Screening of celiac disease in children and adolescents with type 1 diabetes mellitus

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Abstract

Background and aim: Celiac disease is the most frequent autoimmune disease in Type 1 diabetes mellitus. We aimed to determine the celiac disease prevalence in patients with type 1 diabetes mellitus, additionally to evaluate the clinical, serological and molecular characteristics of type 1 diabetes mellitus and celiac disease which have a common genetic predisposition.

Material and methods: A total number of 76 type 1 diabetes mellitus patients aged between 1-18 years, were evaluated retrospectively. Serologic screening for celiac disease was performed via anti-tissue transglutaminase and anti-endomysial antibodies. Presence of human leukocyte antigens (HLA) (DQ2 and DQ8) documented as well. Patients with positive tissue transglutaminase and endomysial antibodies underwent endoscopic biopsy. Histopathological analysis were performed according to the modified Marsh classification. Patients' demographic characteristics, anthropometric measurements, physical examination, laboratory findings, age at type 1 diabetes mellitus and celiac disease onset, and celiac disease prevalence were evaluated. In addition all findings were compared between type 1 diabetes mellitus patients and newly diagnosed celiac disease patients.

Results: Serum tissue transglutaminase was positive in 14.5% (n=11) of all patients and serum endomysial antibodies was positive in 13.2% (n=10). The overall prevalence of celiac disease in type 1 diabetes mellitus was confirmed as 10.5% (n=8) by histopathological examination in present study. Of the celiac disease patients 37.5% were asymptomatic. In addition, 6 were anti-tissue transglutaminase and 7 were endomysial antibodies positive. Moreover, 60.3% (n=41) were HLA-DQ2 and 58.8% (n=40) were HLA-DQ8 positive. Selective IgA deficiency was described in 3 cases. In one of them HLA-DQ2/HLA-DQ8 was found positive and celiac disease diagnosis was confirmed by biopsy. HLA-DQ8 ratio was found significantly higher in patients with celiac disease than the type 1 diabetes mellitus patients. In addition, HLA-DQ2/DQ8 positivity was observed in 62.5% (n=5) of celiac disease patients.

Conclusion: Our findings have demonstrated the increasing prevalence of celiac disease in children with type 1 diabetes mellitus. Particularly the higher risk of asymptomatic celiac disease in type 1 diabetes mellitus patients, revealed the diagnostic value of serological screening. Furthermore, increased HLA-DQ2 and HLA-DQ8 positivity, which are more prominent in case of selective IgA deficiency, clearly demonstrates the requirement of routine total IgA and HLA analysis in serological screening.

Key words: type 1 diabetes mellitus, celiac disease, serological screening, human leukocyte antigens
Введение
Цель: Целиакия является наиболее частым наследственным заболеванием при сахарном диабете 1 типа. Целью нашего исследования стало определение распространенности целиакии у пациентов с сахарным диабетом 1 типа, а также оценка клинических, серологических и молекулярных характеристик сахарного диабета 1 типа и целиакии, которые имеют общую генетическую предрасположенность.

Материалы и методы: Всего 76 больных сахарным диабетом 1 типа в возрасте от 1 до 18 лет были включены в исследование. Серологический скрининг на антитела против трансглутаминазы и антитела к эндомизию проведен по модифицированной классификации Марша. Оценены демографические характеристики, антропометрические показатели, клиническое обследование, лабораторные данные, возраст при появлении сахарного диабета 1 типа и целиакии, а также распространенность целиакии. Кроме того, все результаты были сопоставлены между больными сахарным диабетом 1 типа и пациентами с патологией, диагностированной по консенсусному Consensus.

Результаты: Анализ сыворотки крови на антитела против трансглутаминазы был положительный у 14,5% (n=11) всех пациентов, аальный анализ проведен по моделированной классификации Марша. Оценены демографические характеристики, антропометрические показатели, клиническое обследование, лабораторные данные, возраст при появлении сахарного диабета 1 типа и целиакии, а также распространенность целиакии. Кроме того, все результаты были сопоставлены между больными сахарным диабетом 1 типа и пациентами с патологией, диагностированной по консенсусному Consensus.

