8 | |
| R: 5’ GGA ATA CAG AGC CAG CCA AGC C 3’ | 22/13 (59%) | 65.8 | 1170 |
| Exon2 and exon3 | | | |
| F: 5’ CAT GAG TTC ACT GAG TTC CC 3’ | 20/10 (50%) | 58.4 °C | 1034 |
| R: 5’ TAC CAC TCT CCT TCC TCT TC 3’ | 20/10 (50%) | 58.4 °C | |
| Exon4 | | | |
| F: 5’ CTA GGG ACC CAA TAT GTG TTG 3’ | 21/10 (48%) | 59.5 | |
| R: 5’ AGA AAC ATC CCA GCT CTG TC 3’ | 20/10 (50%) | 58.4 | 360 |
| 3’UTR | | | |
| F1: 5’ CAG CTA GGG ACC CAA TAT GTG TTG AG 3’ | 26/13 (50%) | 59.5 | |
| R1: 5’ GTG AAG TCA ACT CAG ATC CCA CCA GC 3’ | 26/13 (50%) | 59.5 | 1088 |
| F2: 5’ GCT TGG AAA CTG GAT GAG GTC ATA GC 3’ | 26/13 (50%) | 59.5 | |
| R2: 5’ AGA GGA AGA GAC ACA GAC AGA GTG GC 3’ | 26/13 (50%) | 59.5 | 1255 |

F: forward primer; R: reverse primer.

### 2.5. Statistical Analysis

All statistical data analyses were performed by the SPSS 17.0. Genotype and allele frequencies of CTLA4 gene for patients with GO were compared to healthy controls. Chi-square or Fisher’s exact test were used to evaluate the association between GO and CTLA4 SNPs.

### 3. Results

Among 22 GO patients, there were 14 females and 8 males, and the average age of patients was 46 years old. Moreover, half of GO patients was moderate in severity categories [16], and the most common symptom of GO patients was proptosis. The characteristics of patients were shown in Table 2.

### Table 2. Characteristics of patients (n = 22).

| Total, No. (%) | |
|----------------|------------------|
| Median age of the patients | 46.2 ± 16.5 |
| Sex of the patients | |
| Male | 8 (36.4) |
| Female | 14 (63.6) |
| Graves’ ophthalmopathy | |
| Mild | 7 (31.8) |
| Moderate | 14 (63.6) |
| Severe | 1 (4.6) |

The severity of GO was classified via the criterion proposed by Werner [6]. Class 0 to class 2 were classified as “mild”, class 3 and class 4 were classified as “moderate”, class 5 and class 6 were classified as “severe”.

All SNPs were in accordance with the Hardy-Weinberg equilibrium in the control group (p > 0.05) (Table 3). There was no nucleotide substitution in exon 2 and exon 3 of CTLA4 region, and the allele frequencies of CTLA4 polymorphisms had no significant difference between patients with GO and controls. However, the genotype frequency (Table 4) of “TT” genotype in rs733618 significantly differed between patients with GO and healthy controls (OR = 0.421, 95%CI: 0.290–0.611, p = 0.043), and the
“CC” and “CT” genotype in rs16840252 were nearly significantly differed in genotype frequency ($p = 0.052$). Haplotype analysis (Table 5) showed that CTLA4 Crs733618Crs16840252 might increase the risk of GO (OR = 2.375, 95%CI: 1.636-3.448, $p = 0.043$).

Table 3. Allele frequencies in patients and controls and odds ratio of developing Graves’ ophthalmopathy.

