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**Celiac disease in children: A review of the literature**

Sahin Y. Celiac disease in children

Yasin Sahin

**Yasin Sahin,** Pediatric Gastroenterology-Hepatology and Nutrition, Medical Park Gaziantep Hospital, Gaziantep 27560, Turkey

**Author contributions:** Sahin Y wrote the paper and collected the data.

**Corresponding author: Yasin Sahin, MD, Associate Professor,** Pediatric Gastroenterology-Hepatology and Nutrition, Medical Park Gaziantep Hospital, Mucahitler Mah. 52063 sok. no:2 Sehitkamil, Gaziantep 27560, Turkey. ysahin977@gmail.com

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**Abstract**

Celiac disease is an immune-mediated systemic disease triggered by intake of gluten in genetically susceptible individuals. The prevalence of celiac disease in the general population is estimated to be 1% in the world. Its prevalence differs depending on geographical and ethnic variations. The prevalence of celiac disease has increased significantly in the last 30 years due to the increased knowledge and awareness of physicians and the widespread use of highly sensitive and specific diagnostic tests for celiac disease. Despite increased awareness and knowledge about celiac disease, up to 95% of celiac patients still remain undiagnosed. The presentations of celiac disease have significantly changed in the last few decades. Classical symptoms of celiac disease occur in a minority of celiac patients, while older children have either minimal or atypical symptoms. Serologic tests for celiac disease should be done in patients with unexplained chronic or intermittent diarrhea, failure to thrive, weight loss, delayed puberty, short stature, amenorrhea, iron deficiency anemia, nausea, vomiting, chronic abdominal pain, abdominal distension, chronic constipation, recurrent aphthous stomatitis, and abnormal liver enzyme elevation, and in children who belong to specific groups at risk. Early diagnosis of celiac disease is very important to prevent long-term complications. Currently, the only effective treatment is a lifelong gluten-free diet. In this review, we will discuss the epidemiology, clinical findings, diagnostic tests, and treatment of celiac disease in the light of the latest literature.

**Key Words:** Celiac disease; Children; Intestinal biopsy

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**Core Tip:** Celiac disease is a systemic lifelong disease. The prevalence of celiac disease has increased significantly in the last three decades due to the increased awareness of physicians and widespread use of highly sensitive and specific diagnostic tests for celiac disease. Despite increased awareness and widespread use of diagnostic tests, up to 95% of celiac patients still remain undiagnosed. Early diagnosis is very important to prevent long-term complications. The only effective treatment is still a lifelong gluten-free diet. In this review, we will discuss the epidemiology, clinical findings, diagnostic tests, and treatment of celiac disease in the light of the latest literature.

**INTRODUCTION**

Celiac disease is an immune-mediated systemic disease triggered by intake of gluten and related prolamines in genetically susceptible individuals, characterized by presence of various combinations of small intestinal damages, celiac specific antibodies, human leukocyte antigen (HLA)-DQ2 or HLA-DQ8, and gluten-dependent clinical manifestations[1]. Gluten is found in wheat, barley, rye, and oats[2].

**PATHOGENESIS**

The key elements of the celiac disease, an autoimmune disease, are genetics HLA-DQ2 and HLA-DQ8 genotypes, environmental factors (gluten intake), and autoantigen to tissue transglutaminase (tTG), which are known to play an important role in the pathogenesis[3]. In addition to genetic susceptibility and gluten exposure, loss of intestinal barrier function, gluten-induced proinflammatory innate immune response, inappropriate adaptive immune response, and unbalanced gut microbiome all seem to be components of the celiac disease autoimmunity[3]. More than 99% of celiac patients have HLA-DQ2 or HLA-DQ8 compared to 40% in the general population[4].

It has been suggested that breast milk, mode of delivery, and the age of gluten intake in infants are a risk for developing celiac disease and may affect the incidence of celiac disease. However, there is a limited information in retrospective studies that those factors affect the risk of developing celiac disease[5-7].

Furthermore, it has been suggested that gastrointestinal system (GIS) infections such as rotavirus may increase the risk of developing celiac disease, and therefore rotavirus vaccine may significantly reduce the risk of celiac disease especially in infants with gluten intake before 6 mo[8].

**EPIDEMIOLOGY**

The prevalence of celiac disease in the general population is estimated to be 1% in the world[9]. The seroprevalence of celiac disease and a biopsy-proven prevalence of celiac disease in the world is 1.4% and 0.7%, respectively[10]. Its prevalence varies depending on geographical and ethnic variations. The highest prevalence is in Europe (0.8%) and Ocenia (0.8%), and the lowest prevalence is in South America (0.4%). The biopsy-proven prevalence of celiac disease was found to be 1.5 times higher in women than men, and approximately two times higher in children than adults. The reason for this difference may be genetic factors [human leukocyte antigen (HLA) and non-HLA genes], environmental factors such as wheat consumption, age at gluten intake, gastrointestinal infections, proton pump inhibitor and antibiotic use, and the rate of cesarean section[10-12].

Celiac disease can occur at any age from early childhood to old age. It has two peaks; the first peak occurs after gluten intake within the first 2 years of life, the second is seen in the second or third decade of life. The diagnosis of celiac disease is difficult because symptoms vary from patient to patient[13].

The prevalence of celiac disease has increased significantly in the last 30 years, the reason for this is not only the increased knowledge and awareness of physicians about celiac disease but also due to the widespread use of highly sensitive and specific diagnostic tests for celiac disease[14,15]. For example, the incidence of pediatric celiac disease in Canada has increased 3-fold after the use of the endomysial antibody (EMA) test[16]. Despite increased awareness and knowledge about celiac disease, up to 95% of celiac patients still remain undiagnosed[17-19]. The delay in celiac disease diagnosis is reported to be 4-10 years in some studies[20,21]. There are many undiagnosed cases even in developed countries. Very few patients have clinically significant signs of celiac disease. The majority of cases have atypical signs or vague symptoms, so the diagnosis could not be made or diagnosis is delayed[22,23]. The reason for delayed or overlooked diagnosis may be the limited accessibility to serological diagnostic tests in developing countries and the lack of experienced specialists in this field[24].

The risk of developing celiac disease is higher in first- and second-degree relatives of celiac patients, Down syndrome, type 1 diabetes mellitus (DM), selective immunoglobulin (Ig)A deficiency, autoimmune thyroiditis, Turner syndrome, and Williams syndrome (Table 1)[25-28]. Screening tests for celiac disease at risk groups such as type 1 DM, autoimmune thyroid diseases, and first degree relatives of celiac patients also contributed to the increase in prevalence of celiac disease[27,29,30].

The prevalence of celiac disease in first degree relatives of celiac patients is as high as 10%-20%[1,31]. In a recent study of Sahin *et al*[32] the prevalence of celiac disease (CD) in siblings of pediatric celiac patients is reported to be 3.9%. The prevalence of CD in monozygotic twins has been found as high as 75%-80%[33,34].

In recent years, there has been a marked increase in the number of people having gluten-free diet. Furthermore, it has been observed that first-degree relatives of celiac patients start on a gluten-free diet before serologic tests for celiac disease were performed[35]. Therefore, before performing a serological test for celiac disease, it should be paid attention to whether they are on a gluten-free diet. Otherwise, the result of serological tests may be negative, and it would be difficult to diagnose celiac disease. Patients should take gluten-containing foods for 2-8 wk before serological tests[36].

**CLINICAL MANIFESTATIONS**

Symptoms usually occur in children after ingestion of gluten containing grains between 4 and 24 mo. There may be a delay or latent period between gluten intake and the onset of symptoms[37].

GIS and extra-intestinal manifestations are common in celiac disease[38]. The main GIS manifestations of celiac disease are chronic diarrhea, recurrent abdominal pain, nausea, vomiting, and abdominal distension. Common extra-intestinal manifestations are failure to thrive, short stature, chronic anemia, osteopenia, osteoporosis, delayed puberty, dental enamel defect, irritability, chronic fatigue, neuropathy, arthritis, arthralgia, amenorrhea, and increased liver enzymes[1,38].

Symptoms are usually different in infants than older children. Diarrhea, anorexia, abdominal distension, and abdominal pain are usually seen in younger children. If the diagnosis is delayed, failure to thrive, irritability, and severe malnutrition can be seen. GIS symptoms such as diarrhea, nausea, vomiting, abdominal pain, abdominal distension, weight loss, and constipation may occur in older children depending on the amount of gluten intake[28,37]. GIS signs of celiac disease such as diarrhea are seen in approximately 50% of patients[39-41].

The presentations of CD have significantly changed in the last few decades[41-48]. Classical symptoms of celiac disease occur in a minority of celiac patients, while older children have either minimal or atypical symptoms. GIS symptoms are mild or nonspecific[48,49].

It has been shown that pediatric patients diagnosed with celiac disease who are younger age at the diagnosis have less severe symptoms in the last 20 years. Also, it has been reported that the rate of asymptomatic patients, closer follow up, and strict adherence to gluten-free diet is higher in the last 10 years and that normalization of serological tests is faster than in the last decade[42].

Recently, the clinical symptoms of children with celiac disease are observed to change from GIS symptoms to extra-intestinal symptoms[39,50]. The exact reason for this is unclear, but it has been suggested that there may be increased awareness and widespread use of highly sensitive and specific serologic tests. It has been reported that isolated short stature is seen in up to 47.5% of celiac patients[41,51].

**EXTRA-INTESTINAL MANIFESTATIONS**

Extra-intestinal findings are seen in up to 60% of pediatric celiac patients (Table 2)[52]. Short stature is the most common finding in children[52-54]. It has been reported that 10%-47.5% of pediatric celiac patients have short stature at the time of diagnosis[41,54-57]. Nineteen percent to 59% of the non-endocrinologic causes of short stature are reported to be celiac disease[55,56,58-60]. Starting a gluten-free diet in the early period causes rapid growth and weight catch up, especially in the first 6 mo. The target height is usually reached within 3 years after diagnosis. If the target height is not reached despite a strict gluten-free diet, endocrinological evaluation should be done to rule out growth hormone deficiency[55,61-63].

Hypogonadism in girls and delayed puberty in boys due to androgen resistance is a common finding in undiagnosed or untreated pediatric celiac patients[55,64,65]. Delayed puberty is seen in 10%-20% of celiac patients[52,66]. Generally, the development of puberty occurs within 6-8 mo after starting a gluten-free diet. If delayed puberty persists, the patient should be referred to pediatric endocrinology for further evaluation of other disorders of the reproductive system[55,67].

Iron deficiency anemia is seen in up to 40% of pediatric celiac patients[52,53,68,69]. Since iron is absorbed from the first part of the duodenum, which is mainly affected by celiac disease, iron deficiency anemia is common in celiac patients. It has been reported that 84% of pediatric celiac patients have the complete recovery of iron deficiency anemia with a strict gluten-free diet and iron supplementation therapy within 12-24 mo[52].

Hypertransaminasemia is seen in 9%-14% of celiac patients[70]. Mostly, liver damage is reversible, and liver failure rarely occurs[71]. It has been suggested that as a result of exposure to more hepatotoxins through the portal circulation due to the altered intestinal permeability, inflammation and liver damage may occur[54,72]. The response to a strict gluten-free diet is excellent. The increased liver enzymes return to normal by the rate of 75%-90% within 12-24 mo with a strict gluten-free diet[73].