Заключение: Наши результаты показали растущую распространенность целиакии у детей с сахарным диабетом 1 типа. В частности, более высокий риск возникновения бессимптомной целиакии у больших сахарным диабетом 1 типа выявил диагностическую ценность се- рологического скрининга. Кроме того, повышенная позитивность HLA-DQ2 и HLA-DQ8, которая более выражена в случае изолированной целиакии, также подтверждает необходимость проведения стандартного анализа общего IgA и HLA при сахарном диабете 1 типа.

Ключевые слова: сахарный диабет 1 типа, целиакия, серологический скрининг, человеческие лейкоцитарные антитела.
determine the CD prevalence in T1DM-patients, additionally to compare the clinical, serological and molecular characteristics of patients with T1DM and CD.

Material and methods

This study was performed with the Institutional Review Board protocol approval date 18/02/2019 and number 2019-29 in Istanbul Training and Research Hospital, Department of Gastroenterology, Hepatology and Nutrition, 01 January 2017 - 30 June 2018. A total number of 76 T1DM-patients aged between 1-18 years, were evaluated retrospectively. During application, height and weight percentiles of cases, as well as the body mass index (BMI) for patients over two years of age were determined.

Sero logical screening for CH was performed using anti tissue transglutaminase antibodies (tTGAb) via enzyme-linked immunosor bent assay (ELISA). All patients also screened for anti-endomyosal antibodies by the immunofluorescence method using human umbilical cord and fluorescein isothiocyanate conjugated anti human-IgA (INOVA, San Diego, Calif., USA). In case of serum IgA deficiency, total serum IgA levels were measured by nephelometry. Presence of HLA antigens (DQ2 and DQ8) determined via polymerase chain reaction (PCR) technique through genomic DNA isolated from the all patients’ peripheral blood. Patients with positive tTGA and EmA underwent endoscopic biopsy from distal duodenum. Histopathological analysis were performed according to the modified Marsh classification [11]. All CD-confirmed patients were referred for a gluten-free diet and followed.

In addition, cell blood count analysis was performed on patients’ venous blood samples. Haematological parameters were analysed using a haematology analyser (Cell-Dyne 3700, Abbott, Abbott Park, IL, USA). Biochemical analysis performed from serum samples by electro-chemiluminescence immunoassay on Beckman Coulter Unicel DXI 800 analyzer. Serum glycosylated hemoglobin (HbA1c) level <7% as well as moderate metabolic control, between 7-9% as moderate metabolic control and 9% as poor metabolic control.

Patients’ demographic characteristics, age at T1DM and CH onset, anthropometric measurements, physical examination, imaging and laboratory findings, were evaluated. In addition all findings were compared between T1DM-patients and newly diagnosed CH-patients.

Statistical analysis

All the data were analysed with SPSS (Statistical Package for the Social Sciences) software for Windows (v21.0; IBM, Armonk, NY, USA). Individual and aggregate data were summarized using descriptive statistics including mean, standard deviations, medians (min-max), frequency distributions and percentages. Normality of data distribution was verified by Kolmogorov-Smirnov test. Comparison of the variables with normal distribution was made with Student t-test. The variables which were not normally distributed, the Mann Whitney and Kruskal Wallis tests were conducted to compare between groups. Evaluation of categorical variables was performed by Chi-Square test. P-Values of <0.05 were considered statistically significant.

Results

The 76 T1DM-patients included in this study were 45 (59,2%) female, 31 (40,8%) male, and the mean age of symptom onset was 139,16±46,99 months (Ranged=40-120) in our sample group. Additionally, the mean age of total participants was 95,67±52,85 months (Ranged: 0-215) at the time of DM diagnosis. Six of the cases (7,9%) were screened due to the CD history in family members, and one of these patients was diagnosed with CD. The overall prevalence of CD in T1DM was confirmed as 10.5% (n=8) by histopathological examination in present study. The mean age of CD-patients was 109,20±56,10 months (Ranged: 78-188) at the time of CD diagnosis (Table 1).