| SNP         | Position | Allele | Minor Allele Frequency (HWE) | Genotype Frequency | HWE p Value | Odds Ratio (95% CI) | $\chi^2$ p Value |
|-------------|----------|--------|-------------------------------|-------------------|-------------|---------------------|-----------------|
| rs11571315  | 203866178| C/T    | 0.181 (0.275)                | 0                  | 0.237       | 1.707 (0.607–4.799) | 0.308           |
| rs733618    | 203866221| T/C    | 0.409 (0.500)                | 8                  | 0.468       | 0.468 (0.136–1.611) | 0.226           |
| rs4553808   | 203866282| A/G    | 0.068 (0.175)                | 14                 | 2.139       | 2.139 (0.621–7.370) | 0.152           |
| rs11571316  | 203866366| A/G    | 0.114 (0.100)                | 0                  | 0.889       | 0.889 (0.190–4.150) | 0.152           |
| rs62182595  | 203866465| A/G    | 0.068 (0.175)                | 0                  | 0.421       | 0.421 (0.290–0.611) | 0.152           |
| rs16840252  | 203866796| C/T    | 0.068 (0.200)                | 0                  | 0.237       | 0.237 (0.052–1.073) | 0.152           |
| rs5742909   | 203867624| C/T    | 0.068 (0.175)                | 0                  | 0.800       | 0.800 (0.181–3.536) | 0.152           |
| rs231775    | 203867991| A/G    | 0.182 (0.275)                | 0                  | 1.154       | 1.154 (0.287–4.635) | 0.152           |
| rs3087243   | 203874196| A/G    | 0.114 (0.100)                | 0                  | 0.884       | 0.884 (0.190–4.150) | 0.152           |

HWE: Hardy-Weinberg equilibrium; 95%CI: 95% confidence interval.

Table 4. Statistical analysis of CTLA4 (SNPs).

| SNP         | Genotype Frequency (Patient (n)) | Genotype Frequency (Control (n)) | Odds Ratio (95% CI) | $\chi^2$ p Value |
|-------------|----------------------------------|----------------------------------|---------------------|-----------------|
| rs11571315  | CC 0 0                           | NA NA                            | NA NA               | NA              |
|             | CT 8 11                          | 0.468 (0.136–1.611)              | 0.226               |                |
|             | TT 14 9                          | 2.139 (0.621–7.370)              | 0.226               |                |
|             | CC 4 4                           | 0.889 (0.190–4.150)              | 1.000               |                |
| rs733618    | CT 18 12                         | 3.000 (0.736–12.227)             | 0.118               |                |
|             | TT 0 4                           | 0.421 (0.290–0.611)              | 0.043 **            |                |
|             | AA 19 13                         | 3.410 (0.742–15.677)             | 0.152               |                |
| rs4553808   | AG 3 7                           | 0.293 (0.064–1.348)              | 0.152               |                |
|             | GG 0 0                           | NA NA                            | NA NA               |                |
|             | AA 0 0                           | NA NA                            | NA NA               |                |
| rs11571316  | AG 5 4                           | 1.176 (0.267–5.176)              | 0.118               |                |
|             | GG 17 16                         | 0.850 (0.193–3.739)              | 0.152               |                |
|             | AA 0 0                           | NA NA                            | NA NA               |                |
| rs62182595  | AG 3 7                           | 0.293 (0.064–1.348)              | 0.152               |                |
|             | GG 19 13                         | 3.410 (0.742–15.677)             | 0.152               |                |
|             | CC 19 12                         | 1.22 (0.932–19.131)              | 0.052 *             |                |
| rs16840252  | CT 3 8                           | 0.237 (0.052–1.073)              | 0.052 *             |                |
|             | TT 0 0                           | NA NA                            | NA NA               |                |
|             | CC 19 13                         | 3.410 (0.742–15.677)             | 0.152               |                |
| rs5742909   | CT 3 7                           | 0.293 (0.064–1.348)              | 0.152               |                |
|             | TT 0 0                           | NA NA                            | NA NA               |                |
|             | AA 0 0                           | NA NA                            | NA NA               |                |
| rs231775    | AG 8 11                          | 0.468 (0.136–1.611)              | 0.226               |                |
|             | GG 14 9                          | 2.139 (0.621–7.370)              | 0.226               |                |
|             | AA 0 0                           | NA NA                            | NA NA               |                |
| rs3087243   | AG 5 4                           | 1.250 (0.283–5.525)              | 0.152               |                |
|             | GG 17 16                         | 0.800 (0.181–3.536)              | 0.152               |                |

95%CI: 95% confidence interval; NA: Not applicable. * indicates $p < 0.1$, ** indicates $p < 0.05$. 
Table 5. CTLA4 haplotypes and odds ratio of developing Graves’ ophthalmopathy.

