Research Article

TagSNP approach for HLA risk allele genotyping of Saudi celiac disease patients: effectiveness and pitfalls

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Background: Celiac disease (CD) is a genetically complex autoimmune disease which is triggered by dietary gluten. Human leukocyte antigen (HLA) class II genes are known to act as high-risk markers for CD, where >95% of CD patients carry (HLA), DQ2 and/or DQ8 alleles. Therefore, the present study was conducted to investigate the distribution of HLA haplotypes among Saudi CD patients and healthy controls by using the tag single nucleotide polymorphisms (SNP).

Methods: HLA-tag SNPs showing strong linkage value ($r^2 > 0.99$) were used to predict the HLA-DQ2 and DQ8 genotypes in 101 Saudi CD patients and in 103 healthy controls by using real-time polymerase chain reaction technique. Genotype calls were further validated by Sanger sequencing method.

Results: A total of 63.7% of CD cases and of 60.2% of controls were predicted to carry HLA-DQ2 and DQ8 heterodimers, either in the homozygous or heterozygous states. The prevalence of DQ8 in our CD patients was predicted to be higher than the patients from other ethnic populations (35.6%). More than 32% of the CD patients were found to be non-carriers of HLA risk haplotypes as predicted by the tag SNPs.

Conclusion: The present study highlights that the Caucasian specific HLA-tag SNPs would be of limited value to accurately predict CD specific HLA haplotypes in Saudi population, when compared with the Caucasian groups. Prediction of risk haplotypes by tag SNPs in ethnic groups is a good alternate approach as long as the tag SNPs were identified from the local population genetic variant databases.

Introduction

Celiac disease (CD) is an immune-related disorder of gastrointestinal system, which is triggered by ingestion of gluten peptide found in cereals like wheat, rye and barley. Originally, CD was thought to exclusively affect white Europeans [1], but recent reports indicate its increasing prevalence in diverse ethnic groups like Caucasians, Africans, Arabs and South Asians [2–6]. This increasing frequency of CD could be attributed to the rapid changes in lifestyle and diet and also due to the recent developments in diagnostic procedures. Recent studies indicate that the prevalence of CD ranges from 0.6% to 1.1% among the Middle Eastern arab countries [7]. Although CD is considered a major health problem in the Middle Eastern region, exact frequency of CD remains elusive due to the lack of large scale data [2]. Patients with classical CD presents gastrointestinal (GI) manifestations like diarrhea, malabsorption, abdominal...
pain and distension, bloating, vomiting, and weight loss [3,8–10]. Currently, gluten-free food is the standard dietary restriction to manage the disease complications [11].

The strongest genetic predisposing factors known to explain 25–40% of CD’s heritability are, human leukocyte antigen DQ (HLA-DQ) class II haplotypes, which are formed by variants in the highly polymorphic HLA -DQA1 and -DQB1 genes [12,13]. HLA genes encode cell surface receptors of most antigen presenting cells, which forms a cleft that binds to gliadin peptides. The genetics of the various HLA haplotypes that contribute to CD development is complex, as the disease risk is basically determined by the number and configuration of the DQA1 and DQB1 alleles. About 90–95% of CD patients share HLA-DQ2 heterodimer (encoded by HLA-DQA1*0501 and HLA-DQB1*0201 alleles) and the remaining patients carry HLA-DQ8 heterodimer (encoded by HLA-DQA1*0301 and HLA-DQB1*0302 alleles) [14]. It is extremely rare for individuals negative for both DQ2 and DQ8 risk alleles to develop CD [15]. Therefore, due to its very high negative predictive value, HLA typing has become a standard exclusion criteria in CD diagnosis [6,15,16]. Even though HLA-DQ haplotypes are major predisposing genetic factors, they are not sufficient to develop the disease because only 20–30% of the normal population carry these HLA-DQ variants [17,18]. This fact supports the contribution of other HLA and non-HLA genetic loci regions in CD predisposition [17,19,20].

