

Extra-intestinal manifestations of non-celiac gluten sensitivity: An expanding paradigm

Giuseppe Losurdo, Mariabeatrice Principi, Andrea Iannone, Annacinzia Amoruso, Enzo Ierardi, Alfredo Di Leo, Michele Barone

Giuseppe Losurdo, Mariabeatrice Principi, Andrea Iannone, Annacinzia Amoruso, Enzo Ierardi, Alfredo Di Leo, Michele Barone, Section of Gastroenterology, Department of Emergency and Organ Transplantation, University "Aldo Moro" of Bari, Bari 70124, Italy

ORCID number: Giuseppe Losurdo (0000-0001-7038-3287); Mariabeatrice Principi (0000-0003-0545-5656); Andrea Iannone (0000-0002-5468-9515); Annacinzia Amoruso (0000-0002-4161-3378); Enzo Ierardi (0000-0001-7275-5080); Alfredo Di Leo (0000-0003-2026-1200); Michele Barone (0000-0003-2026-1200).

Author contributions: Losurdo G, Barone M and Di Leo A planned the study; Principi M, Iannone A, Amoruso A and Barone M collected the data; Losurdo G and Ierardi E wrote the paper; Barone M and Di Leo A revised the article; Losurdo G and Principi M contributed equally and are co-first authors.

Conflict-of-interest statement: None declared.

Open-Access: This article is an open-access article which was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

Manuscript source: Invited manuscript

Correspondence to: Giuseppe Losurdo, MD, Doctor, Section of Gastroenterology, Department of Emergency and Organ Transplantation, University "Aldo Moro" of Bari, Piazza Giulio Cesare 11, Bari 70124, Italy. giuseppelos@alice.it
Telephone: +39-80-5593452
Fax: +39-80-5593088

Received: March 8, 2018

Peer-review started: March 9, 2018

First decision: March 14, 2018

Revised: March 27, 2018

Accepted: March 31, 2018

Article in press: March 31, 2018

Published online: April 14, 2018

Abstract

Non celiac gluten sensitivity (NCGS) is a syndrome characterized by a cohort of symptoms related to the ingestion of gluten-containing food in subjects who are not affected by celiac disease (CD) or wheat allergy. The possibility of systemic manifestations in this condition has been suggested by some reports. In most cases they are characterized by vague symptoms such as 'foggy mind', headache, fatigue, joint and muscle pain, leg or arm numbness even if more specific complaints have been described. NCGS has an immune-related background. Indeed there is a strong evidence that a selective activation of innate immunity may be the trigger for NCGS inflammatory response. The most commonly autoimmune disorders associated to NCGS are Hashimoto thyroiditis, dermatitis herpetiformis, psoriasis and rheumatologic diseases. The predominance of Hashimoto thyroiditis represents an interesting finding, since it has been indirectly confirmed by an Italian study, showing that autoimmune thyroid disease is a risk factor for the evolution towards NCGS in a group of patients with minimal duodenal inflammation. On these bases, an autoimmune stigma in NCGS is strongly supported; it could be a characteristic feature that could help the diagnosis and be simultaneously managed. A possible neurological involvement has been underlined by NCGS association with gluten ataxia, gluten neuropathy and gluten encephalopathy. NCGS patients may show even psychiatric diseases such as depression, anxiety and psychosis. Finally, a link with functional disorders (irritable bowel syndrome and fibromyalgia) is a topic under discussion. In conclusion, the novelty of this matter has generated an expansion of literature data with the unavoidable consequence that some

reports are often based on low levels of evidence. Therefore, only studies performed on large samples with the inclusion of control groups will be able to clearly establish whether the large information from the literature regarding extra-intestinal NCGS manifestations could be supported by evidence-based agreements.

Key words: Non celiac gluten sensitivity; Celiac disease; Gluten; Gluten ataxia; Autoimmunity; Gluten-related disorders; Thyroiditis; Extra-intestinal

© **The Author(s) 2018.** Published by Baishideng Publishing Group Inc. All rights reserved.