Osteopenia and osteoporosis are usually seen in patients with celiac disease. Approximately 75% of celiac patients have osteopenia and 10%-30% have osteoporosis[74]. Secondary hyperparathyroidism occurs due to the insufficient absorption of vitamin D and calcium from the damaged duodenal mucosa. It is commonly seen in 12%-54% of celiac patients[75]. Normal blood levels of vitamin D and calcium is observed within the first year after a strict gluten-free diet[76,77].

The most common joint and muscle disorders seen in celiac disease are myopathy, arthralgia, and non-erosive arthritis[55,78]. Since arthralgia is mostly seen after the age of 12, the most common finding in pediatric celiac patients is subclinical synovitis. It is most commonly seen in the knee joint. Its incidence is 5%-10%[54]. Since symptoms are mild, ultrasonography is important in the diagnosis of joint disorders.

The most common finding of neurological manifestations is headache, which is seen in up to 20% of celiac patients. More rarely, ataxia and neuropathy (0.1%-7.4%) are seen[79,80]. The prevalence of epilepsy is reported to be 1.43 times higher in children with celiac disease compared to the general population[81]. The relationship between epilepsy and CD is still unclear.

The exact prevalence of enamel defects in celiac disease is unknown. In recent studies, it has been reported that enamel defects are seen in 55%-64% of celiac patients[82,83].

Aphthous stomatitis is seen in up to 46% of celiac patients[84]. Although its mechanism is not known exactly, it is usually completely cured with a strict gluten-free diet[52].

Dermatitis herpetiformisis thought to be an extra-intestinal manifestation of celiac disease, but it is relatively rare in pediatric celiac patients in Finland[85]. Unlike celiac disease, its annual incidence is decreasing. The reason for this is unknown exactly[85]. In contrast to that study, it has been reported that it is more common in childhood[86].

**ASSOCIATED DISEASES WITH CELIAC DISEASE**

The risk of another autoimmune disease is three to 10 times higher in patients with celiac disease compared to the general population[87,88].

The most common accompanying disease is type 1 DM since it has common genetic factors and pathogenic mechanisms with celiac disease[89]. HLA-DQ2 is present in approximately 90%-95% of celiac patients and 50% of type 1 DM patients, but HLA-DQ8 is detected in approximately 10% of celiac patients and approximately 70% of type 1 DM patients[90]. In a systematic review, the prevalence of celiac disease in patients with type 1 DM was reported to be approximately six times higher than in the general population[91]. The prevalence of celiac disease was reported to be 2.4%-16.4% in children with type 1 DM[92-95]. There is consensus about initial screening for celiac disease in newly diagnosed DM patients, but it is not clear when and how often to screen for celiac disease and initiate a gluten-free diet in asymptomatic patients[93]. It has been recommended that screening test for CD should be done at the time of type 1 DM diagnosis and then every 2 years[96]. In another study, it was recommended that children diagnosed with type 1 DM should be screened for celiac disease once a year for the first 5 years[92]. In other studies, it has been recommended that serological screening tests for celiac disease should be done within the first 2 years when the diagnosis is made, then 5 years after the diagnosis and if there is any symptom suggestive of CD[93,97]. Since 58%-85% of type 1 DM patients diagnosed with CD are asymptomatic, early diagnosis of CD is very important to prevent long-term complications such as failure to thrive, osteopenia, infertility, and malignancy[29,77,92,93,98,99].

There is good evidence that autoimmune thyroid diseases are associated with celiac disease[1,100]. The prevalence of celiac disease in patients with autoimmune thyroid disease is found to be 3.0%-4.8%[30,101,102].

Also, the prevalence of celiac disease in patients with selective IgA deficiency is reported to be 10-20 times higher than in the general population[103].

There is a close relationship between Down syndrome and celiac disease. The prevalence of celiac disease in patients diagnosed with Down syndrome is reported to be 5%-12%[104-108]. The North American Society for Paediatric Gastroenterology, Hepatology and Nutrition and The European Society for Paediatric Gastroenterology, Hepatology and Nutrition (ESPGHAN) recommend screening tests for celiac disease in children with Down syndrome due to the increased risk of developing celiac disease[28]. In a study conducted in 2020, involving 1317 pediatric patients with Down syndrome aged 3 and over, the prevalence of celiac disease was found to be 9.8% in children with Down syndrome[109]. If screening test for celiac disease is not done, the diagnosis of celiac disease is either overlooked or delayed in 82% of the patients with Down syndrome, thus causing increased morbidity[109].

The increased prevalence of celiac disease is also seen in autoimmune liver disease, Turner syndrome, and Williams syndrome[1,110-116].

**THE DEFINITONS RELATED TO CELIAC DISEASE**

***Silent celiac disease***

Silent celiac disease is defined by the presence of celiac antibodies and HLA-DQ2 or HLA-DQ8 and small intestinal biopsy findings compatible with celiac disease especially in patients with autoimmune disease or a genetic disorder or relatives of celiac disease but without any symptoms suggestive of CD[1].

***Potential celiac disease***

Potential celiac disease is defined by the presence of celiac antibodies, HLA-DQ2 or HLA-DQ8, but intestinal biopsy is not compatible with celiac disease. Marsh classification score 0 or 1 is detected in intestinal biopsy, and the risk of developing celiac disease is increased[117].

Clinical symptoms and signs of the celiac disease are not always seen. Even if there are clinical findings, they are usually mild. The diagnosis of potential CD has increased significantly in recent years due to increased use of serological screening for celiac disease in the general population. A lower prevalence of HLA-DQ2 and a higher prevalence of HLA-DQ8 are detected in potential celiac patients compared to active celiac patients[118].

It should be considered that the cause of negative intestinal biopsy may be the patchy involvement of the small intestinal mucosa, low gluten intake, and inappropriate biopsy orientation[119].

Its treatment is still uncertain and controversial. There is no consensus about how often celiac serological tests should be performed in potential celiac patients on a gluten-containing diet, and how often they should be evaluated clinically[120]. It has been reported that villous atrophy is observed in 33% of symptomatic potential celiac patients after 3 years[121]. Therefore, it has been suggested that symptomatic patients should be given a gluten free diet.

***Refractory celiac disease***

Refractory celiac disease is characterized by the persistence of symptoms and intestinal villous atrophy despite a strict gluten-free diet for at least 12 mo. Generally, celiac antibodies are negative in most patients at the time of diagnosis, but the presence of high-titer antibodies does not rule out the refractory celiac disease. In all cases, dietary adherence should be carefully questioned. It can cause complications such as ulcerative jejunoileitis, collagenous sprue, and intestinal lymphoma[117].

***Seronegative celiac disease***

It is characterized by the presence of clinical signs of severe malabsorption and intestinal villous atrophy and negative celiac antibodies[122]. It constitutes approximately 2%-3% of celiac patients. Seronegative celiac disease can be confirmed with improvement in both symptoms and histology 1 year after starting a gluten-free diet[122]. Compared with classical celiac disease, seronegative celiac patients are associated with a higher rate of autoimmune disease, and these patients have a higher risk of developing refractory celiac disease[122].

In this form of celiac disease, genetic analysis is the key step for the diagnosis, because if it is found as negative, celiac disease is ruled out. Other diseases causing villous atrophy are parasitic infections (*e.g.*, *Giardia lamblia*), autoimmune enteropathy, small intestinal bacterial overgrowth, common variable immunodeficiency, eosinophilic gastroenteritis, drug induced enteropathy (*e.g.*, olmesartan, mycophenolate), intestinal lymphoma, Crohn's disease, tropical sprue, human immunodeficiency virus enteropathy, and Whipple disease should be considered in the differential diagnosis (Table 3)[122-124].

***Non-responsive celiac disease***

Non-responsive celiac disease is defined by the persistence of GI symptoms more than 12 mo despite a strict gluten-free diet. The most common causes of non-responsive celiac disease are persistent gluten ingestion and incorrect diagnosis[125,126]. It needs to be differentiated from active celiac disease and other conditions associated with celiac disease.

**DIAGNOSIS**

The clinical symptoms of celiac disease are very diverse. Celiac patients may present with symptoms of GIS or extra-intestinal symptoms or no symptoms at all. Therefore, serologic tests for celiac disease should be done in patients with unexplained chronic or intermittent diarrhea, failure to thrive, weight loss, delayed puberty, short stature, amenorrhea, iron deficiency anemia, nausea, vomiting, chronic abdominal pain, abdominal distension, chronic constipation, recurrent aphthous stomatitis, and abnormal liver enzyme elevation[1].

Furthermore, celiac disease should be investigated in patients with high risk of developing celiac disease, such as type 1 DM, Down syndrome, autoimmune thyroid disease, Turner syndrome, selective IgA deficiency, autoimmune liver disease, and first-degree relatives of celiac patients, even if they are asymptomatic[1].

Celiac disease is diagnosed by a variable combination of symptoms, positive celiac antibodies, presence of HLA-DQ2/DQ8, and duodenal histology[1].

ESPGHAN guidelines from 2012 recommend tissue tTG-IgA test, which is highly sensitive and specific and less costly compared to EMA IgA antibody test, as an initial screening test for suspected celiac disease, and the total IgA test to rule out selective IgA deficiency. The analysis of deamidated gliadin peptide (DGP) IgA test is recommended for children under 2 years of age. If there is IgA deficiency, the tTG-IgG test or the EMA-IgG test or the DGP-IgG test should be performed[1].

If serological tests are negative for tTG-IgA and total IgA level is normal, celiac disease is unlikely. In this condition, the reasons leading to the false negative tTG result should be considered. Those are low gluten intake, protein-losing enteropathy, use of immunosuppressive drugs, and patients under 2 years of age. If the tTG is found as positive [lower than 10 times upper limit of normal (ULN)], gastroduodenoscopy and multiple biopsies of the small intestine should be performed to confirm the diagnosis[1].

If the tTG is higher than 10 times ULN in a symptomatic patient, it should be discussed with the parents in order to make a diagnosis of celiac disease without biopsy. If the parents agree, EMA test and HLA-DQ2/DQ8 analysis are performed. To rule out false positivity of the tTG test, an EMA test is performed from a second blood sample. If EMA and HLA-DQ2 or HLA-DQ8 are positive, celiac disease is diagnosed without biopsy[1]. In practice, it has been reported that this reduces the need for endoscopy by 30%-50%[127].

Since celiac disease causes patchy involvement in the small intestine, at least four biopsies from the duodenum and at least one biopsy from the bulbus should be performed by gastroduodenoscopy. Biopsies are evaluated according to modified Marsh-Oberhuber classification (Table 4)[128]. Since the lesion of celiac disease can only be seen in the bulb, at least one biopsy should be taken from the bulb[129].

While interpreting the serological test results of celiac disease, serum total IgA levels, the amount of gluten consumption, use of immunosuppressive drugs, and age of the patient should be considered[1]. IgG class celiac antibody tests should be performed in patients with low serum IgA levels (total serum IgA < 0.2 g/L)[1].

If the patient has the gluten-free diet for a long time or gluten-free diet for a short time before testing, false negative results may occur[130]. Therefore, patients should take definitely gluten-containing foods before the test. Gluten challenge test should be performed for patients with a gluten-free diet before serological tests, 3-7.5 g/d gluten-containing diet (approximately two slices of bread) is recommended for 2 wk[131].

If the patient is strongly suspected of celiac disease, multiple intestinal biopsy and HLA-DQ2/DQ8 analysis are recommended, even if the serological tests for celiac disease are negative. If the histology is compatible with celiac disease but HLA-DQ2/8 negative, celiac disease is unlikely and other causes of enteropathy should be investigated (Table 3)[1]. Celiac disease is diagnosed if the celiac serological tests are positive and the biopsy is compatible with celiac disease.