Traditionally, CD linked HLA risk variants are genotyped by PCR-based HLA typing using sequence-specific oligonucleotide probes (SSOP), sequence-specific primers (SSP), and Sanger sequencing-based typing (SBT) methods. No doubt that these methods have improved the HLA typing, but several inherent limitations like time-consuming and expensive protocols, low throughput, unphased data and ambiguity of results limits their wide-spread use in molecular diagnosis. In this regard, Monsuur et al. has developed a simple, high-throughput allelic discrimination method to rapidly predict the DQ2.5, DQ2.2, DQ7, and DQ8 risk alleles using tagSNPs. This tag SNP approach has shown >94.0% predictive value for CD diagnosis, with >96.8% of sensitivity and >99.4% of specificity, when tested among European population [14]. This method was then eventually used for population screening to determine the prevalence of CD HLA risk alleles in few other ethnic groups [14,21–25].

To the best of our knowledge, data on the distribution of HLA locus gene variants and their relevance to CD diagnosis among Saudi population are limited. Therefore, our study is aimed to assess the utility of real-time PCR based tagSNPs to provide new information on HLA-DQ risk haplotypes associated with CD in Saudi Arabia. Moreover, the present study has also aimed to investigate the distribution of these HLA risk alleles among CD patients and healthy population in Saudi Arabia.

Materials and methods
Recruitment of study subjects and sampling
Ethical approval for this study was granted from the Research Ethics Committee, King Abdulaziz University Hospital (KAUH), Jeddah. Unrelated Saudi nationals with CD were recruited from a Pediatric Gastroenterology clinic; all cases were examined at the Department of Genetic Medicine for obtaining information about the prevalence of disease among other family members and the comorbidities. A total of 101 sporadic CD patients were clinically diagnosed based on the guidelines of European Society for Paediatric Gastroenterology, Hepatology and Nutrition (ESPHGAN) which includes serology testing for antibodies against gliadin and endomysium (EMA) or tissue transglutaminase (tTG) followed by small bowel biopsy to confirm the diagnosis for serology positive results [26].

Additionally, 103 healthy Saudi controls, who are over 20 years old, having no personal or family history for allergies, autoimmune or inflammatory disorders (Diabetes, Rheumatic arthritis, and Systemic Lupus Erythematosus) were randomly recruited from volunteers. All the enrolled participants and the parents of the children (<16 years) were informed about the study processes before obtaining written informed consent to participate in the present study. We collected 5 ml of peripheral blood samples in EDTA vacutainers from all participants and stored at −20°C until DNA extraction procedure was conducted.

Genotyping
Genomic DNA isolation
We isolated DNA from the 200 μl blood samples with QIAamp DNA Mini Kit (Catalogue # 51306). DNA Quality and quantity measurements were done using NanoDrop™ 2000c Spectrophotometer.

Genotyping for HLA-DQ tag SNPs
TaqMan® Genotyping assay (Applied Biosystems) was run by using 7500 FAST Real-Time PCR machine (Applied Biosystem, Int., U.S.A.) to genotype the individuals for the four HLA tag SNPs (rs7775228, rs2395182, rs2187668, and
### Table 1

Selected HLA TagSNPs associated with Celiac disease

| Detected HLA Haplotype | SNP ID | Gene       | Variant Type            | Chr. Position | Allele Change | TaqMan Assay ID |
|------------------------|--------|------------|-------------------------|---------------|---------------|-----------------|
| HLA-DQ2.2              | rs2395182 | HLA-DRA    | intergenic variant      | 6:32445540    | [G/T]         | C_29315313,10   |
|                        | rs7775228 | HLA-DQB1   | regulatory region variant| 6:32690302    | [T/C]         | C_11409966,10   |
| HLA-DQ8                | rs7454108 | NR         | regulatory region variant| 6:32713706    | [T/C]         | C_58662585,10   |
| HLA-DQ2.5              | rs2187668 | HLA-DQA1   | intronic                | 6:32638107    | [C/T]         | C_29817179,10   |

NR, not reported.