| CTLA4 Haplotypes | Patient(n) | Control (n) | OR (95%CI) | p Value |
|------------------|------------|-------------|------------|---------|
| C<sub>rs733618</sub>C<sub>rs16840252</sub> | 22 | 16 | 2.375 (1.636–3.448) | 0.043 ** |
| C<sub>rs733618</sub>T<sub>rs16840252</sub> | 3 | 5 | 0.474 (0.097–2.307) | 0.445 |
| T<sub>rs733618</sub>C<sub>rs16840252</sub> | 18 | 16 | 1.125 (0.241–5.252) | 1 |
| T<sub>rs733618</sub>T<sub>rs16840252</sub> | 3 | 7 | 0.293 (0.064–1.348) | 0.152 |

OR: odds ratio; 95%CI: 95% confidence interval; NA: Not applicable. ** indicates p < 0.05.

Our data suggested that the genotype of −1722 (rs733618) in the promoter region of CTLA4 was associated with GO, the p value of TT vs. (CT + CC) was 0.043. We further combined −1722 (rs733618) and −1147 (rs16840252) together to observe the correlation between GO and controls, because the genotype of −1147 was very close to the critical point, p = 0.052. As a result, the two-marker combination was found to be more significantly related to GO (p = 0.025).

4. Discussion

Previous research had found that rs733618 [17], rs231775 [17], rs3087243 [18,19] and a two-marker combination containing rs733618 and rs16840252 [17] were related to GD. Our results showed that rs733618 was specifically related to GO. Comparing family-based research [17] to our results, only rs733618 in promoter was related to GO but not rs231775 in exon 1, and it indicated that major transcription level of CTLA4 was affected by SNPs in promoter region. Moreover, SNPs in exon 1 did not lead to significant reduced protein expression [20]. Therefore, rs733618 were associated with GD and GO, and “TT” genotype of rs733618 which was found to increasing the risk of GO in our study may affect major transcription level of CTLA4.

W. Tang et al., used expectation-maximization (EM) algorithm to create several feasible haplotypes, and haplotype compounded of G<sub>rs733618</sub>C<sub>rs16840252</sub>A<sub>rs231775</sub>G<sub>rs3087243</sub> was associated with gastric cardia adenocarcinoma (p = 0.012) [21]. Our results indicated that C<sub>rs733618</sub>C<sub>rs16840252</sub> was related to GO. Therefore, these two SNPs might be involved in both cancer and autoimmune disease. The immune dysfunction would be affected by CTLA4 genetic variation, and the risk of cancer and the development of autoimmune disease would be increased [22]. The rs16840252 mutation would increase the risk of cancer as well [23]. The most central function of CTLA4 is negative regulation of T cell activation, and it is expressed on the surface of activated T cells and competes to B7 with CD28 [24,25]. On the contrary, autoimmune disease would develop by reducing CTLA4 transcription level because of the immune responses are overreacting [26]. Investigation of rs733618 and rs16840252 in T cell activation would uncover the underlying mechanism in development of GO. In Table 6, disease association of rs733618 [23,27–34] and rs16840252 [35,36] were summarized.

Several studies in GD-related CTLA4 SNPs were summarized in Table 7. We found that those GD-related SNPs were located at different positions in the CTLA4 region within different populations. The majority of patients with GD from different ethnicities were associated with rs231775 and/or rs3087243 (Polish [37,38], Chinese [39–44], Saudi Arabian [45,46], Iranian [47,48], Italian [49], Japanese [50], and Taiwanese [17–19]). However, rs733618 and its relationship with GO have been found merely in Taiwan population. Thus, the correlation between GO and SNPs would diversely result from racial specificity, severity of GO and various inclusion criteria. Further investigation on racial specific GD-related SNPs associated with GO would be indicated.
Table 6. Summary of rs733618 and rs16840252-related disease.