ESPGHAN guidelines from 2020 report that the tTG-IgA test and total IgA test combination give more accurate results than other test combinations as the initial test for suspected celiac disease regardless of age. If total IgA level is found to be low, tTG-IgG test or EMA-IgG test or DGP-IgG test should be performed (Figure 1)[119].

If the tTG test is found as positive (> 10 times ULN), HLA-DQ2/8 analysis is not recommended in the ESPGHAN 2020 guidelines even if the patient is asymptomatic. It has been suggested that the EMA test should be checked in a second blood sample and if the EMA test is detected positive and the family agrees, celiac disease can be diagnosed without biopsy. In other words, the presence of HLA-DQ2/8 analysis and clinical symptoms are not mandatory for celiac diagnosis in last guideline in 2020 (Figure 1)[119].

If HLA-DQ2/DQ8 test is negative, the probability of celiac disease is low, but a positive HLA-DQ2/DQ8 test does not confirm the diagnosis of celiac disease[132]. If the tTG test is detected positive (< 10 times ULN), multiple intestinal biopsy is recommended to rule out false positivity. It is not recommended to diagnose without biopsy in patients with selective IgA deficiency even if IgG-based antibody positivity is detected[119].

It has been considered that villous atrophy may be seen in other GIS diseases such as parasitic infections, autoimmune diseases, bacterial overgrowth in the small intestine, and Crohn's disease (Table 3)[133].

It has been reported that the pooled sensitivity and specificity of tTG or DGP or tTG + antigliadin antibodies for diagnosing celiac disease is 94.0% and 94.4%, respectively, in a systematic review[134]. It has been suggested that those tests can be used in places where access to laboratory tests is limited.

**MANAGEMENT**

Currently, the only effective treatment is a lifelong gluten-free diet. Significant improvements in symptoms, normalization of biochemical tests, and improvement in quality of life with a strict gluten-free diet are seen[135].

Rapid improvement in clinical symptoms is observed within 2-4 wk in children. Serological and histological responses are slower compared to clinical symptoms[136]. Although histological response in children is observed within 2 years by a rate of 95%, this rate is 60% in adults[137].

The amount of tolerable gluten varies from patient to patient. As little as 50 mg of gluten, present in a few amounts of bread crumbs or a small piece of cake or traces of contamination, may cause symptoms and/or enteropathy in asymptomatic patients[135,138]. It is unlikely that a gluten intake of less than 10 mg/d will cause significant histological abnormality[139].

Adherence to the gluten-free diet is better in children diagnosed with CD at an early age and those who continue to follow up regularly. It is less in adolescents compared to adults[135].

It has been reported that there is a direct relationship between the duration of exposure to the gluten-free diet and increased autoimmune disorders[140].

In a multicenter prospective study involving 6605 children with the HLA genotype associated with celiac disease, it was shown that the amount of gluten exposure in the first 5 years of life is associated with the development of celiac disease and celiac autoimmunity[141]. Since celiac disease is a multisytemic disease that affects multiple organs, a lifelong gluten-free diet may reduce malignant and non-malignant complications[142].

**FOLLOW-UP**

Currently, there are no standard evidence-based recommendations for the follow-up of pediatric celiac disease[143].

Patients with celiac disease should be followed up 6 mo after diagnosis and every 6 mo in terms of improvement in symptoms, compliance with the gluten-free diet, quality of life, and progressive normalization of celiac-associated antibodies. Screening tests should be done in terms of autoimmune thyroid disease. A control duodenal biopsy is not required after a gluten-free diet. However, if there is a partial or no response to the gluten-free diet, careful examination should be done for involuntary gluten contamination or poor compliance with the gluten-free diet. If the response to a strict gluten-free diet is poor, duodenal biopsy can be performed[135,143,144].

Earlier diagnosis of celiac disease in asymptomatic patients is associated with better quality of life as well as better compliance with the gluten-free diet[42,145,146].

It has been shown that pediatric patients who are lost to follow up are less adherent to the gluten-free diet and have positive celiac serological antibodies[147]. It has been shown that the regular control is very important.

Routine testing for vitamin and mineral deficiency is reported to be unnecessary in the vast majority of children who follow up to regular controls and have normal growth and development and have no symptoms[148].

The essential marker of the success of the gluten-free diet is still satisfactory height and weight gain in children and adolescents[135].

The best marker of proper follow-up and management is the decline in the antibody levels and the return of antibody levels to normal in follow-up. The presence of persistent positive antibodies usually indicates ongoing intestinal damage and gluten exposure. Serological follow-up should be done within 6 mo and 12 mo after diagnosis and then once a year[149].

tTG-IgA test is reported to be best test in follow up[150]. It has been shown that the average time to return to normal levels of the tTG test in patients with strictly adherent to the gluten-free diet is 1 year[151].

It has been detected that there is no correlation between symptoms and mucosal healing[152]. Gluten challenge test can be performed in cases when there is a doubt about the initial diagnosis of celiac disease. However, HLA typing should be done before evaluation of mucosal damage. Gluten challenge is not recommended under 5 years of age and during pubertal development[1].

In recent studies, it has been reported that gluten consumption can be shown in symptomatic and asymptomatic patients who are unaware of gluten intake by gluten immunogenic peptide tests in stool and urine[153,154]. Gluten intake of more than 50 mg/d for stool test and more than 25 mg/d for urine test seems to be necessary for the sensitivity of the test[153]. Dietary adherence to the gluten-free diet can be evaluated with this test. It can replace serological tests in follow-up, but its use in routine practice is still uncertain and further studies are needed.

**DIETS AND NEW TREATMENTS**

Currently, the only effective treatment is still to avoid gluten completely for life. The adherence to the gluten-free diet has some disadvantages; negative impact on quality of life, psychological problems, involuntary gluten contamination, possible vitamin and mineral deficiencies, metabolic syndrome, increased cardiovascular risk, and severe constipation[153,155-157].

Approximately 40% of celiac patients are not satisfied with the gluten-free diet due to the negative effect on their quality of life and seek alternative treatments[158,159].

Clinical studies are still ongoing in the treatment of celiac disease. Larazotide acetate is a zonulin antagonist that blocks the tight junction, thus restricting the passage of gluten through the permeable intestinal mucosa[160]. This drug is shown to be effective in controlling gluten-related symptoms[160]. There is also limited information that larazotide may allow patients to tolerate minimal amounts of gluten (involuntary gluten contamination or short-term feeding with a small amount of gluten).

ALV003 (latiglutenase) reduces gluten into small pieces in the stomach before it passes into the duodenum[161]. In a study involving 494 celiac patients, latiglutenase was compared with placebo. It has been shown that latiglutenase did not improve histological findings or symptoms[162]. Further studies are needed.

Vaccination (Nexvax2) is another therapeutic option intended to be used for desensitization in celiac patients against gliadin peptides. Although its major side effects are abdominal pain and vomiting, it passed phase 1. Given the effectiveness of vaccines, it can be a definitive cure for celiac disease[163].

**COMPLICATIONS**

Complication are usually manifested in late-diagnosed celiac patients (after the age of 50) and in patients not adhering to a strict gluten-free diet. These patients have a higher mortality than the general population[164], but complications are rare (< 1%)[165].

Complications of celiac disease include hyposplenism, refractory celiac disease, intestinal lymphoma, small bowel adenocarcinoma, and ulcerative jejunoileitis[166].

Despite adhering to a gluten-free diet and having complaints that cannot be explained by any other reason, complications should be considered in every patient whose symptoms persist.

**CONCLUSION**

Celiac disease is a lifelong multi-systemic disease triggered by intake of gluten in genetically susceptible individuals.

Serologic tests for CD should be done in patients with unexplained chronic or intermittent diarrhea, failure to thrive, weight loss, delayed puberty, short stature, amenorrhea, iron deficiency anemia, nausea, vomiting, chronic abdominal pain, abdominal distension, chronic constipation, recurrent aphthous stomatitis, and abnormal liver enzyme elevation.

Since tTG-IgA test and total IgA test combination give more accurate results than other test combinations, ESPGHAN 2020 guideline recommends this combination as the initial test for suspected celiac disease regardless of age. While interpreting the serological test results of celiac disease, serum total IgA levels, the amount of gluten consumption, use of immunosuppressive drugs, and age of the patient should be considered.

Early diagnosis of CD is very important to prevent long-term complications such as failure to thrive, osteopenia, infertility, and malignancy.

Currently, the only effective treatment is a lifelong gluten-free diet.

**REFERENCES**

1 **Husby S**, Koletzko S, Korponay-Szabó IR, Mearin ML, Phillips A, Shamir R, Troncone R, Giersiepen K, Branski D, Catassi C, Lelgeman M, Mäki M, Ribes-Koninckx C, Ventura A, Zimmer KP; ESPGHAN Working Group on Coeliac Disease Diagnosis; ESPGHAN Gastroenterology Committee; European Society for Pediatric Gastroenterology, Hepatology, and Nutrition. European Society for Pediatric Gastroenterology, Hepatology, and Nutrition guidelines for the diagnosis of coeliac disease. *J Pediatr Gastroenterol Nutr* 2012; **54**: 136-160 [PMID: 22197856 DOI: 10.1097/MPG.0b013e31821a23d0]

2 **Fasano A**, Catassi C. Clinical practice. Celiac disease. *N Engl J Med* 2012; **367**: 2419-2426 [PMID: 23252527 DOI: 10.1056/NEJMcp1113994]

3 **Caio G**, Volta U, Sapone A, Leffler DA, De Giorgio R, Catassi C, Fasano A. Celiac disease: a comprehensive current review. *BMC Med* 2019; **17**: 142 [PMID: 31331324 DOI: 10.1186/s12916-019-1380-z]

4 **Hadithi M**, von Blomberg BM, Crusius JB, Bloemena E, Kostense PJ, Meijer JW, Mulder CJ, Stehouwer CD, Peña AS. Accuracy of serologic tests and HLA-DQ typing for diagnosing celiac disease. *Ann Intern Med* 2007; **147**: 294-302 [PMID: 17785484 DOI: 10.7326/0003-4819-147-5-200709040-00003]

5 **Lionetti E**, Castellaneta S, Francavilla R, Pulvirenti A, Catassi C; SIGENP Working Group of Weaning and CD Risk. Mode of Delivery and Risk of Celiac Disease: Risk of Celiac Disease and Age at Gluten Introduction Cohort Study. *J Pediatr* 2017; **184**: 81-86.e2 [PMID: 28196682 DOI: 10.1016/j.jpeds.2017.01.023]

6 **Koletzko S**, Lee HS, Beyerlein A, Aronsson CA, Hummel M, Liu E, Simell V, Kurppa K, Lernmark Å, Hagopian W, Rewers M, She JX, Simell O, Toppari J, Ziegler AG, Krischer J, Agardh D; TEDDY Study Group. Cesarean Section on the Risk of Celiac Disease in the Offspring: The Teddy Study. *J Pediatr Gastroenterol Nutr* 2018; **66**: 417-424 [PMID: 28753178 DOI: 10.1097/MPG.0000000000001682]

7 **Dydensborg Sander S**, Hansen AV, Størdal K, Andersen AN, Murray JA, Husby S. Mode of delivery is not associated with celiac disease. *Clin Epidemiol* 2018; **10**: 323-332 [PMID: 29593435 DOI: 10.2147/CLEP.S152168]