### Figure 1

Tag SNP positions with reference to HLA loci (in kb)

(DQ2.2) rs2395182 + rs7775228: \( r^2 = 0.971 \), (DQ2.5) rs2187668: \( r^2 = 0.994 \), (DQ8) rs7454108: \( r^2 = 0.892 \).

### Table 2

Interpretation of HLA tagSNPs to HLA-DQ alleles genotypes

| SNP          | Genotype | DQ2.5 Type | SNP          | Genotype | DQ8 Type |
|--------------|----------|------------|--------------|----------|----------|
| rs2187668    | CC       | DQx        | rs7454108    | CC       | DQx/DQx  |
| rs2187668    | CT       | DQ2.5/DQx  | rs7454108    | CT       | DQ2.5/DQx|
| rs2187668    | TT       | DQ2.5/DQ2.5| rs7454108    | TT       | DQx      |

- SNP          | Genotype | rs7775228/ DQ2.2 Type | SNP          | Genotype | rs7775228/ DQ2.2 Type |
|--------------|----------|------------------------|--------------|----------|-----------------------|
| rs2395182    | GG       | DQx                    | DQx          | DQx      |
| rs2395182    | GT       | DQx                    | DQ2.2/DQx    | DQ2.2/DQx|
| rs2395182    | TT       | DQx                    | DQ2.2/DQx    | DQ2.2/DQ2.2|

*DOX, not DQ2.2, DQ2.5, DQ8.

rs7454108 listed in Table 1 with chromosomal positions shown in Figure 1. TaqMan® Genotyper Software (Applied Biosystem, int. U.S.A.) was used to analyze scatter plots of test samples.

### HLA genotype notation using the tagSNPs

The genotype status of the tagSNPs is used to interpret HLA-DQ haplotypes as presented in Table 2. As per the original study published the Mansuur et al. [14], the homozygous (for major or minor allele) or heterozygous (both major and minor alleles) status of the queried tag SNP will predict the individual’s HLA-DQ haplotype status due to its strong Linkage Disequilibrium (LD) value seen among Caucasian population (\( r^2 \) value > 0.99).
Table 3 Comparison of HLA-DQ tagging SNPs allele frequencies

| rs ID   | Alleles | Frequency | OR   | 95% CI          | X2        | P-value |
|---------|---------|-----------|------|-----------------|-----------|---------|
|         |         | Cases (n=101) | Control (n=103) |                  |           |         |
| rs2395182 | Minor  | G         | 0.104 | 0.189           | 2.013     | [1.138--3.562] | 5.925 | 0.015* |
|         | Major   | T         | 0.896 | 0.811           |           |         |
| rs7775228 | Minor  | C         | 0.144 | 0.218           | 1.667     | [0.998--2.787] | 3.852 | 0.048* |
|         | Major   | T         | 0.856 | 0.782           |           |         |
| rs7454108 | Minor  | C         | 0.188 | 0.18            | 0.945     | [0.572--1.56]  | 0.049 | 0.824 |
|         | Major   | T         | 0.812 | 0.82            |           |         |
| rs2187668 | Minor  | T         | 0.1   | 0.03            | 4.074     | [1.615--10.273] | 10.157 | 0.001* |
|         | Major   | C         | 0.9   | 0.97            |           |         |

*Values are statistically significant P ≤0.05.

The determination of HLA-DQ2.5 and DQ8 haplotypes is straightforward and is predicted by the genotype status of rs2187668 and rs7454108 SNPs, respectively. Whereas DQ2.2 haplotype was determined based on the genotype status of 2 tagSNPs i.e., ‘T’ (major allele) for rs2395182 and ‘C’ (minor allele) for rs7775228. The homozygous statuses of TT (rs2395182) and CC (rs7775228) for both the tag SNPs suggest that the individual is homozygous to DQ2.2 haplotype. In case of DQ2.2 heterozygous haplotype, individuals will have either of the GT-CT, GT-CC or TT-CT genotype combinations for the rs2395182 and rs7775228 tag SNPs. The individuals who are not carrying either ‘T’ (rs2395182) or ‘C’ (rs7775228) alleles in the abovementioned combination were considered to be negative for DQ2.5/DQ2.2 and DQ8 haplotypes and carrying a different HLA-DQ haplotype (DQx), Table 2.

