

Is screening for Celiac Disease Needed in Patients with Liver Disease?

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Abstract Celiac disease is known for its classic clinical manifestations such as anemia and diarrhea. Liver involvement in celiac disease has been studied for more than thirty years. We review different etiologies of liver diseases and their possible association with celiac disease and seek to define in which situations serological screening should be performed.

Keywords: *celiac disease, liver diseases, liver cirrhosis, alanine transaminase, autoimmune hepatitis*

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1. Introduction

Celiac disease is a chronic immune-mediated inflammatory intestinal disorder induced by gluten ingestion in genetically susceptible individuals [1]. It differs substantially from other autoimmune diseases as immunosuppressants are not part of the treatment. The diagnosis requires a joint clinicopathological approach; the recommended first-line test is serology with immunoglobulin A (IgA) tissue transglutaminase (tTG) and IgA endomysial (EmA) antibodies. These serological tests have high levels of sensitivity and specificity, but the presence of lymphocyte infiltrate and villous atrophy in small bowel biopsy is still the gold standard for confirming diagnosis [2]. As in many developing countries the tests for celiac disease are not available, clinicians and institutions should find ways to make confirmatory tests for celiac disease available. Wakim-Fleming *et al* recently demonstrated that in cirrhotic patients, high titers for human tTG and EmA can diagnose celiac disease in the absence of a small bowel biopsy [3]. However, Carocio *et al* pointed out that false positive human tTG can be observed in patients with chronic liver diseases [4]. These facts must be taken into account when facing suspected celiac disease.

Patients may present classical clinical manifestations (weight loss, anemia, diarrhea or general weakness), metabolic bone disease, iron, or folate deficiencies (osteoporosis, fractures, or anemia), or neurologic symptoms (ataxia, encephalopathy, or myelopathy) and associated autoimmune diseases that are often of greater clinical significance than celiac disease [5,6]. When gluten is removed from the diet, most patients present clinical remission and test negative for autoantibodies [7]. High

aminotransferase levels of unknown etiology may also normalize with a gluten-free diet [8].

We aimed to perform a theoretical study of literature on screening of celiac disease in patients with liver disease.

2. Material and Methods

A review for liver involvement in celiac disease was performed by conducting a broad search for "celiac disease" AND "alanine transaminase" "liver cirrhosis" "portal hypertension" "autoimmune hepatitis" "primary biliary cirrhosis" "sclerosing cholangitis" "non-alcoholic fatty liver disease" in Pubmed. Additionally the references of the selected articles were also consulted for relevant articles on the subject. Only relevant full papers were included.

3. Celiac Disease and the Liver

Liver involvement in celiac disease has been studied for more than thirty years [9]. At present, celiac disease screening is recommended for all patients with abnormal liver blood tests, autoimmune liver diseases, prior interferon-based therapy for viral hepatitis, and cirrhosis of various etiologies [10,11]. A Czech study observed that 3% (29/962) of the individuals with liver diseases and 0.8% (5/523) who underwent liver transplant were seropositive for anti-tTG antibodies. However, celiac disease was biopsy-diagnosed in 16 patients: 4 with autoimmune hepatitis type I, 3 with Wilson's disease, 3 with "celiac hepatitis", 2 with primary sclerosing cholangitis, 1 with primary biliary cirrhosis, 1 with Budd-Chiari syndrome, 1 with toxic hepatitis, and 1 with non-alcoholic steatohepatitis. This study demonstrated that celiac disease may be present with any etiology of liver

disease [11]. The fact that two diseases coexist does not mean that there is an association, and the most important questions here are whether there is evidence to recommend celiac disease screening in all patients with liver disease and whether there are possible benefits.

Among patients with cryptogenic hypertransaminasemia, about 3.6% to 4.1% present biopsy-proven celiac disease [12], and in most cases abnormal liver tests normalize with a gluten-free diet [8,12]. When they do not, liver biopsies are mandatory for further investigation of hepatic involvement, which has previously been related to chronic active hepatitis [8]. Nonetheless, when aminotransferase levels remain elevated, screening for autoimmune liver diseases such as autoimmune hepatitis (AIH), primary biliary cirrhosis (PBC), and primary sclerosing cholangitis (PSC) must be performed [13,14,15].

Celiac disease is ten times more prevalent among patients with AIH than in the general population [16]. When both diseases are present, the introduction of a gluten-free diet in immunosuppressed patients can cause a high AIH remission rate. When clinical remission is reached, a prolonged immunosuppressive regimen induces a high sustained remission rate after treatment withdrawal [17].

The association between PBC and celiac disease has long been suspected since Logan *et al* first reported both conditions diagnosed simultaneously in 4 patients [18]. The association remains controversial [19,20,21]. Nevertheless, a gluten-free diet may be helpful in restoration of liver function in patients with PBC and celiac disease [19].