8 **Silvester JA**, Leffler DA. Is Autoimmunity Infectious? The Effect of Gastrointestinal Viral Infections and Vaccination on Risk of Celiac Disease Autoimmunity. *Clin Gastroenterol Hepatol* 2017; **15**: 703-705 [PMID: 28017844 DOI: 10.1016/j.cgh.2016.12.014]

9 **Lindfors K**, Ciacci C, Kurppa K, Lundin KEA, Makharia GK, Mearin ML, Murray JA, Verdu EF, Kaukinen K. Coeliac disease. *Nat Rev Dis Primers* 2019; **5**: 3 [PMID: 30631077 DOI: 10.1038/s41572-018-0054-z]

10 **Singh P**, Arora A, Strand TA, Leffler DA, Catassi C, Green PH, Kelly CP, Ahuja V, Makharia GK. Global Prevalence of Celiac Disease: Systematic Review and Meta-analysis. *Clin Gastroenterol Hepatol* 2018; **16**: 823-836.e2 [PMID: 29551598 DOI: 10.1016/j.cgh.2017.06.037]

11 **Lebwohl B**, Murray JA, Verdú EF, Crowe SE, Dennis M, Fasano A, Green PH, Guandalini S, Khosla C. Gluten Introduction, Breastfeeding, and Celiac Disease: Back to the Drawing Board. *Am J Gastroenterol* 2016; **111**: 12-14 [PMID: 26259710 DOI: 10.1038/ajg.2015.219]

12 **Choung RS**, Ditah IC, Nadeau AM, Rubio-Tapia A, Marietta EV, Brantner TL, Camilleri MJ, Rajkumar SV, Landgren O, Everhart JE, Murray JA. Trends and racial/ethnic disparities in gluten-sensitive problems in the United States: findings from the National Health and Nutrition Examination Surveys from 1988 to 2012. *Am J Gastroenterol* 2015; **110**: 455-461 [PMID: 25665935 DOI: 10.1038/ajg.2015.8]

13 **Fasano A**. Celiac disease--how to handle a clinical chameleon. *N Engl J Med* 2003; **348**: 2568-2570 [PMID: 12815143 DOI: 10.1056/NEJMe030050]

14 **Liu E**, Dong F, Barón AE, Taki I, Norris JM, Frohnert BI, Hoffenberg EJ, Rewers M. High Incidence of Celiac Disease in a Long-term Study of Adolescents With Susceptibility Genotypes. *Gastroenterology* 2017; **152**: 1329-1336.e1 [PMID: 28188747 DOI: 10.1053/j.gastro.2017.02.002]

15 **King JA**, Jeong J, Underwood FE, Quan J, Panaccione N, Windsor JW, Coward S, deBruyn J, Ronksley PE, Shaheen AA, Quan H, Godley J, Veldhuyzen van Zanten S, Lebwohl B, Ng SC, Ludvigsson JF, Kaplan GG. Incidence of Celiac Disease Is Increasing Over Time: A Systematic Review and Meta-analysis. *Am J Gastroenterol* 2020; **115**: 507-525 [PMID: 32022718 DOI: 10.14309/ajg.0000000000000523]

16 **McGowan KE**, Castiglione DA, Butzner JD. The changing face of childhood celiac disease in north america: impact of serological testing. *Pediatrics* 2009; **124**: 1572-1578 [PMID: 19948628 DOI: 10.1542/peds.2008-2373]

17 **Hujoel IA**, Van Dyke CT, Brantner T, Larson J, King KS, Sharma A, Murray JA, Rubio-Tapia A. Natural history and clinical detection of undiagnosed coeliac disease in a North American community. *Aliment Pharmacol Ther* 2018; **47**: 1358-1366 [PMID: 29577349 DOI: 10.1111/apt.14625]

18 **Murray JA**, Van Dyke C, Plevak MF, Dierkhising RA, Zinsmeister AR, Melton LJ 3rd. Trends in the identification and clinical features of celiac disease in a North American community, 1950-2001. *Clin Gastroenterol Hepatol* 2003; **1**: 19-27 [PMID: 15017513 DOI: 10.1053/jcgh.2003.50004]

19 **Lebwohl B**, Rubio-Tapia A, Assiri A, Newland C, Guandalini S. Diagnosis of celiac disease. *Gastrointest Endosc Clin N Am* 2012; **22**: 661-677 [PMID: 23083985 DOI: 10.1016/j.giec.2012.07.004]

20 **Sanders DS**, Hurlstone DP, Stokes RO, Rashid F, Milford-Ward A, Hadjivassiliou M, Lobo AJ. Changing face of adult coeliac disease: experience of a single university hospital in South Yorkshire. *Postgrad Med J* 2002; **78**: 31-33 [PMID: 11796869 DOI: 10.1136/pmj.78.915.31]

21 **Lo W**, Sano K, Lebwohl B, Diamond B, Green PH. Changing presentation of adult celiac disease. *Dig Dis Sci* 2003; **48**: 395-398 [PMID: 12643621 DOI: 10.1023/a:1021956200382]

22 **Lebwohl B**, Sanders DS, Green PHR. Coeliac disease. *Lancet* 2018; **391**: 70-81 [PMID: 28760445 DOI: 10.1016/S0140-6736(17)31796-8]

23 **Singh P**, Wadhwa N, Chaturvedi MK, Bhatia V, Saini S, Tandon N, Makharia GK, Maki M, Not T, Phillips A, Bhatnagar S. Validation of point-of-care testing for coeliac disease in children in a tertiary hospital in north India. *Arch Dis Child* 2014; **99**: 1004-1008 [PMID: 24942708 DOI: 10.1136/archdischild-2013-305567]

24 **Nenna R**, Tiberti C, Petrarca L, Lucantoni F, Mennini M, Luparia RP, Panimolle F, Mastrogiorgio G, Pietropaoli N, Magliocca FM, Bonamico M. The celiac iceberg: characterization of the disease in primary schoolchildren. *J Pediatr Gastroenterol Nutr* 2013; **56**: 416-421 [PMID: 23149808 DOI: 10.1097/MPG.0b013e31827b7f64]

25 **Nellikkal SS**, Hafed Y, Larson JJ, Murray JA, Absah I. High Prevalence of Celiac Disease Among Screened First-Degree Relatives. *Mayo Clin Proc* 2019; **94**: 1807-1813 [PMID: 31447136 DOI: 10.1016/j.mayocp.2019.03.027]

26 **Fasano A**, Berti I, Gerarduzzi T, Not T, Colletti RB, Drago S, Elitsur Y, Green PH, Guandalini S, Hill ID, Pietzak M, Ventura A, Thorpe M, Kryszak D, Fornaroli F, Wasserman SS, Murray JA, Horvath K. Prevalence of celiac disease in at-risk and not-at-risk groups in the United States: a large multicenter study. *Arch Intern Med* 2003; **163**: 286-292 [PMID: 12578508 DOI: 10.1001/archinte.163.3.286]

27 **Singh P**, Arora S, Lal S, Strand TA, Makharia GK. Risk of Celiac Disease in the First- and Second-Degree Relatives of Patients With Celiac Disease: A Systematic Review and Meta-Analysis. *Am J Gastroenterol* 2015; **110**: 1539-1548 [PMID: 26416192 DOI: 10.1038/ajg.2015.296]

28 **Hill ID**, Dirks MH, Liptak GS, Colletti RB, Fasano A, Guandalini S, Hoffenberg EJ, Horvath K, Murray JA, Pivor M, Seidman EG; North American Society for Pediatric Gastroenterology, Hepatology and Nutrition. Guideline for the diagnosis and treatment of celiac disease in children: recommendations of the North American Society for Pediatric Gastroenterology, Hepatology and Nutrition. *J Pediatr Gastroenterol Nutr* 2005; **40**: 1-19 [PMID: 15625418 DOI: 10.1097/00005176-200501000-00001]

29 **Pham-Short A**, Donaghue KC, Ambler G, Phelan H, Twigg S, Craig ME. Screening for Celiac Disease in Type 1 Diabetes: A Systematic Review. *Pediatrics* 2015; **136**: e170-e176 [PMID: 26077482 DOI: 10.1542/peds.2014-2883]

30 **Sahin Y**, Evliyaoglu O, Erkan T, Cokugras FC, Ercan O, Kutlu T. The frequency of celiac disease in children with autoimmune thyroiditis. *Acta Gastroenterol Belg* 2018; **81**: 5-8 [PMID: 29562371]

31 **Rubio-Tapia A**, Van Dyke CT, Lahr BD, Zinsmeister AR, El-Youssef M, Moore SB, Bowman M, Burgart LJ, Melton LJ 3rd, Murray JA. Predictors of family risk for celiac disease: a population-based study. *Clin Gastroenterol Hepatol* 2008; **6**: 983-987 [PMID: 18585974 DOI: 10.1016/j.cgh.2008.04.008]

32 **Sahin Y**. The Frequency of Celiac Disease in Siblings of Celiac Patients. *EC Paediatrics* 2019; **2**: 154-157

33 **Greco L**, Romino R, Coto I, Di Cosmo N, Percopo S, Maglio M, Paparo F, Gasperi V, Limongelli MG, Cotichini R, D'Agate C, Tinto N, Sacchetti L, Tosi R, Stazi MA. The first large population based twin study of coeliac disease. *Gut* 2002; **50**: 624-628 [PMID: 11950806 DOI: 10.1136/gut.50.5.624]

34 **Lundin KE**, Wijmenga C. Coeliac disease and autoimmune disease-genetic overlap and screening. *Nat Rev Gastroenterol Hepatol* 2015; **12**: 507-515 [PMID: 26303674 DOI: 10.1038/nrgastro.2015.136]

35 **Kim HS**, Patel KG, Orosz E, Kothari N, Demyen MF, Pyrsopoulos N, Ahlawat SK. Time Trends in the Prevalence of Celiac Disease and Gluten-Free Diet in the US Population: Results From the National Health and Nutrition Examination Surveys 2009-2014. *JAMA Intern Med* 2016; **176**: 1716-1717 [PMID: 27598396 DOI: 10.1001/jamainternmed.2016.5254]

36 **Rubio-Tapia A**, Hill ID, Kelly CP, Calderwood AH, Murray JA; American College of Gastroenterology. ACG clinical guidelines: diagnosis and management of celiac disease. *Am J Gastroenterol* 2013; **108**: 656-76; quiz 677 [PMID: 23609613 DOI: 10.1038/ajg.2013.79]

37 **Gallegos C**, Merkel R. Current Evidence in the Diagnosis and Treatment of Children With Celiac Disease. *Gastroenterol Nurs* 2019; **42**: 41-48 [PMID: 30688706 DOI: 10.1097/SGA.0000000000000365]

38 **Van Kalleveen MW**, de Meij T, Plötz FB. Clinical spectrum of paediatric coeliac disease: a 10-year single-centre experience. *Eur J Pediatr* 2018; **177**: 593-602 [PMID: 29392394 DOI: 10.1007/s00431-018-3103-4]

39 **Garampazzi A**, Rapa A, Mura S, Capelli A, Valori A, Boldorini R, Oderda G. Clinical pattern of celiac disease is still changing. *J Pediatr Gastroenterol Nutr* 2007; **45**: 611-614 [PMID: 18030243 DOI: 10.1097/MPG.0b013e31814c3d79]