Validation of genotype calls
To check the accuracy and reproducibility of SNP genotyping assay, we performed Sanger sequencing of 100 random DNA samples from both CD patients and controls and compared the results between these two approaches. Prior to Sanger sequencing, the PCR products were purified using QIA quick PCR Purification Kit following the manufacturer instructions (Qiagen, Alameda, CA, U.S.A.). Purified PCR products were used as a DNA template for cycle sequencing reactions using ABI 3500 Genetic Analyzer (Life Technologies, U.S.A.). The reaction mixture of cycle sequencing PCR consists of 1 μl of big dye, 2 μl of 5× big dye buffer, 1 μl of either forward or reverse primer, and 1 μl of purified PCR product and 5μl nuclease-free water. Bioedit software 6 version was used for alignment and identifying the sequence variants.

Statistical analysis
To assess the CD risk conferred by different HLA genotypes, we conducted the analysis using Statistical Package for Social Sciences (SPSS) software version 14.0. Statistically significant difference in allele and genotypes was determined using Pearson’s standard chi-squared test, odds ratio (OR), and 95% confidence interval (CI) and P value <0.05 was considered significant.

Results
Clinical analysis
The present study included a total of 101 CD patients (45 males and 56 females), and the 103 healthy controls (43 males and 60 females). The mean age of patients was found to be 28.8 ± 13.9 years and for controls it was 31.6 ± 8.8 years. Of the study participants, 30% of the patients and 50% of the controls were born to consanguineous parents. The common clinical symptoms seen in CD patients were as follows: chronic diarrhea, abdominal pain, anorexia and abdominal distension. The common autoimmune manifestations seen in our patient group were type 1 diabetes mellitus (32%), autoimmune thyroiditis (8%), and systemic lupus erythematosus (3%). We also observed few non-autoimmune disease manifestations like osteomalacia (5%), seizure disorders (4%), and Down syndrome (4%).

Real time PCR genotyping results
HLA tagSNPs- allelic frequency distribution analysis
In Table 3, of the 4 HLA-tag SNPs tested, only three (rs2395182, rs7775228, and rs2187668) have shown the significant difference in minor allele frequency distribution among CD patients in comparison with healthy controls. Our statistical results for rs2395182 have indicated that the minor ‘G’ allele is more prevalent in healthy controls (18%)
controls and 24.75% CD patients were heterozygous for HLA-DRB1.

The predicted homozygous DQA1/DRB1-B2 (DQ2.2) haplotype was also frequent in control (5.82%) group than in CD patients (0.99%). A total of 29.13% of CD patients were heterozygous for the DQA1/DRB1-B2 haplotype between patients and control group, (32.67% vs 14.56% respectively, p-value = 0.024). No heterozygous DQA1/DRB1-B2 (DQ2.2) carriers were predicted by the tag SNP combinations in either patient or control groups. The DQB1 heterozygous haplotypes (DQB1/DQ2.2 and DQB1/DQX) were highly frequent among CD patients (33.66%) with significant difference in DQB1/DQX between patients and control group, (32.67% vs 14.56% respectively, p = 0.014).

Interestingly, the homozygous DQB1 high risk haplotype was predicted by the tag SNPs more frequently in control groups compared with the 3% of the healthy controls. This difference in minor allelic frequency is statistically significant (p-value = 0.001; OR = 4.074; 95% CI = 1.615–10.273). For rs7454108, there was no statistically significant difference between case and control groups (P-value > 0.5).