| SNP        | Disease                              | Subjects and Results                                                                                                                                                                                                 | Ref.  |
|------------|--------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|-------|
| rs733618   | Systemic lupus                       | Asian: C allele was strongly associated with SLE and also CC genotype was significantly associated with the risk of SLE, \( p = 0.000 \).                                                                                | [23]  |
|            | Breast cancer                        | Chinese: CC genotype and C allele showed an increased risk of breast cancer \( (p = 0.030, \text{odds ratio (OR)} = 1.457, 95\% \text{CI 1.036–2.051}; p = 0.024, \text{OR} = 1.214, 95\% \text{CI 1.026–1.436}, \text{respectively}) \). | [27]  |
|            | Polycystic ovary syndrome             | Chinese Han population: significantly different between case and control groups in either genotypic or allelic distribution, \( p = 0.01 \) and \( 0.009 \), respectively.                                               | [28]  |
|            | Survival in patients with sepsis      | Adult Caucasian patients with sepsis: lower 90-day mortality was observed for T rs733618 A rs231775 A rs3087243 haplotype-negative patients than for patients carrying the TAA haplotype, \( p = 0.0265 \). | [29]  |
|            | Survival in patients with sepsis      | Unrelated Chinese Han population: GG genotype reduced the progression-free survival and the overall survival of patients with multiple myeloma who received bortezomib-based therapy, \( p = 0.002 \). | [30]  |
|            | Survival in patients with sepsis      | Non-small-cell lung cancer Chinese: T > C polymorphism was associated with the development of NSCLC in \( \geq 60 \) years and even drinking subgroups.                                                                 | [31]  |
|            | Survival in patients with sepsis      | Myasthenia gravis (MG) Chinese Han population: C allele were more frequent in MG patients, \( p = 0.042 \).                                                                                                          | [32]  |
|            | Urinary schistosomiasis               | Gabonese children: T allele and TT genotype were significantly overrepresented in the patient group, \( p = 0.001 \).                                                                                   | [33]  |
|            | Lymphatic filariasis (LF)             | Sarawak, Malaysia: CT genotype \( (p = 0.02) \) and those with combined minor allele C carriers \( (CT + CC; p = 0.01) \) exhibited a significantly decreased risk for LF.               | [34]  |
| rs16840252 | Colorectal cancer                     | Chinese: polymorphism was associated with an increased risk of colon cancer in homozygote model \( p = 0.040 \) and recessive model \( p = 0.037 \).                                                     | [35]  |
|            | Recurrent schizophrenia               | Chinese Han population: A significant association with schizophrenia, \( p \) (allele) = 0.0081, \( p \) (genotype) = 0.0117.                                                                             | [36]  |
Table 7. Summary of several studies for CTLA4 SNPs associated with Graves’ disease.