40 **Simmons JH**, Klingensmith GJ, McFann K, Rewers M, Taylor J, Emery LM, Taki I, Vanyi S, Liu E, Hoffenberg EJ. Impact of celiac autoimmunity on children with type 1 diabetes. *J Pediatr* 2007; **150**: 461-466 [PMID: 17452216 DOI: 10.1016/j.jpeds.2006.12.046]

41 **Sahin Y**. Clinical evaluation of children with celiac disease: a single-center experience. *Arch Clin Gastroenterol* 2020; **6**: 26-30

42 **Krauthammer A**, Guz-Mark A, Zevit N, Marderfeld L, Waisbourd-Zinman O, Silbermintz A, Mozer-Glassberg Y, Nachmias Friedler V, Rozenfeld Bar Lev M, Matar M, Assa A, Shamir R. Two decades of pediatric celiac disease in a tertiary referral center: What has changed? *Dig Liver Dis* 2020; **52**: 457-461 [PMID: 32111387 DOI: 10.1016/j.dld.2020.02.001]

43 **Fasano A**. Clinical presentation of celiac disease in the pediatric population. *Gastroenterology* 2005; **128**: S68-S73 [PMID: 15825129 DOI: 10.1053/j.gastro.2005.02.015]

44 **Beattie RM**. The changing face of coeliac disease. *Arch Dis Child* 2006; **91**: 955-956 [PMID: 17119070 DOI: 10.1136/adc.2006.099671]

45 **Ravikumara M**, Tuthill DP, Jenkins HR. The changing clinical presentation of coeliac disease. *Arch Dis Child* 2006; **91**: 969-971 [PMID: 16887861 DOI: 10.1136/adc.2006.094045]

46 **Lebenthal E,** Shteyer E, Branski D. The changing clinical presentation of celiac disease. In: Fasano A, Troncone R, Branski D. Frontiers in celiac disease. Pediatr Adolesc Med Basel: Karger, 2008: 18-22 [DOI: 10.1159/000128609]

47 **Khatib M**, Baker RD, Ly EK, Kozielski R, Baker SS. Presenting Pattern of Pediatric Celiac Disease. *J Pediatr Gastroenterol Nutr* 2016; **62**: 60-63 [PMID: 26111294 DOI: 10.1097/MPG.0000000000000887]

48 **Kivelä L**, Kaukinen K, Lähdeaho ML, Huhtala H, Ashorn M, Ruuska T, Hiltunen P, Visakorpi J, Mäki M, Kurppa K. Presentation of Celiac Disease in Finnish Children Is No Longer Changing: A 50-Year Perspective. *J Pediatr* 2015; **167**: 1109-15.e1 [PMID: 26316370 DOI: 10.1016/j.jpeds.2015.07.057]

49 **Almallouhi E**, King KS, Patel B, Wi C, Juhn YJ, Murray JA, Absah I. Increasing Incidence and Altered Presentation in a Population-based Study of Pediatric Celiac Disease in North America. *J Pediatr Gastroenterol Nutr* 2017; **65**: 432-437 [PMID: 28151767 DOI: 10.1097/MPG.0000000000001532]

50 **Bottaro G**, Cataldo F, Rotolo N, Spina M, Corazza GR. The clinical pattern of subclinical/silent celiac disease: an analysis on 1026 consecutive cases. *Am J Gastroenterol* 1999; **94**: 691-696 [PMID: 10086653 DOI: 10.1111/j.1572-0241.1999.00938.x]

51 **van Rijn JC**, Grote FK, Oostdijk W, Wit JM. Short stature and the probability of coeliac disease, in the absence of gastrointestinal symptoms. *Arch Dis Child* 2004; **89**: 882-883 [PMID: 15321874 DOI: 10.1136/adc.2004.057851]

52 **Jericho H**, Sansotta N, Guandalini S. Extraintestinal Manifestations of Celiac Disease: Effectiveness of the Gluten-Free Diet. *J Pediatr Gastroenterol Nutr* 2017; **65**: 75-79 [PMID: 28644353 DOI: 10.1097/MPG.0000000000001420]

53 **Nurminen S**, Kivelä L, Huhtala H, Kaukinen K, Kurppa K. Extraintestinal manifestations were common in children with coeliac disease and were more prevalent in patients with more severe clinical and histological presentation. *Acta Paediatr* 2019; **108**: 681-687 [PMID: 29569302 DOI: 10.1111/apa.14324]

54 **Jericho H**, Guandalini S. Extra-Intestinal Manifestation of Celiac Disease in Children. *Nutrients* 2018; **10**: 755 [PMID: 29895731 DOI: 10.3390/nu10060755]

55 **Nardecchia S**, Auricchio R, Discepolo V, Troncone R. Extra-Intestinal Manifestations of Coeliac Disease in Children: Clinical Features and Mechanisms. *Front Pediatr* 2019; **7**: 56 [PMID: 30891436 DOI: 10.3389/fped.2019.00056]

56 **Bonamico M**, Sciré G, Mariani P, Pasquino AM, Triglione P, Scaccia S, Ballati G, Boscherini B. Short stature as the primary manifestation of monosymptomatic celiac disease. *J Pediatr Gastroenterol Nutr* 1992; **14**: 12-16 [PMID: 1573504 DOI: 10.1097/00005176-199201000-00003]

57 **Gokce S**, Arslantas E. Changing face and clinical features of celiac disease in children. *Pediatr Int* 2015; **57**: 107-112 [PMID: 25040342 DOI: 10.1111/ped.12448]

58 **Hyer W**, Cotterill AM, Savage MO. Common causes of short stature detectable by a height surveillance programme. *J Med Screen* 1995; **2**: 150-153 [PMID: 8536185 DOI: 10.1177/096914139500200310]

59 **Saari A**, Harju S, Mäkitie O, Saha MT, Dunkel L, Sankilampi U. Systematic growth monitoring for the early detection of celiac disease in children. *JAMA Pediatr* 2015; **169**: e1525 [PMID: 25730696 DOI: 10.1001/jamapediatrics.2015.25]

60 **Singh P**, Sharma PK, Agnihotri A, Jyotsna VP, Das P, Gupta SD, Makharia GK, Khadgawat R. Coeliac disease in patients with short stature: A tertiary care centre experience. *Natl Med J India* 2015; **28**: 176-180 [PMID: 27132724]

61 **Troncone R**, Kosova R. Short stature and catch-up growth in celiac disease. *J Pediatr Gastroenterol Nutr* 2010; **51** Suppl 3: S137-S138 [PMID: 21088537 DOI: 10.1097/MPG.0b013e3181f1dd66]

62 **Patwari AK**, Kapur G, Satyanarayana L, Anand VK, Jain A, Gangil A, Balani B. Catch-up growth in children with late-diagnosed coeliac disease. *Br J Nutr* 2005; **94**: 437-442 [PMID: 16176616 DOI: 10.1079/bjn20051479]

63 **Giovenale D**, Meazza C, Cardinale GM, Sposito M, Mastrangelo C, Messini B, Citro G, Delvecchio M, Di Maio S, Bozzola M. The prevalence of growth hormone deficiency and celiac disease in short children. *Clin Med Res* 2006; **4**: 180-183 [PMID: 16988097 DOI: 10.3121/cmr.4.3.180]

64 **Leffler DA**, Green PH, Fasano A. Extraintestinal manifestations of coeliac disease. *Nat Rev Gastroenterol Hepatol* 2015; **12**: 561-571 [PMID: 26260366 DOI: 10.1038/nrgastro.2015.131]

65 **Abaci A**, Esen I, Unuvar T, Arslan N, Bober E. Two cases presenting with pubertal delay and diagnosed as Celiac disease. *Clin Pediatr (Phila)* 2008; **47**: 607-609 [PMID: 18566358 DOI: 10.1177/0009922808316185]

66 **Philip R**, Patidar P, Saran S, Agarwal P, Arya T, Gupta K. Endocrine manifestations of celiac disease. *Indian J Endocrinol Metab* 2012; **16**: S506-S508 [PMID: 23565481 DOI: 10.4103/2230-8210.104149]

67 **Traggiai C**, Stanhope R. Disorders of pubertal development. *Best Pract Res Clin Obstet Gynaecol* 2003; **17**: 41-56 [PMID: 12758225 DOI: 10.1053/ybeog.2003.0360]

68 **Kalayci AG**, Kanber Y, Birinci A, Yildiz L, Albayrak D. The prevalence of coeliac disease as detected by screening in children with iron deficiency anaemia. *Acta Paediatr* 2005; **94**: 678-681 [PMID: 16188768 DOI: 10.1111/j.1651-2227.2005.tb01964.x]

69 **Baydoun A**, Maakaron JE, Halawi H, Abou Rahal J, Taher AT. Hematological manifestations of celiac disease. *Scand J Gastroenterol* 2012; **47**: 1401-1411 [PMID: 22861356 DOI: 10.3109/00365521.2012.706828]

70 **Äärelä L**, Nurminen S, Kivelä L, Huhtala H, Mäki M, Viitasalo A, Kaukinen K, Lakka T, Kurppa K. Prevalence and associated factors of abnormal liver values in children with celiac disease. *Dig Liver Dis* 2016; **48**: 1023-1029 [PMID: 27338852 DOI: 10.1016/j.dld.2016.05.022]

71 **Kaukinen K**, Halme L, Collin P, Färkkilä M, Mäki M, Vehmanen P, Partanen J, Höckerstedt K. Celiac disease in patients with severe liver disease: gluten-free diet may reverse hepatic failure. *Gastroenterology* 2002; **122**: 881-888 [PMID: 11910339 DOI: 10.1053/gast.2002.32416]

72 **Anania C**, De Luca E, De Castro G, Chiesa C, Pacifico L. Liver involvement in pediatric celiac disease. *World J Gastroenterol* 2015; **21**: 5813-5822 [PMID: 26019445 DOI: 10.3748/wjg.v21.i19.5813]

73 **Lee GJ**, Boyle B, Ediger T, Hill I. Hypertransaminasemia in Newly Diagnosed Pediatric Patients With Celiac Disease. *J Pediatr Gastroenterol Nutr* 2016; **63**: 340-343 [PMID: 27548248 DOI: 10.1097/MPG.0000000000001153]

74 **Pantaleoni S**, Luchino M, Adriani A, Pellicano R, Stradella D, Ribaldone DG, Sapone N, Isaia GC, Di Stefano M, Astegiano M. Bone mineral density at diagnosis of celiac disease and after 1 year of gluten-free diet. *ScientificWorldJournal* 2014; **2014**: 173082 [PMID: 25379519 DOI: 10.1155/2014/173082]

75 **Keaveny AP**, Freaney R, McKenna MJ, Masterson J, O'Donoghue DP. Bone remodeling indices and secondary hyperparathyroidism in celiac disease. *Am J Gastroenterol* 1996; **91**: 1226-1231 [PMID: 8651176]

76 **Margoni D**, Chouliaras G, Duscas G, Voskaki I, Voutsas N, Papadopoulou A, Panayiotou J, Roma E. Bone health in children with celiac disease assessed by dual x-ray absorptiometry: effect of gluten-free diet and predictive value of serum biochemical indices. *J Pediatr Gastroenterol Nutr* 2012; **54**: 680-684 [PMID: 22094895 DOI: 10.1097/MPG.0b013e31823f5fc5]

77 **Björck S**, Brundin C, Karlsson M, Agardh D. Reduced Bone Mineral Density in Children With Screening-detected Celiac Disease. *J Pediatr Gastroenterol Nutr* 2017; **65**: 526-532 [PMID: 28319607 DOI: 10.1097/MPG.0000000000001568]