The most common symptom reported in our patients was abdominal pain with a ratio of 32,9% (n=25) followed by constipation (25,0%), and diarrhea (11,8%) respectively. The 46,1% (n=35) of all the patients and 37,5% (n=3) of the CD patients were asymptomatic (Table 2).

Table 1

| Symptoms                  | n (%) | Age (Mean±SD) | P-value |
|---------------------------|-------|---------------|---------|
| Male                      | 31 (40,8) | 142,90±50,40 | 0,485   |
| Female                    | 45 (59,2) | 136,58±44,88 |         |
| Mean Age at Application   | 76 (%100) | 139,16±46,99 |         |
| Mean Age at DM Diagnosis  | 76 (%100) | 95,67±52,85  |         |
| Mean Age at CD Diagnosis  | 8 (%1,0)  | 109,20±56,10 |         |

SD=Standard Deviation.

According to the evaluation of anthropometric measurements obtained from all cases; mean height was 146,20±20,47 cm, mean weight was 44,09±18,39 kg, and mean BMI was 19,08±4,75 (Ranged=12,0-38,0). In addition, laboratory outcomes are represented in Table 3. There were no statistically significant differences found between the CD and T1DM groups according to the age, BMI and laboratory findings (p>0,05) (Table 3). Additionally, vitamin D deficiency was detected in 88,2% (n=67) of patients, folic acid deficiency in 1,3% (n=1) and vitamin B12 deficiency in 9,2% (n=7). Iron-deficiency anemia detected in 1 patient and selective IgA deficiency was detected in 3 (3,9%) patients. Only 2.7% of patients (n=2) had normal levels of glycosylated hemoglobin; well-controlled HbA1c was 10,5%, moderate controlled HbA1c was 40,8% and poorly controlled HbA1c was 48,7%.

In addition, anti-tTGA was positive in 14,5% (n=11), anti-transglutaminase IgG (anti-tTG) in 17,1% (n=13), anti-EMA in 13,2% (n=10) and antiendomisyum IgG (anti-EMG) in 7,9% (n=6) of all patients. Of the CD patients, 6 were anti-tTGA and 7 were EMA positive.

The distribution of patients according to the tissue groups analysis (n=68); 60,3% (n=41) were HLA-DQ2 and 58,8% (n=40) were HLA-DQ8 positive. Moreover, rates of HLA-DQ8, anti-tTGA, anti-tTG, anti-EMA and anti-EMG were found significantly higher in patients with CD than the T1DM patients (Table 4).
Clinical characteristics of celiac patients are presented in Table 5. Selective IgA deficiency was described in 1 CD-case. Anti-tTGA and EMA negativity was observed in a CD-patient due to the selective IgA deficiency, then HLA-DQ2/HLA-DQ8 was found positive in this patient. In addition, HLA-DQ2/DQ8 positivity was observed in 62.5% (n=5) of CH patients. HLA-DQ8 was found to be positive in all CD patients except a single patient with missing data. While a high rate of vitamin D deficiency was detected in the CD-cases (75%), B12 deficiency was detected in only one CD-case. All patients with positive laboratory findings for CD underwent duodenal biopsy by gastroscopy and after histopathological evaluation; 2 were identified as Marsh Type 2, 2 were Marsh Type 3b and 2 were Marsh Type 3c.

Table 3
Comparison of clinical features and laboratory outcomes between CD and DM patients.

|                      | DM Group (Mean±SD) | CD Group (Mean±SD) | P-value |
|----------------------|--------------------|--------------------|---------|
| Age                  | 140.56±47.04       | 127.25±47.95       | 0.515   |
| BMI                  | 19.29±4.80         | 16.83±3.76         | 0.146   |
| HbA1c (%)            | 9.10±2.06          | 9.61±2.33          | 0.542   |
| Hemoglobin (g/dL)    | 13.45±1.77         | 12.88±1.04         | 0.115   |
| MCV (fL)             | 80.74±7.07         | 82.32±6.57         | 0.747   |
| Iron (ug/dL)         | 81.01±3.51         | 63.96±3.51         | 0.189   |
| Vitamin B12 (pg/mL) | 46.06±231.47       | 40.53±170.53       | 0.602   |
| Folate (mg/L)        | 11.22±3.51         | 13.56±5.69         | 0.291   |
| Vitamin D (ng/ml)    | 17.50±8.22         | 22.33±3.51         | 0.231   |
| Total IgA (mg/dL)    | 15.04±66.94        | 165.31±82.75       | 0.564   |