**HLA-DQ haplotype results**

The risk classification of the HLA-DQ genotypes (Table 4) is based on previous study from Saudi Arabia [27]. In our study, tag SNP predicted homozygous HLA-DQ2.5 haplotype is significantly high in CD patients compared to healthy controls (10.89 vs 2.91%; p = 0.024). No heterozygous DQ2.5 (in combination with DQ8 or DQ2.2) carriers were predicted by the tag SNP combinations in either patient or control groups. The DQ8 heterozygous haplotypes (DQ8/DQ2.2 and DQ8/DQX) were highly frequent among CD patients (33.66%) with significant difference in DQ8/DQX between patients and control group, (32.67% vs 14.56% respectively, p = 0.014). Surprisingly, the homozygous DQ8 high risk haplotype was predicted by the tag SNPs more frequently in control groups (7.76%) than in patients (1.98%). However, the difference was not statistically significant. The predicted homozygous HLA-DQ2.2 haplotype was also frequent in control (5.82%) group than in CD patients (0.99%). A total of 29.13% controls and 24.75% CD patients were heterozygous for DQ2.2. A total of 74 (36.27%) individuals predicted to be lacking all the high risk HLA alleles and seen as the extremely low risk group (DQX).

**Validation by sanger sequencing results**

The sequencing analysis with sanger method showed the accuracy (100%) of Real-time PCR based tagSNPs approach in determining the HLA-DQ haplotype among randomly selected individuals for each SNP genotypes from both study groups (Figure 2).

**Discussion**

In the present study, we assessed the transferability of the Real-time PCR based TaqMan SNP Genotyping Assay to accurately predict the HLA risk haplotypes associated with CD using four of the six tag SNPs in the Saudi population for the first time. Two HLA-tagSNPs (rs4713586 for DQ4 and rs4639334 for DQ7) were withdrawn from the present study because they were not polymorphic in the Saudi population (as per data from Saudi Human Genome Project -SHGP) and are not useful in tagging the targeted HLA haplotypes.

Our results shows that, in total 67.3% of Saudi CD patients were predicted to be the carriers of HLA-DRB1-associated major risk alleles. We observed that Saudi individuals with homozygous HLA-DQ2.5 haplotype have a 4-fold higher risk to develop CD (OR = 4.074). This finding further confirms previous studies, which revealed a high risk associated with two copies of DQ2.5 among Europeans, Africans and Arabs (Table 5) [17,22,24,27–31]. A recent study among Saudi children has reported that the homozygous DQ2.5 was seen in more CD cases than healthy controls. In that study, presence of either HLA-DQ8 or HLA-DQ2.2 alone did not confer a risk of CD in the Saudi children; however, the combination of DQ2.5 with either DQ8 or DQ2.2 significantly increases the disease risk in general population [27]. In contrast, no heterozygous DQ2.5 individuals have been predicted in this study. It can be explained by the low

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**Table 4 HLA-DQ Genotyping Frequency by TagSNP Method**

| Haplotype                  | Controls (n = 103) | Cases (n = 101) | OR       | CI         | P-Value  |
|----------------------------|-------------------|----------------|----------|------------|----------|
| Very high risk             | DQ2.5/DQ8         | –              | –        | –          | NA       |
| High risk                  | DQ2.5/DQ2.5       | (3) 2.91       | (11) 10.89| 4.07       | (1.1015–15.0691)| 0.024*   |
|                           | DQ2.5/DQ2.2       | –              | –        | –          | –        | NA       |
|                           | DQ8/DQ8           | (8) 7.76       | (2) 1.98 | 0.24       | (0.0497–1.1587)| 0.556    |
| Intermediate risk          | DQ2.5/DQX         | –              | –        | –          | –        | NA       |
| Low Risk                   | DQ8/DQ2.2         | (6) 5.83       | (5) 4.95 | 0.84       | (0.2486–2.8518)| 0.782    |
|                           | DQ8/DQX           | (15) 14.56     | (29) 32.67| 2.36       | (1.1771–4.7434)| 0.014*   |
| Very low risk              | DQ2.2/DQ2.2       | (8) 5.82       | (1) 0.99 | 0.16       | (0.0191–1.3678)| 0.578    |
|                           | DQ2.2/DQX         | (24) 23.3      | (20) 19.8| 0.81       | (0.4161–1.5875)| 0.543    |