78 **Garg K**, Agarwal P, Gupta RK, Sitaraman S. Joint Involvement in Children with Celiac Disease. *Indian Pediatr* 2017; **54**: 946-948 [PMID: 28849767 DOI: 10.1007/s13312-017-1188-x]

79 **Casella G**, Bordo BM, Schalling R, Villanacci V, Salemme M, Di Bella C, Baldini V, Bassotti G. Neurological disorders and celiac disease. *Minerva Gastroenterol Dietol* 2016; **62**: 197-206 [PMID: 26619901]

80 **Lionetti E**, Francavilla R, Pavone P, Pavone L, Francavilla T, Pulvirenti A, Giugno R, Ruggieri M. The neurology of coeliac disease in childhood: what is the evidence? A systematic review and meta-analysis. *Dev Med Child Neurol* 2010; **52**: 700-707 [PMID: 20345955 DOI: 10.1111/j.1469-8749.2010.03647.x]

81 **Ludvigsson JF**, Zingone F, Tomson T, Ekbom A, Ciacci C. Increased risk of epilepsy in biopsy-verified celiac disease: a population-based cohort study. *Neurology* 2012; **78**: 1401-1407 [PMID: 22517096 DOI: 10.1212/WNL.0b013e3182544728]

82 **Zoumpoulakis M**, Fotoulaki M, Topitsoglou V, Lazidou P, Zouloumis L, Kotsanos N. Prevalence of Dental Enamel Defects, Aphthous-Like Ulcers and Other Oral Manifestations in Celiac Children and Adolescents: A Comparative Study. *J Clin Pediatr Dent* 2019; **43**: 274-280 [PMID: 31283894 DOI: 10.17796/1053-4625-43.4.9]

83 **Macho VMP**, de Barros Menéres Manso MCA, E Silva DMV, de Andrade DJC. The difference in symmetry of the enamel defects in celiac disease versus non-celiac pediatric population. *J Dent Sci* 2020; **15**: 345-350 [PMID: 32952893 DOI: 10.1016/j.jds.2020.02.006]

84 **Bucci P**, Carile F, Sangianantoni A, D'Angiò F, Santarelli A, Lo Muzio L. Oral aphthous ulcers and dental enamel defects in children with coeliac disease. *Acta Paediatr* 2006; **95**: 203-207 [PMID: 16449028 DOI: 10.1080/08035250500355022]

85 **Graziano M**, Rossi M. An update on the cutaneous manifestations of coeliac disease and non-coeliac gluten sensitivity. *Int Rev Immunol* 2018; **37**: 291-300 [PMID: 30516407 DOI: 10.1080/08830185.2018.1533008]

86 **Reunala T**, Salmi TT, Hervonen K, Kaukinen K, Collin P. Dermatitis Herpetiformis: A Common Extraintestinal Manifestation of Coeliac Disease. *Nutrients* 2018; **10** [PMID: 29757210 DOI: 10.3390/nu10050602]

87 **Kahaly GJ**, Frommer L, Schuppan D. Celiac Disease and Glandular Autoimmunity. *Nutrients* 2018; **10**: 814 [PMID: 29941778 DOI: 10.3390/nu10070814]

88 **Assa A**, Frenkel-Nir Y, Tzur D, Katz LH, Shamir R. Large population study shows that adolescents with celiac disease have an increased risk of multiple autoimmune and nonautoimmune comorbidities. *Acta Paediatr* 2017; **106**: 967-972 [PMID: 28247429 DOI: 10.1111/apa.13808]

89 **Akirov A**, Pinhas-Hamiel O. Co-occurrence of type 1 diabetes mellitus and celiac disease. *World J Diabetes* 2015; **6**: 707-714 [PMID: 26069719 DOI: 10.4239/wjd.v6.i5.707]

90 **Hermann R**, Turpeinen H, Laine AP, Veijola R, Knip M, Simell O, Sipilä I, Akerblom HK, Ilonen J. HLA DR-DQ-encoded genetic determinants of childhood-onset type 1 diabetes in Finland: an analysis of 622 nuclear families. *Tissue Antigens* 2003; **62**: 162-169 [PMID: 12889996 DOI: 10.1034/j.1399-0039.2003.00071.x]

91 **Elfström P**, Sundström J, Ludvigsson JF. Systematic review with meta-analysis: associations between coeliac disease and type 1 diabetes. *Aliment Pharmacol Ther* 2014; **40**: 1123-1132 [PMID: 25270960 DOI: 10.1111/apt.12973]

92 **Sahin Y,** Cakir MD, Isakoca M, Sahin DA. Prevalence of Celiac Disease in Children with Type 1 Diabetes Mellitus in the South of Turkey. *Iran J Ped* 2020; **30**: e97306 [DOI: 10.5812/ijp.97306]

93 **Weiss B**, Pinhas-Hamiel O. Celiac Disease and Diabetes: When to Test and Treat. *J Pediatr Gastroenterol Nutr* 2017; **64**: 175-179 [PMID: 27574884 DOI: 10.1097/MPG.0000000000001388]

94 **Hansen D**, Brock-Jacobsen B, Lund E, Bjørn C, Hansen LP, Nielsen C, Fenger C, Lillevang ST, Husby S. Clinical benefit of a gluten-free diet in type 1 diabetic children with screening-detected celiac disease: a population-based screening study with 2 years' follow-up. *Diabetes Care* 2006; **29**: 2452-2456 [PMID: 17065683 DOI: 10.2337/dc06-0990]

95 **Salardi S**, Volta U, Zucchini S, Fiorini E, Maltoni G, Vaira B, Cicognani A. Prevalence of celiac disease in children with type 1 diabetes mellitus increased in the mid-1990 s: an 18-year longitudinal study based on anti-endomysial antibodies. *J Pediatr Gastroenterol Nutr* 2008; **46**: 612-614 [PMID: 18493223 DOI: 10.1097/MPG.0b013e31815d697e]

96 **Pham-Short A**, Donaghue KC, Ambler G, Chan AK, Craig ME. Coeliac disease in Type 1 diabetes from 1990 to 2009: higher incidence in young children after longer diabetes duration. *Diabet Med* 2012; **29**: e286-e289 [PMID: 22672045 DOI: 10.1111/j.1464-5491.2012.03720.x]

97 **Chiang JL**, Maahs DM, Garvey KC, Hood KK, Laffel LM, Weinzimer SA, Wolfsdorf JI, Schatz D. Type 1 Diabetes in Children and Adolescents: A Position Statement by the American Diabetes Association. *Diabetes Care* 2018; **41**: 2026-2044 [PMID: 30093549 DOI: 10.2337/dci18-0023]

98 **van der Pals M**, Myléus A, Norström F, Hammarroth S, Högberg L, Rosén A, Ivarsson A, Carlsson A. Body mass index is not a reliable tool in predicting celiac disease in children. *BMC Pediatr* 2014; **14**: 165 [PMID: 24981433 DOI: 10.1186/1471-2431-14-165]

99 **Poulain C**, Johanet C, Delcroix C, Lévy-Marchal C, Tubiana-Rufi N. Prevalence and clinical features of celiac disease in 950 children with type 1 diabetes in France. *Diabetes Metab* 2007; **33**: 453-458 [PMID: 17964843 DOI: 10.1016/j.diabet.2007.06.004]

100 **Midhagen G**, Järnerot G, Kraaz W. Adult coeliac disease within a defined geographic area in Sweden. A study of prevalence and associated diseases. *Scand J Gastroenterol* 1988; **23**: 1000-1004 [PMID: 3201123 DOI: 10.3109/00365528809090160]

101 **Collin P**, Salmi J, Hällström O, Reunala T, Pasternack A. Autoimmune thyroid disorders and coeliac disease. *Eur J Endocrinol* 1994; **130**: 137-140 [PMID: 8130887 DOI: 10.1530/eje.0.1300137]

102 **Sategna-Guidetti C**, Bruno M, Mazza E, Carlino A, Predebon S, Tagliabue M, Brossa C. Autoimmune thyroid diseases and coeliac disease. *Eur J Gastroenterol Hepatol* 1998; **10**: 927-931 [PMID: 9872614 DOI: 10.1097/00042737-199811000-00005]

103 **Korponay-Szabó IR**, Dahlbom I, Laurila K, Koskinen S, Woolley N, Partanen J, Kovács JB, Mäki M, Hansson T. Elevation of IgG antibodies against tissue transglutaminase as a diagnostic tool for coeliac disease in selective IgA deficiency. *Gut* 2003; **52**: 1567-1571 [PMID: 14570724 DOI: 10.1136/gut.52.11.1567]

104 **Carlsson A**, Axelsson I, Borulf S, Bredberg A, Forslund M, Lindberg B, Sjöberg K, Ivarsson SA. Prevalence of IgA-antigliadin antibodies and IgA-antiendomysium antibodies related to celiac disease in children with Down syndrome. *Pediatrics* 1998; **101**: 272-275 [PMID: 9445503 DOI: 10.1542/peds.101.2.272]

105 **Gale L**, Wimalaratna H, Brotodiharjo A, Duggan JM. Down's syndrome is strongly associated with coeliac disease. *Gut* 1997; **40**: 492-496 [PMID: 9176077 DOI: 10.1136/gut.40.4.492]

106 **Bonamico M**, Mariani P, Danesi HM, Crisogianni M, Failla P, Gemme G, Quartino AR, Giannotti A, Castro M, Balli F, Lecora M, Andria G, Guariso G, Gabrielli O, Catassi C, Lazzari R, Balocco NA, De Virgiliis S, Culasso F, Romano C; SIGEP (Italian Society of Pediatric Gastroenterology and Hepatology) and Medical Genetic Group. Prevalence and clinical picture of celiac disease in italian down syndrome patients: a multicenter study. *J Pediatr Gastroenterol Nutr* 2001; **33**: 139-143 [PMID: 11568513 DOI: 10.1097/00005176-200108000-00008]

107 **Book L**, Hart A, Black J, Feolo M, Zone JJ, Neuhausen SL. Prevalence and clinical characteristics of celiac disease in Downs syndrome in a US study. *Am J Med Genet* 2001; **98**: 70-74 [PMID: 11426458]

108 **Zachor DA**, Mroczek-Musulman E, Brown P. Prevalence of celiac disease in Down syndrome in the United States. *J Pediatr Gastroenterol Nutr* 2000; **31**: 275-279 [PMID: 10997372 DOI: 10.1097/00005176-200009000-00014]

109 **Liu E**, Wolter-Warmerdam K, Marmolejo J, Daniels D, Prince G, Hickey F. Routine Screening for Celiac Disease in Children With Down Syndrome Improves Case Finding. *J Pediatr Gastroenterol Nutr* 2020; **71**: 252-256 [PMID: 32304557 DOI: 10.1097/MPG.0000000000002742]

110 **Bonamico M**, Pasquino AM, Mariani P, Danesi HM, Culasso F, Mazzanti L, Petri A, Bona G; Italian Society Of Pediatric Gastroenterology Hepatology (SIGEP); Italian Study Group for Turner Syndrom (ISGTS). Prevalence and clinical picture of celiac disease in Turner syndrome. *J Clin Endocrinol Metab* 2002; **87**: 5495-5498 [PMID: 12466343 DOI: 10.1210/jc.2002-020855]