Table 4
Results of serological screening in DM and CD patients.

|                      |          | DM Group (Mean±SD) | CD Group (Mean±SD) | P-value |
|----------------------|----------|--------------------|--------------------|---------|
| TG IgA (mg/dL)       | -        | 8.08±33.86         | 171.46±127.28      | 0.000*  |
| (Mean±SD)            |          |                    |                    |         |
| TG IgG (mg/dL)       | -        | 6.86±14.35         | 46.20±37.63        | 0.001*  |
| (Mean±SD)            |          |                    |                    |         |
| EMA-IgA              | Negative | 65 (%95.6)         | 1 (%12.5)          | 0.000*  |
| n (%)                |          | 3 (%4.4)           | 7 (%87.5)          |         |
| EMA-IgG              | Negative | 67 (%98.5)         | 3 (%37.5)          | 0.000*  |
| n (%)                |          | 1 (%1.5)           | 5 (%62.5)          |         |
| HLA-DQ2              | Negative | 25 (%41.0)         | 2 (%28.6)          | 0.420   |
| n (%)                |          | 36 (%59.0)         | 5 (%40.4)          |         |
| HLA-DQ8              | Negative | 28 (%45.9)         | 0 (%0.0)           | 0.019*  |
| n (%)                |          | 33 (%54.1)         | 7 (%100.0)         |         |

*= p<0.05 statistically significant.

Discussion
The risk of CD is significantly associated with female gender, young age and DM diagnosis especially at early ages [3]. In addition, patients mostly diagnosed with T1DM (75-80%) before the onset of CD. Classic CD disease often diagnose aged between 2 to 3 years, while the mean age of T1DM comorbid CD is about 8 years at diagnosis [12]. Rami et al. reported a mean age of T1DM diagnosis as 6.5 ± 4.1 years and CD diagnosis as 10.0 ± 5.4 years in 98 children [13]. Similarly, Deja et al. reported a mean age of 7.39 years for T1DM and 8.43 years for CD diagnosis in 27 children. Researchers also documented no statistically significant differences according to the gender[14]. In accordance with these data, the mean age at diagnosis of DM was 95.67±52.85 months and CD was 109.20±56.10 months except a single patient with missing data. While a high rate of vitamin D deficiency was detected in the CD-cases (75%), B12 deficiency was detected in only one CD-case. All patients with positive laboratory findings for CD underwent duodenal biopsy by gastroscopy and after histopathological evaluation; 2 were identified as Marsh Type 2, 2 were Marsh Type 3b and 2 were Marsh Type 3c.
in our study. Additionally, there was no significant differences found according to the gender.

In recent years it has been reported that the prevalence of CD in patients with T1DM has increased in worldwide and ranged from 4.4% to 11.1% in studies both with children and adults[12]. The CD prevalence in T1DM ranging from 4.4 to 6.4% in European countries and 10.5-11.1% in South America and India [15]. Ergür et al. documented a CD prevalence of 7.8% in 38 children with T1DM[16]. Supportively, the overall prevalence of CD in T1DM was found as 10.5% in present study.

The clinical presentation of the CD in T1DM is generally silent. Joshi et al. found positive TGA in 11 of 71 screened children, and 5 were (7.0%) diagnosed with CD histopathologically. The researchers reported that the majority of CD-positive patients (64%) were asymptomatic and frequently reported symptoms were puffiness, abdominal pain, and diarrhea. Moreover, researchers reported 45.5% vitamin D deficiency and 54.5% iron deficiency anemia in CD patients [17]. Tsouka et al. revealed 41-undiagnosed CD patients as a result of 771 serologically screened children with T1DM between 2005 and 2011 years. In same study 21 of 41 patients were asymptomatic and 10 of them had vitamin D deficiency [18]. Similarly in our study the 46.1% of the all patients and 37.5% of the CD patients were asymptomatic. The most common symptom reported in our patients was abdominal pain with a ratio of 32.9% followed by constipation (25.9%), and diarrhea (11.8%) respectively. There were no statistically significant differences found between the CD and T1DM groups according to the age, BMI and laboratory findings. Vitamin D deficiency was detected in 88.2% of all patients, folic acid deficiency in 1.3% and vitamin B12 deficiency in 9.2%. Iron-deficiency anemia detected in only 1 patient and a high rate of vitamin D deficiency was detected in the CD-cases (75%).