NA: Not Applicable *Values are statistically significant P<0.05.
Table 5 HLA Risk haplotype for Celiac disease distribution among Middle Eastern countries

| Population | Groups | Highest | High | Intermediate | Low | Very Low | *Overall frequency (%) | Reference |
|------------|--------|---------|------|--------------|-----|----------|------------------------|-----------|
| Egypt      | Cases = 31 | 16.13 | 41.94 | 6.45 | 9.68 | 12.9 | 9.68 | NR | NR | 3.23 | 3.25 | 77.42 | 35.49 | Mohammed, M., et al. 2014 [31] |
| Iran       | Cases = 59 | 11.9 | 13.6 | 11.9 | 3.3 | 27.1 | 0 | 8.5 | 1.7 | 5.08 | 3.3 | 64.5 | 23.7 | Rostami-Nejad, M., et al. 2014 [22] |
| Israel     | Cases = 44 | 3.3 | 0.6 | 3.3 | 2.6 | 6.6 | 5.3 | 7 | 0 | 14.5 | 21.2 | 13.8 | 18.2 | Pallav, K., et al. 2014 [43] |
| Jordan     | Cases = 173 | 4.6 | NR | NR | NR | NR | NR | NR | NR | NR | NR | NR | NR | NR | NR | 57.8 | 61.8 | 22.5 | El-Akawi, Z., 2015 [44] |
| Libya      | Cases = 31 | NR | NR | 32 | 3 | 52 | 10 | NR | NR | NR | 3 | NR | 84 | 13 | Alarida, K., et al. 2010 [37] |
| Morocco    | Cases = 115 | 3.2 | NR | 7.7 | 12.2 | 36 | 4.5 | NR | 6.4 | 19.2 | 14.7 | 59.6 | 19.9 | Piancatelli, D., et al. 2017 [59] |
| Gaza Strip Palestine | Cases = 65 | 4.6 | NR | NR | NR | NR | NR | NR | NR | NR | NR | NR | NR | 7.9 | 70.8 | 15.4 | Ayesh, et al. 2017 [40] |
| Saudi Arabia | Cases = 101 | 3.1 | NR | NR | NR | NR | NR | NR | NR | NR | NR | NR | NR | 17.5 | 27.8 | Al-Hussaini, A., et al. 2019 [27] |
| Saudi Arabia | Cases = 103 | 0 | 10.89 | 0 | 1.98 | 0 | 3.96 | 29.7 | 0.99 | 16.83 | 35.6 | 10.89 | 35.64 | Present study |
| Gaza Strip Palestine | Cases = 110 | 0 | 2.91 | 0 | 7.76 | 0 | 3.88 | 15.5 | 5.82 | 17.47 | 46.6 | 2.91 | 27.14 | Bouguerra, F., et al., 1997 [45] |
| Turkey     | Cases = 78 | 28.2 | 35.8 | NR | 11.5 | 15.3 | NR | 6.4 | NR | NR | 2.6 | 78.5 | 46.1 | Çakır, M., et al. 2014 [46] |
| Control    | n = 103 | 0 | 2.91 | 0 | 7.76 | 0 | 3.88 | 15.5 | 5.82 | 17.47 | 46.6 | 2.91 | 27.14 | Bouguerra, F., et al., 1997 [45] |
| Control    | n = 13 | 15.3 | 38.4 | NR | 15.3 | 7.6 | NR | 15.3 | NR | NR | NR | 61.3 | 45.9 | Present study |

NR, Not Reported/ specific allele statues is not clear in the published articles.
* Overall frequency for (a) DQ2 encoded by DQA1*0501 and DQB1*02 alleles either in homozygous or heterozygous state. (b) DQ8 encoded by DQA1*0301 and DQB1*0302 alleles either in homozygous or heterozygous state.
Figure 2. Sanger sequencing results for the 4 HLA tagSNPs used in the present study

(A) rs2395182

(B) rs7775228

(C) rs7454108

(D) rs2187668*

* No heterozygous form was detected for SNP (rs2187668).

minor allele frequency for the predictive 'T' allele of the tagging rs2187668 SNP in this population (0.1%, 0.03%) in CD patients and control respectively, (Table 3).