111 **Gillett PM**, Gillett HR, Israel DM, Metzger DL, Stewart L, Chanoine JP, Freeman HJ. Increased prevalence of celiac disease in girls with Turner syndrome detected using antibodies to endomysium and tissue transglutaminase. *Can J Gastroenterol* 2000; **14**: 915-918 [PMID: 11125180 DOI: 10.1155/2000/172914]

112 **Ivarsson SA**, Carlsson A, Bredberg A, Alm J, Aronsson S, Gustafsson J, Hagenäs L, Häger A, Kriström B, Marcus C, Moëll C, Nilsson KO, Tuvemo T, Westphal O, Albertsson-Wikland K, Aman J. Prevalence of coeliac disease in Turner syndrome. *Acta Paediatr* 1999; **88**: 933-936 [PMID: 10519331 DOI: 10.1080/08035259950168397]

113 **Rujner J**, Wisniewski A, Gregorek H, Wozniewicz B, Młynarski W, Witas HW. Coeliac disease and HLA-DQ 2 (DQA1\* 0501 and DQB1\* 0201) in patients with Turner syndrome. *J Pediatr Gastroenterol Nutr* 2001; **32**: 114-115 [PMID: 11176342 DOI: 10.1097/00005176-200101000-00033]

114 **Nadeem M**, Roche EF. Coeliac disease in Turner syndrome. *Arch Dis Child* 2013; **98**: 649-650 [PMID: 23723336 DOI: 10.1136/archdischild-2013-304126]

115 **Mårild K**, Størdal K, Hagman A, Ludvigsson JF. Turner Syndrome and Celiac Disease: A Case-Control Study. *Pediatrics* 2016; **137**: e20152232 [PMID: 26746404 DOI: 10.1542/peds.2015-2232]

116 **Giannotti A**, Tiberio G, Castro M, Virgilii F, Colistro F, Ferretti F, Digilio MC, Gambarara M, Dallapiccola B. Coeliac disease in Williams syndrome. *J Med Genet* 2001; **38**: 767-768 [PMID: 11694549 DOI: 10.1136/jmg.38.11.767]

117 **Ludvigsson JF**, Leffler DA, Bai JC, Biagi F, Fasano A, Green PH, Hadjivassiliou M, Kaukinen K, Kelly CP, Leonard JN, Lundin KE, Murray JA, Sanders DS, Walker MM, Zingone F, Ciacci C. The Oslo definitions for coeliac disease and related terms. *Gut* 2013; **62**: 43-52 [PMID: 22345659 DOI: 10.1136/gutjnl-2011-301346]

118 **Biagi F**, Bianchi PI, Vattiato C, Marchese A, Trotta L, Badulli C, De Silvestri A, Martinetti M, Corazza GR. Influence of HLA-DQ2 and DQ8 on severity in celiac Disease. *J Clin Gastroenterol* 2012; **46**: 46-50 [PMID: 21694611 DOI: 10.1097/MCG.0b013e318221077e]

119 **Husby S**, Koletzko S, Korponay-Szabó I, Kurppa K, Mearin ML, Ribes-Koninckx C, Shamir R, Troncone R, Auricchio R, Castillejo G, Christensen R, Dolinsek J, Gillett P, Hróbjartsson A, Koltai T, Maki M, Nielsen SM, Popp A, Størdal K, Werkstetter K, Wessels M. European Society Paediatric Gastroenterology, Hepatology and Nutrition Guidelines for Diagnosing Coeliac Disease 2020. *J Pediatr Gastroenterol Nutr* 2020; **70**: 141-156 [PMID: 31568151 DOI: 10.1097/MPG.0000000000002497]

120 **Trovato CM**, Montuori M, Valitutti F, Leter B, Cucchiara S, Oliva S. The Challenge of Treatment in Potential Celiac Disease. *Gastroenterol Res Pract* 2019; **2019**: 8974751 [PMID: 31772571 DOI: 10.1155/2019/8974751]

121 **Tosco A**, Salvati VM, Auricchio R, Maglio M, Borrelli M, Coruzzo A, Paparo F, Boffardi M, Esposito A, D'Adamo G, Malamisura B, Greco L, Troncone R. Natural history of potential celiac disease in children. *Clin Gastroenterol Hepatol* 2011; **9**: 320-5; quiz e36 [PMID: 20851213 DOI: 10.1016/j.cgh.2010.09.006]

122 **Volta U**, Caio G, Boschetti E, Giancola F, Rhoden KJ, Ruggeri E, Paterini P, De Giorgio R. Seronegative celiac disease: Shedding light on an obscure clinical entity. *Dig Liver Dis* 2016; **48**: 1018-1022 [PMID: 27352981 DOI: 10.1016/j.dld.2016.05.024]

123 **Shah VH**, Rotterdam H, Kotler DP, Fasano A, Green PH. All that scallops is not celiac disease. *Gastrointest Endosc* 2000; **51**: 717-720 [PMID: 10840307 DOI: 10.1067/mge.2000.104977]

124 **Greenson JK**. The biopsy pathology of non-coeliac enteropathy. *Histopathology* 2015; **66**: 29-36 [PMID: 25234408 DOI: 10.1111/his.12522]

125 **Mooney PD**, Evans KE, Singh S, Sanders DS. Treatment failure in coeliac disease: a practical guide to investigation and treatment of non-responsive and refractory coeliac disease. *J Gastrointestin Liver Dis* 2012; **21**: 197-203 [PMID: 22720310]

126 **Dewar DH**, Donnelly SC, McLaughlin SD, Johnson MW, Ellis HJ, Ciclitira PJ. Celiac disease: management of persistent symptoms in patients on a gluten-free diet. *World J Gastroenterol* 2012; **18**: 1348-1356 [PMID: 22493548 DOI: 10.3748/wjg.v18.i12.1348]

127 **Werkstetter KJ**, Korponay-Szabó IR, Popp A, Villanacci V, Salemme M, Heilig G, Lillevang ST, Mearin ML, Ribes-Koninckx C, Thomas A, Troncone R, Filipiak B, Mäki M, Gyimesi J, Najafi M, Dolinšek J, Dydensborg Sander S, Auricchio R, Papadopoulou A, Vécsei A, Szitanyi P, Donat E, Nenna R, Alliet P, Penagini F, Garnier-Lengliné H, Castillejo G, Kurppa K, Shamir R, Hauer AC, Smets F, Corujeira S, van Winckel M, Buderus S, Chong S, Husby S, Koletzko S; ProCeDE study group. Accuracy in Diagnosis of Celiac Disease Without Biopsies in Clinical Practice. *Gastroenterology* 2017; **153**: 924-935 [PMID: 28624578 DOI: 10.1053/j.gastro.2017.06.002]

128 **Oberhuber G**, Granditsch G, Vogelsang H. The histopathology of coeliac disease: time for a standardized report scheme for pathologists. *Eur J Gastroenterol Hepatol* 1999; **11**: 1185-1194 [PMID: 10524652 DOI: 10.1097/00042737-199910000-00019]

129 **Rashid M**, MacDonald A. Importance of duodenal bulb biopsies in children for diagnosis of celiac disease in clinical practice. *BMC Gastroenterol* 2009; **9**: 78 [PMID: 19835611 DOI: 10.1186/1471-230X-9-78]

130 **Hill ID**, Fasano A, Guandalini S, Hoffenberg E, Levy J, Reilly N, Verma R. NASPGHAN Clinical Report on the Diagnosis and Treatment of Gluten-related Disorders. *J Pediatr Gastroenterol Nutr* 2016; **63**: 156-165 [PMID: 27035374 DOI: 10.1097/MPG.0000000000001216]

131 **Bascuñán KA**, Roncoroni L, Branchi F, Doneda L, Scricciolo A, Ferretti F, Araya M, Elli L. The 5 Ws of a gluten challenge for gluten-related disorders. *Nutr Rev* 2018; **76**: 79-87 [PMID: 29325090 DOI: 10.1093/nutrit/nux068]

132 **Poddighe D**, Turganbekova A, Baymukasheva D, Saduakas Z, Zhanzakova Z, Abdrakhmanova S. Genetic predisposition to celiac disease in Kazakhstan: Potential impact on the clinical practice in Central Asia. *PLoS One* 2020; **15**: e0226546 [PMID: 31895924 DOI: 10.1371/journal.pone.0226546]

133 **Volta U,** Caio G, De Giorgio R. Mistakes in coeliac disease diagnosis and how to avoid them. *UEG Education* 2016; **16**: 1–3

134 **Singh P**, Arora A, Strand TA, Leffler DA, Mäki M, Kelly CP, Ahuja V, Makharia GK. Diagnostic Accuracy of Point of Care Tests for Diagnosing Celiac Disease: A Systematic Review and Meta-Analysis. *J Clin Gastroenterol* 2019; **53**: 535-542 [PMID: 29912751 DOI: 10.1097/MCG.0000000000001081]

135 **Husby S**, Bai JC. Follow-up of Celiac Disease. *Gastroenterol Clin North Am* 2019; **48**: 127-136 [PMID: 30711205 DOI: 10.1016/j.gtc.2018.09.009]

136 **Bishop J**, Ravikumara M. Coeliac disease in childhood: An overview. *J Paediatr Child Health* 2020; **56**: 1685-1693 [PMID: 33197972 DOI: 10.1111/jpc.14674]

137 **Newton KP**, Singer SA. Celiac disease in children and adolescents: special considerations. *Semin Immunopathol* 2012; **34**: 479-496 [PMID: 22549889 DOI: 10.1007/s00281-012-0313-0]

138 **Catassi C**, Fabiani E, Iacono G, D'Agate C, Francavilla R, Biagi F, Volta U, Accomando S, Picarelli A, De Vitis I, Pianelli G, Gesuita R, Carle F, Mandolesi A, Bearzi I, Fasano A. A prospective, double-blind, placebo-controlled trial to establish a safe gluten threshold for patients with celiac disease. *Am J Clin Nutr* 2007; **85**: 160-166 [PMID: 17209192 DOI: 10.1093/ajcn/85.1.160]

139 **Akobeng AK**, Thomas AG. Systematic review: tolerable amount of gluten for people with coeliac disease. *Aliment Pharmacol Ther* 2008; **27**: 1044-1052 [PMID: 18315587 DOI: 10.1111/j.1365-2036.2008.03669.x]

140 **Ventura A**, Magazzù G, Greco L. Duration of exposure to gluten and risk for autoimmune disorders in patients with celiac disease. SIGEP Study Group for Autoimmune Disorders in Celiac Disease. *Gastroenterology* 1999; **117**: 297-303 [PMID: 10419909 DOI: 10.1053/gast.1999.0029900297]

141 **Andrén Aronsson C**, Lee HS, Hård Af Segerstad EM, Uusitalo U, Yang J, Koletzko S, Liu E, Kurppa K, Bingley PJ, Toppari J, Ziegler AG, She JX, Hagopian WA, Rewers M, Akolkar B, Krischer JP, Virtanen SM, Norris JM, Agardh D; TEDDY Study Group. Association of Gluten Intake During the First 5 Years of Life With Incidence of Celiac Disease Autoimmunity and Celiac Disease Among Children at Increased Risk. *JAMA* 2019; **322**: 514-523 [PMID: 31408136 DOI: 10.1001/jama.2019.10329]

142 **Ludvigsson JF**. Mortality and malignancy in celiac disease. *Gastrointest Endosc Clin N Am* 2012; **22**: 705-722 [PMID: 23083988 DOI: 10.1016/j.giec.2012.07.005]