Although histopathological evaluation for CD diagnosis in T1DM patients is the gold standard, serological screening via TGA and EMA is highly valuable particularly in detecting asymptomatic patients. As a result of serologic screening in 268 patients with T1DM, Ozdemir et al. reported 8 EMA positive, 13 TGA positive patients and normal IgA levels in all patients. Researchers confirmed CD diagnosis in 5 (1.9%) patients. They revealed positive EMA and TGA in all of these 5 patients, additionally reported 100%, 100% sensitivity, and 95%, 86% specificity for these two tests respectively [19]. Yildirmaz et al. reported 16 (7.3%) TGA positive cases in 218 T1DM patients and CD diagnosis confirmed in 11 (5%) of these TGA positive cases. As a result of histopathological evaluation; 3 were identified as Marsh Type 2, 1 were Marsh Type 3a, 4 were Marsh Type 3b and 3 were Marsh Type 3c in same study [20]. Bolad et al. documented 10,1% TG and 7,2% EMA positivity in a study comparing 69 T1DM patients with healthy controls. Researchers also noted significantly higher TGA-G titres than healthy controls [21]. Singh et al. detected positive CD serology in 43 (34.1%) of 126 T1DM patients. The CD diagnosis confirmed in 17 patients (13.5%) by histopathological examination [22]. Supportively in present study, anti-tTGA was positive in 14,5%, and anti-EMA was positive in 13,2% of all patients. Of the CD patients, 6 were anti-tTGA and 7 were EMA positive. According to the histopathological evaluation of CD patients; 2 were identified as Marsh Type 2, 2 were Marsh Type 3b and 2 were Marsh Type 3c.

HLA plays a prominent role in the genetic predisposition to celiac disease, supportively there is a strong association between HLA-DQ2/DQ8 and CD. This relationship is so strong that CD is rarely found in individuals with HLA-DQ2 and HLA-DQ8 negative. HLA-DQ2 is responsible for the prevalence of CD in T1DM[23]. Ghawil et al. detected 75% HLADQ2, 21% HLA-DQ2 / HLA-DQ8 and 4% HLA-DQ8 in 24 CD patients with T1DM (n=218) [24]. Dezsofi et al. defined DQ2 / DQ8 heterozygosity as a risk factor for both CD and T1DM in a study consisted of 80 children with T1DM, 100 children with CD, and 47 children with CD+T1DM. Researchers stated that homozygous HLA-DQ8 genotype in T1DM patients and HLA-DQ2 / DQ8 heterozygosity in CD+T1DM patients were significantly higher than CD- patients and healthy controls [25]. In addition, HLA-DQ8 was the most frequently detected genotype in a study conducted by Ergür et al. among CD patients [16]. In accordance with these data, HLADQ2 was positive in 60.3% and HLA-DQ8 was positive in 58.8% of all patients in our study. HLA-DQ8 ratio was significantly higher in patients diagnosed with CD than the T1DM patients. Moreover, HLADQ2/DQ8 positivity was observed in 62.5% (n=5) of CD patients. Additionally, HLA-DQ8 was found to be positive in all CD patients except a single patient with missing data. Furthermore, HLA-DQ2/HLA-DQ8 was found positive in a patient with selective IgA deficiency.

In conclusion, our findings have demonstrated the increasing prevalence of CD in children with T1DM. Particularly the higher risk of asymptomatic CD in T1DM patients, revealed the diagnostic value of serological screening. Furthermore, increased HLA-DQ2 and HLA-DQ8 positivity, which are more prominent in case of selective IgA deficiency, clearly demonstrates the requirement of routine total IgA and HLA analysis in serological screening.

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