Associations between PSC and celiac disease have been described by many authors [22,23]. One case series showed that anti-smooth muscle antibody (ASMA) was present in 3 patients with PSC, indicating that screening for celiac disease may be of value in patients with PSC specially in the presence of ASMA [22]. More studies including a large number of patients are necessary to define the association between autoimmune liver diseases and celiac disease. This question has not been answered because eating habits and celiac disease prevalence vary around the world. The primary concern at this point is whether patients with autoimmune liver disease should be actively screened for EmA and tTG antibodies, especially those with persistent altered liver tests, in regions where the prevalence of celiac disease is significant. In this context, a gluten-free diet may contribute to normalization of liver biochemistry, may halt progression of the liver disease, and may even prevent complications such as the onset of other autoimmune diseases or lymphoma.

It has been reported that nonalcoholic fatty liver disease (NAFLD) without metabolic syndrome could be related to the concomitant presence of celiac disease. Bardella *et al* evaluated 59 patients with hypertransaminasemia and biopsy-proven NAFLD, and 3.4% were found to have celiac disease [24]. After 6 months of a gluten-free diet, liver enzymes normalized. It was later observed that a significant proportion of NAFLD patients, including obese individuals or those with metabolic syndrome, were seropositive for celiac disease, but EmA and tTG were more commonly found in those with body mass indices less than 27 kg/m². Additionally, their aminotransferase levels and ultrasound steatosis regressed with gluten-free diet [25,26]. A pathogenetic link between NAFLD and celiac disease involving gut permeability, microbiota, and

diet has been proposed, but the pathogenesis of liver steatosis in celiac disease remains uncertain [27,28]. Patients with NAFLD should be screened for celiac disease when steatohepatitis is present in the absence of metabolic risk factors and once other causes of the liver disease are excluded [27].

Few studies have addressed the relationship between celiac disease and chronic hepatitis B [29,30], and the results are controversial. It has been demonstrated that vaccines against the hepatitis B virus are less efficacious in patients with celiac disease [31,32]. While celiac disease is considered an autoimmune disorder, this reduced efficacy may reflect defective mechanisms in antibody production and viral clearance. A higher prevalence of celiac disease has been observed in those infected with hepatitis C virus (HCV) than those without it [33,34]. However, a French multi-center study failed to demonstrate such association, probably because of the low prevalence of celiac disease in France [35]. Despite the controversy over the link between these two diseases, the primary concern is that patients may present severe cases of overt celiac disease during HCV treatment, leading to discontinuation of interferon-alfa (IFN) [36]. Patients with both diseases may experience severe diarrhea with weight loss during IFN treatment as well as dermatitis herpetiformis, hypoferritinemia, and refractory anemia that persist after treatment has stopped. Until recently, the association of pegylated interferon-alfa (PEG) with ribavirin was the gold standard treatment for hepatitis C [37]. All HCV patients should be screened for celiac disease prior to anti-viral treatment. For those with positive antibodies, IFN-free regimens are the logical choice. If the new drugs (protease inhibitors of 2nd and 3rd generation) are not available, a gluten-free diet must be started preemptively, and patients should be carefully monitored during IFN treatment [34,36,38].

Some case reports have suggested that celiac disease may be a trigger for the development of idiopathic noncirrhotic intrahepatic portal hypertension (NCIHPH) [39,40]. In India, 10% of NCIHPH patients had biopsy-proven celiac disease [41]. Furthermore, the presence of celiac disease predicts reduced transplant-free survival [42]. Current data suggests the need to search for celiac disease in all patients with unexplained portal hypertension [41], although we do not know if a gluten-free diet can change the evolution of the disease or improve survival.

Celiac disease is at least twice more common in cirrhotic patients than in the general population [3]. The absence of a common histological pattern of liver injury in patients with celiac disease does not favor the assumption that this disease directly damages the liver [43]. Kaukinen *et al* described patients with chronic liver disease before or after liver transplant in which detection and treatment of celiac disease prevented progression to end-stage liver failure. Three were in consideration for liver transplant. Hepatic dysfunction reversed in all patients when a gluten-free diet was adopted [44]. Besides these, other cases were reported where decompensated cirrhosis reversed after the introduction of a gluten-free diet [43,45,46]. These data indicate that celiac disease should be screened in cirrhotic patients, especially in those with hypoalbuminemia and ascites [3,43]. Independent of the etiology of liver cirrhosis, patients with advanced liver disease and celiac disease may benefit from a gluten-free diet.

4. Conclusion

Although major advances were made over the last decades, future studies are still needed to define the prevalence of celiac disease in various etiologies of liver diseases and to prove any alleged association. Also, the impact of celiac disease in several etiologies of liver disease is yet to be determined. Based on the current knowledge, serologic screening for celiac disease should be done in the setting of cryptogenic hypertransaminasemia, autoimmune liver diseases, NAFLD in the absence of classical risk factors, idiopathic noncirrhotic portal hypertension, and advanced cirrhosis of any etiology.

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