| SNP          | Subjects                              | Results                                                                 | Ref. |
|--------------|---------------------------------------|------------------------------------------------------------------------|------|
| rs733617     | Han population of Taiwan (family-based)| C allele over-transmitted to affected individuals ($\chi^2 = 6.714$, nominal $p = 0.0096$). | [17] |
| rs5742909    | Polish Caucasian                      | genotype and allele were differentially distributed ($p = 0.0002$; $p = 4.07 \times 10^{-5}$), and lack of the rare T allele increased GD in patients with familial autoimmune thyroid incidence ($p = 0.00005$). | [37] |
|              | Chinese                               | genotype frequencies of CT and allele frequencies of T were much higher in GD patients with ophthalmopathy than that in the group without ophthalmopathy ($p = 0.020$, $p = 0.019$). | [39] |
|              | Chinese                               | variant allele carriers might have decreased risks of GD when compared with the homozygote carriers TT + TC vs. CC: OR = 0.78, 95% CI = 0.62–0.97. | [44] |
| rs231775     | Han population of Taiwan (family-based)| CTLA4_+49_G/A ($p = 0.0219$), with its minor allele (G allele) over-transmitted to affected individuals ($\chi^2 = 5.252$, nominal $p = 0.0219$). | [17] |
|              | Taiwanese                             | significant differences in the frequencies of the genotypes and alleles, $p < 0.05$. | [18] |
|              | Polish populations                    | a significantly lower frequency of the AA genotype in the group of patients with clinically evident GO ($p = 0.02$, OR = 2.6). | [38] |
|              | Chinese                               | genotype GG and allele frequencies of G in patients with Graves’ disease were significantly increased as compared with control group ($p = 0.008$, $p = 0.007$). | [39] |
|              | Han population of Chinese (unrelated)  | allele G was significantly associated with GD in adults ($p < 0.001$) and children ($p = 0.002$). genotype GG ($p = 0.005$) and allele G ($p = 0.03$) were more prevalent in GD. | [40] |
| rs231775     | Chinese children                      | genotype and allele frequencies of children with GD differed significantly from those of the controls ($p = 0.0023$ and $p = 0.022$, respectively). | [42] |
|              | Saudi Arabian                         | G allele was more frequent in patients with GD than in the control group, $p = 0.003$. | [45] |
|              | Iranian                               | G allele was significantly higher in patients with Graves’ disease than in the control group (27.1% vs. 15.1%, OR = 2.096, 95%CI = 1.350–3.253 and $p < 0.01$). | [47] |
|              | Iranian                               | a significant increase of GG genotype and G allele was observed in patients ($p = 0.012$ and $p = 0.025$, respectively). | [48] |
|              | Italian                               | G allele frequency was significantly higher compared to control subjects ($p = 0.04$). | [49] |
|              | Caucasian and Asian                   | G allele vs. A allele, $p < 0.00001$; genotype: GG vs. AG + AA, $p < 0.00001$; GG + AG vs. AA, $p < 0.00001$; GG vs. AA, $p < 0.00001$; AG vs. AA, $p < 0.00001$. | [50] |
| rs3087243    | Taiwanese children                    | genotype GG was significantly associated with GD (OR = 1.71, 95% CI 1.20–2.44, $p = 0.006$); Allele G was significantly more frequent (OR = 1.61, 95% CI 1.18–2.19, $p = 0.0049$). | [19] |
|              | Han population of Chinese (unrelated)  | G allele was associated with susceptibility to Graves’ disease ($p = 0.011$). | [18] |
|              | Chinese children                      | G allele was significantly associated with GD in adults ($p < 0.001$) and children ($p < 0.001$). G allele was more prevalent in GD $p = 0.02$. | [40] |
|              | Southern China                        | G allele was significantly associated with an increased risk of GD development, $p < 0.001$. | [43] |
|              | Chinese                               | $G > A$ allele frequencies between the patient and control groups, $p = 0.014$. | [44] |
|              | Saudi Arabian                         | $A$ allele and the AA genotypes were significantly increased in patients with GD. | [45] |
|              | Italian                               | G allele was higher in GD patients than those in controls, $p = 0.004$. | [46] |
|              | Japanese                              | allelic frequency of the G allele was also significantly higher in patients with GD ($p = 0.02$). | [49] |
|              | Russian                               | for the TBII-positive GD, G allele carriers in patients had significant association with GD, OR = 2.97, 95%CI = 1.29–6.87, $p = 0.008$. | [51] |
|              | Russian                               | significantly higher frequencies of A allele and AA genotype and a lower proportion of G allele and GG genotype. | [52] |
Herein, Crs733618Crs16840252 of the CTLA4 promoter was shown to be associated with increasing risk of GO. Promoter polymorphism can influence gene transcription, as well as the disease severity [53]. The signal peptide is encoded by exon 1, and mutations in this exon suppress CTLA-4 protein expression [54]. Regulation of mRNA-based processes, such as mRNA localization, mRNA stability, translation, as well as regulation of protein features not encoded in the amino acid sequence are most commonly regulated by 3′UTR [55]. Consequently, there would be other significant SNPs located including but not limited to promoter region associated with GO. Further clinical application for prediction of GO in GD patients needs a feasibility study, and investigation of the mechanism of CTLA4 SNPs and haplotype of GO should be performed.

5. Conclusions

In conclusion, CTLA4 Crs733618Crs16840252 was found to be a potential marker for GO, and these haplotypes would be ethnicity-specific. Clinical application of CTLA4 Crs733618Crs16840252 in predicting GO in GD patients may be beneficial.

Author Contributions: D.-P.C. conceived and designed the experiments, and wrote draft of the manuscript. Y.-C.C. analyzed the data, contributed reagents/materials/analysis tools. Y.-H.W. reviewed literature. W.-T.L. performed the experiments. A.-L.H. analyzed all data. W.-T.W. reviewed and approved the final draft. All authors read and approved the final manuscript.

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