Based on our HLA predictions and from previous studies, it is clear that the frequency of the HLA haplotypes in Arabs is not necessarily different from Europeans, but tagSNPs used for Caucasian population may not accurately predict the HLA genotype in this population. Significant differences in the patterns of linkage disequilibrium in the genomes of Saudis with the highest consanguineous marriages in the world might explain limited benefit of using the same tagSNPs to predict the heterozygote HLA-DQ2.5 allele. Several studies have shown that social factors such as the differences in consanguinity levels in populations is responsible for changes in genotype frequencies and results in the loss of heterozygosity with increasing homozygous genotypes [32–34]. First cousin marriages among Arabs is high and the overall rates of consanguinity in Saudi Arabia ranges from 52.1% to 67.7% for many generations in the past [34,35]. The frequency of consanguinity within our study is 30% in CD patients and 50% in controls.

We found that the most frequent predicted haplotype among Saudi CD patients was DQ8 (33.66%) in heterozygous form with a statistically significant P<0.014 for DQ8/DQX that increases the possible risk by more than 2 folds than the general population (OR = 2.36). Similar high frequency of HLA-DQ8 is seen in CD patients and Amerindian groups in Chile [36]. Our findings showed a significant difference of predicted DQ8 among CD patients in comparison with other Arab population listed in Table 5, as well as other populations such as Cameroon, Italy, Hungary, United States, Finland and Japan [23,37] with low frequency of heterozygotes. Furthermore, among healthy Saudi children fewer heterozygote DQ8/DQX and homozygous DQ8 haplotypes were reported [27]. This finding suggests that the tag SNP rs7454108 in our study predicting excessively more DQ8 haplotype in the Saudi population than the other studies which used classical HLA genotyping methods.

Previous study reported about 17–20% of the general Saudi population carry HLA-DQ8, higher than in the Caucasians, 1–9% [35,38]. These differences propose that variable combinations of HLA-DQ risk alleles among Saudi CD patients might confer different risk gradients for some HLA-DQ molecules compared to Caucasian CD patients. Although homozygous DQ8 haplotype is considered to be strongly CD-associated high risk molecule [39], surprisingly in our population, it was seen more frequently predicted in healthy controls (7.76%) than in CD patients (1.98%).
Such differences, though not significant, were seen in Finnish, Hungarian and in some Arab population like Gaza strip [23,40].

The *DQ2.2* haplotype is a low risk haplotype in CD patients in many countries [21,24,41,42]. In our study, predicted homozygous *DQ2.2* was more frequent in control group (5.82%) than in cases (0.99%). Although *DQ2.2* is known to raise the risk for CD when associated with *DQ2.5* or *DQ8*, it did not confer the high risk for CD in our population. This low frequency of predicted *DQ2.2* among CD patients may suggest it plays a minor role in triggering the autoimmune process in our population [42]. This finding was also supported by Al-Hussaini’s study that *DQ2.2* alone did not confer the risk for CD in the Saudi children [27]. This might also suggest the protective role of *DQ2.2* allele in the Saudi population, which requires to be tested in a larger study. High frequency of *DQA1* \( ^{\ast}02:01 \) allele that is associated with HLA-*DQ2.2* haplotype in Santiago, Chile among control subjects suggests that they protect the population against the CD development [36]. However, it was not possible in our study to determine whether predicted *DQ2.2* individuals were carrying the *DQA1* \( ^{\ast}02:01 \) allele or not because of the low frequency of the predictive minor allele of tagSNPs in this population.

To date, most HLA data on CD patients studies has come from only few Arab countries, that too on a smaller sample size. In Table 5 our predicted HLA genotype frequency among CD patients and across different Arab countries shows some similarities as well as differences.