143 **Valitutti F**, Trovato CM, Montuori M, Cucchiara S. Pediatric Celiac Disease: Follow-Up in the Spotlight. *Adv Nutr* 2017; **8**: 356-361 [PMID: 28298278 DOI: 10.3945/an.116.013292]

144 **Leonard MM**, Fasano A. Zero, One, or Two Endoscopies to Diagnose and Monitor Pediatric Celiac Disease? The Jury Is Still Out. *J Pediatr Gastroenterol Nutr* 2017; **65**: 270-271 [PMID: 28829342 DOI: 10.1097/MPG.0000000000001666]

145 **Mahadev S**, Gardner R, Lewis SK, Lebwohl B, Green PH. Quality of Life in Screen-detected Celiac Disease Patients in the United States. *J Clin Gastroenterol* 2016; **50**: 393-397 [PMID: 26501877 DOI: 10.1097/MCG.0000000000000433]

146 **Webb C**, Myléus A, Norström F, Hammarroth S, Högberg L, Lagerqvist C, Rosén A, Sandström O, Stenhammar L, Ivarsson A, Carlsson A. High adherence to a gluten-free diet in adolescents with screening-detected celiac disease. *J Pediatr Gastroenterol Nutr* 2015; **60**: 54-59 [PMID: 25238121 DOI: 10.1097/MPG.0000000000000571]

147 **Barnea L**, Mozer-Glassberg Y, Hojsak I, Hartman C, Shamir R. Pediatric celiac disease patients who are lost to follow-up have a poorly controlled disease. *Digestion* 2014; **90**: 248-253 [PMID: 25531121 DOI: 10.1159/000368395]

148 **Wessels MM**, van Veen II, Vriezinga SL, Putter H, Rings EH, Mearin ML. Complementary Serologic Investigations in Children with Celiac Disease Is Unnecessary during Follow-Up. *J Pediatr* 2016; **169**: 55-60 [PMID: 26547400 DOI: 10.1016/j.jpeds.2015.09.078]

149 **Husby S**, Murray JA, Katzka DA. AGA Clinical Practice Update on Diagnosis and Monitoring of Celiac Disease-Changing Utility of Serology and Histologic Measures: Expert Review. *Gastroenterology* 2019; **156**: 885-889 [PMID: 30578783 DOI: 10.1053/j.gastro.2018.12.010]

150 **Dipper CR**, Maitra S, Thomas R, Lamb CA, McLean-Tooke AP, Ward R, Smith D, Spickett G, Mansfield JC. Anti-tissue transglutaminase antibodies in the follow-up of adult coeliac disease. *Aliment Pharmacol Ther* 2009; **30**: 236-244 [PMID: 19438848 DOI: 10.1111/j.1365-2036.2009.04039.x]

151 **Isaac DM**, Rajani S, Yaskina M, Huynh HQ, Turner JM. Antitissue Transglutaminase Normalization Postdiagnosis in Children With Celiac Disease. *J Pediatr Gastroenterol Nutr* 2017; **65**: 195-199 [PMID: 27906802 DOI: 10.1097/MPG.0000000000001480]

152 **Mahadev S**, Murray JA, Wu TT, Chandan VS, Torbenson MS, Kelly CP, Maki M, Green PH, Adelman D, Lebwohl B. Factors associated with villus atrophy in symptomatic coeliac disease patients on a gluten-free diet. *Aliment Pharmacol Ther* 2017; **45**: 1084-1093 [PMID: 28220520 DOI: 10.1111/apt.13988]

153 **Moreno ML**, Cebolla Á, Muñoz-Suano A, Carrillo-Carrion C, Comino I, Pizarro Á, León F, Rodríguez-Herrera A, Sousa C. Detection of gluten immunogenic peptides in the urine of patients with coeliac disease reveals transgressions in the gluten-free diet and incomplete mucosal healing. *Gut* 2017; **66**: 250-257 [PMID: 26608460 DOI: 10.1136/gutjnl-2015-310148]

154 **Syage JA**, Kelly CP, Dickason MA, Ramirez AC, Leon F, Dominguez R, Sealey-Voyksner JA. Determination of gluten consumption in celiac disease patients on a gluten-free diet. *Am J Clin Nutr* 2018; **107**: 201-207 [PMID: 29529159 DOI: 10.1093/ajcn/nqx049]

155 **West J**, Logan RF, Card TR, Smith C, Hubbard R. Risk of vascular disease in adults with diagnosed coeliac disease: a population-based study. *Aliment Pharmacol Ther* 2004; **20**: 73-79 [PMID: 15225173 DOI: 10.1111/j.1365-2036.2004.02008.x]

156 **Hallert C**, Grant C, Grehn S, Grännö C, Hultén S, Midhagen G, Ström M, Svensson H, Valdimarsson T. Evidence of poor vitamin status in coeliac patients on a gluten-free diet for 10 years. *Aliment Pharmacol Ther* 2002; **16**: 1333-1339 [PMID: 12144584 DOI: 10.1046/j.1365-2036.2002.01283.x]

157 **Midhagen G**, Hallert C. High rate of gastrointestinal symptoms in celiac patients living on a gluten-free diet: controlled study. *Am J Gastroenterol* 2003; **98**: 2023-2026 [PMID: 14499782 DOI: 10.1111/j.1572-0241.2003.07632.x]

158 **Aziz I**, Evans KE, Papageorgiou V, Sanders DS. Are patients with coeliac disease seeking alternative therapies to a gluten-free diet? *J Gastrointestin Liver Dis* 2011; **20**: 27-31 [PMID: 21451794]

159 **Rashid M**, Cranney A, Zarkadas M, Graham ID, Switzer C, Case S, Molloy M, Warren RE, Burrows V, Butzner JD. Celiac disease: evaluation of the diagnosis and dietary compliance in Canadian children. *Pediatrics* 2005; **116**: e754-e759 [PMID: 16322131 DOI: 10.1542/peds.2005-0904]

160 **Leffler DA**, Kelly CP, Green PH, Fedorak RN, DiMarino A, Perrow W, Rasmussen H, Wang C, Bercik P, Bachir NM, Murray JA. Larazotide acetate for persistent symptoms of celiac disease despite a gluten-free diet: a randomized controlled trial. *Gastroenterology* 2015; **148**: 1311-9.e6 [PMID: 25683116 DOI: 10.1053/j.gastro.2015.02.008]

161 **Lähdeaho ML**, Kaukinen K, Laurila K, Vuotikka P, Koivurova OP, Kärjä-Lahdensuu T, Marcantonio A, Adelman DC, Mäki M. Glutenase ALV003 attenuates gluten-induced mucosal injury in patients with celiac disease. *Gastroenterology* 2014; **146**: 1649-1658 [PMID: 24583059 DOI: 10.1053/j.gastro.2014.02.031]

162 **Murray JA**, Kelly CP, Green PHR, Marcantonio A, Wu TT, Mäki M, Adelman DC; CeliAction Study Group of Investigators. No Difference Between Latiglutenase and Placebo in Reducing Villous Atrophy or Improving Symptoms in Patients With Symptomatic Celiac Disease. *Gastroenterology* 2017; **152**: 787-798.e2 [PMID: 27864127 DOI: 10.1053/j.gastro.2016.11.004]

163 **Anderson RP**, Jabri B. Vaccine against autoimmune disease: antigen-specific immunotherapy. *Curr Opin Immunol* 2013; **25**: 410-417 [PMID: 23478068 DOI: 10.1016/j.coi.2013.02.004]

164 **Rubio-Tapia A**, Ludvigsson JF, Choung RS, Brantner TL, Rajkumar SV, Landgren O, Murray JA. Increased mortality among men aged 50 years old or above with elevated IgA anti-transglutaminase antibodies: NHANES III. *BMC Gastroenterol* 2016; **16**: 136 [PMID: 27809801 DOI: 10.1186/s12876-016-0547-8]

165 **Biagi F**, Gobbi P, Marchese A, Borsotti E, Zingone F, Ciacci C, Volta U, Caio G, Carroccio A, Ambrosiano G, Mansueto P, Corazza GR. Low incidence but poor prognosis of complicated coeliac disease: a retrospective multicentre study. *Dig Liver Dis* 2014; **46**: 227-230 [PMID: 24268568 DOI: 10.1016/j.dld.2013.10.010]

166 **Al-Toma A**, Goerres MS, Meijer JW, Peña AS, Crusius JB, Mulder CJ. Human leukocyte antigen-DQ2 homozygosity and the development of refractory celiac disease and enteropathy-associated T-cell lymphoma. *Clin Gastroenterol Hepatol* 2006; **4**: 315-319 [PMID: 16527694 DOI: 10.1016/j.cgh.2005.12.011]

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**P-Reviewer:** Pavlovic M, Wierzbicka A **S-Editor:** Liu M **L-Editor:** Filipodia **P-Editor:** Wang LYT

**Figure Legends**



**Figure 1 Algorithm for diagnosis of celiac disease.** CD: Celiac disease; DGP: Deamidated gliadin peptide; EMA: Endomysial antibody; tTG:Tissue transglutaminase antibody; ULN: Upper limit of normal.

**Table 1 Groups with higher risk of developing celiac disease**

|  |
| --- |
| **Groups with higher risk of developing celiacdisease** |
| First-degree relatives of celiac patients |
| Second-degree relatives of celiac patients |
| Type 1 diabetes mellitus |
| Autoimmune thyroid disease |
| Autoimmune liver disease |
| Down syndrome |
| Turner syndrome |
| Williams syndrome |
| Selective IgA deficiency |
| Systemic lupus eryhtematosus |
| Juvenile chronic arthritis |

**Table 2 Extra-intestinal manifestations of celiac disease**

|  |
| --- |
| **Extra-intestinal manifestations of celiac disease** |
| Short stature |
| Anemia |
| Osteopenia/osteoporosis |
| Delayed puberty |
| Dental enamel defects |
| Dermatitis herpetiformis |
| Recurrent aphtous stomatitis |
| Neurological manifestations; peripheral neuropathy, epilepsy, ataxia, headache |
| Arthritis, arthralgia |
| Infertility |
| Amenorrhea |
| Elevated liver enzymes |
| Alopecia |
| Anxiety, depression |

**Table 3 Other diseases causing villous atrophy**

|  |
| --- |
| **Other diseases causing villous atrophy** |
| Parasitic infections (*Giardia lamblia*) |
| Autoimmune enteropathy |
| Small intestinal bacterial overgrowth |
| Common variable immunodeficiency |
| Cow's milk or soya protein hypersensitivity |
| Intractable diarrhea of infancy |
| Eosinophilic gastroenteritis |
| Drug induced enteropathy (*e.g.*, olmesartan, mycophenolate) |
| Intestinal lymphoma |
| Crohn's disease |
| Human immunodeficiency virus enteropathy |
| Tropical disease |

**Table 4 The modified Marsh classification**

|  |  |  |  |
| --- | --- | --- | --- |
|  | **IEL** | **Crypts** | **Villi** |
| Type 0 | < 40 | Normal | Normal |
| Type 1 | > 40 | Normal | Normal |
| Type 2 | > 40 | Hypertrophic | Normal |
| Type 3a | > 40 | Hypertrophic | Mild atrophy |
| Type 3b | > 40 | Hypertrophic | Marked atrophy |
| Type 3c | > 40 | Hypertrophic | Absent |

IEL: Intraepithelial lymphocyte count/100 epithelial cells.



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