The present study has provided information on the predicted HLA genetic background of CD in Saudi population. The differences in HLA’s association with CD as observed in the present study population compared to non-Arab populations could be due to the different ethnic and cultural practices like first cousin marriages, which will in turn influence the polymorphic nature of SNPs. The present study also suggests that transferability of tagSNP approach in populations (like Arabs, African, Japanese and Chinese etc.) which have known differences in LD structure, still needs to be determined. Therefore, an immediate search for other tag SNPs with higher \( r^2 \) value for disease association needs to be identified using population specific genetic database such as SHGP.

In conclusion, tagSNP typing is a reliable and easy alternate approach to rapidly genotype highly polymorphic HLA region. The present study represents the first investigation to test the applicability of tag SNPs to determine the HLA status of CD patients in Saudi population. Our findings reveal that, tag SNPs predicted homozygous *DQ2.5* and heterozygous *DQ8* haplotypes, of HLA are associated with CD development among Saudi patients. The findings of this study highlight that Caucasian specific tagSNPs would be of limited value to accurately predict CD specific HLA haplotypes in Saudi population. More than 32% of the CD patients were predicted to be not carrying the HLA risk alleles, highlighting the low predictive value for them in Saudi population. Large-scale HLA typing of Arab CD patients with different highly polymorphic population specific tagSNPs might reveal the accurate picture of HLA risk haplotypes in the disease diagnosis and treatment.

**Data Availability**

The data generated by us are presented in the form of tables and figures in the manuscript. Individual participant SNP genotypes or HLA haplotypes cannot be released due to the Institutional ethical committee rules and regulation to protect the privacy of the participants and to maintain the confidentiality of their clinical information. All the pooled data is presented in the manuscript to protect the privacy of the participants and maintain the confidentiality of their personal data.

**Competing Interests**

The authors declare that there are no competing interests associated with the manuscript.

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**Author Contribution**

Reham H. Baaqeel, Ramu Elango and Noor Ahmad Shaik conceived and designed the study. Funding acquisition was conceived by Reham H. Baaqeel, Ramu Elango. The participant recruitment and clinical data was collected by Omar I. Saadah, Bakr H. Al-hussaini, Meshari A. Alafan and Yagoub Bin-Taleb. The methodology was done by Reham H. Baaqeel, Mohammed A. Salama, Hadiah Bassam Al Mahdi. Statistical analysis was prepared by Mohammed A. Salama, Ramu Elango. The validation of the results was analyzed by Reham H. Baaqeel, Hadiah Bassam Al Mahdi, Ramu Elango and Noor Ahmad Shaik. The original draft was prepared and edited by Reham H. Baaqeel and reviewed by Ramu Elango and Noor Ahmad Shaik. The study was done under the supervision of Omar I. Saadah, Ramu Elango and Babajan Banaganapalli. This project was administrated by Jumana Yousuf Al-Aama.
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Abbreviations
χ2, Chi-square; bp, Base Pair; CD, Celiac Disease; CI, Confidence Interval; DGP, De-amidated Gliadin Peptide Antibody; DNA, Deyoxyribonucleic Acid; dNTP, Deyoxyribonucleotide Triphosphate; EDTA, Ethylenediaminetetraacetic Acid; EMA, Anti-endomysium; ESPGHAN, European Society for Paediatric Gastroenterology, Hepatology and Nutrition; GFD, Gluten Free Diet; GI, Gastrointestinal; GWAS, Genome-Wide Association Studies; HLA, Human Leukocyte Antigen; IgA/IgG, Anti-Gliadin Antibodies; KAUiH, King Abdulaziz University Hospital; LD, Linkage Disequilibrium; MHC, Major Histocompatibility Complex; OR, Odds Ratio; PCR, Polymerase Chain Reaction; SHGP, Saudi Human Genome Program; SNP, Single-Nucleotide Polymorphism; SS, Statistical Package for the Social Sciences; tTG, Anti-Tissue Transglutaminase.

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