Core tip: Non celiac gluten sensitivity is an expanding field of investigation within gluten-related disorders. Similarly to celiac disease, it shows a systemic involvement, therefore several extra-intestinal manifestations have been hypothesized and investigated in many studies. They may involve many districts and have neurological/psychiatric, dermatological, rheumatologic and nutritional implications. Moreover, the possibility of association with other autoimmune diseases should not be underestimated. However, the large data amount from the literature often requires to be supported by evidence-based agreements.

Losurdo G, Principi M, Iannone A, Amoruso A, Ierardi E, Di Leo A, Barone M. Extra-intestinal manifestations of non-celiac gluten sensitivity: An expanding paradigm. *World J Gastroenterol* 2018; 24(14): 1521-1530 Available from: URL: <http://www.wjgnet.com/1007-9327/full/v24/i14/1521.htm> DOI: <http://dx.doi.org/10.3748/wjg.v24.i14.1521>

INTRODUCTION

Non celiac gluten sensitivity (NCGS) is a syndrome characterized by a set of symptoms related to the ingestion of gluten-containing food in subjects who are not affected by celiac disease (CD) or wheat allergy^[1]. Despite it has been included in the spectrum of gluten related disorders, it shows a peculiar picture with some elements resembling CD, *i.e.*, immunological involvement and response to gluten free diet, and some features close to irritable bowel syndrome^[2].

In detail, NCGS is distinguished by symptoms that typically take place soon after gluten ingestion, withdraw with gluten exclusion, and relapse following gluten challenge within hours or days. The "classical" clinical picture of NCGS is a combination of irritable bowel syndrome-like manifestations, such as abdominal pain, bloating, diarrhea or alterations in bowel habit with alternation of constipation and loose stools.

However, the possibility of systemic manifestations in this condition has been suggested by some reports. In most cases they are characterized by vague symptoms such as 'foggy mind', headache, fatigue, joint and

muscle pain, leg or arm numbness even if more specific complaints have been described, such as dermatitis, (eczema or skin rash), depression, neurological symptoms and anemia^[3-8]. Moreover, the possibility of association with other autoimmune diseases has been hypothesized. Indeed, similarly to CD, NCGS can be considered as an immune system-related disease and this aspect should be of relevance.

In conclusion, the spectrum of NCGS extra-intestinal manifestations is constantly expanding with new reports. Therefore, we aimed to summarize the main extra-intestinal manifestations of NCGS in a narrative review. In particular, in this review we focused on the associations supported by an evidence-based link more than single case reports, where it is difficult to differentiate a casual association from a real relationship. For this reason we searched in PubMed database in February 2018 using the following terms: gluten sensitivity, extra-intestinal, autoimmune, thyroid, neurology, psychiatry, rheumatology, skin, dermatology, nutrition, irritable bowel syndrome and fibromyalgia. In this way, 880 articles were found, and, as reported in the flow chart in Figure 1, we selected 86 studies for this review. Other studies which were not focused on NCGS or reporting an unclear definition of NCGS, or in which results about extra-intestinal manifestation were not listed have been excluded. Additionally, we graded the level of evidence on the association between NCGS and systemic manifestations using the Oxford consensus^[9].

ASSOCIATION WITH AUTOIMMUNE DISEASES

On the base of convincing evidence, NCGS has an immune-related background. Indeed it has been demonstrated that a selective activation of innate immunity may be the trigger for NCGS inflammatory response^[10,11]. It is unclear whether gliadin is the real responsible for the autoimmune event onset, since some other components of wheat, such as amylase-trypsin inhibitors or fermentable oligo-di-mono-saccharides and polyols (FODMAPs) have been invoked^[12-14]. For this reason some Authors consider the term "non celiac wheat sensitivity" more appropriate than the current one^[15].

CD, which is the most common and studied gluten-related disorder, is often associated to several other autoimmune diseases, such as type 1 diabetes, autoimmune thyroiditis or dermatitis herpetiformis^[16]. For this reason it is conceivable that also patients with NCGS could show autoimmune disorders. In a cohort of 131 NCGS patients^[17], the prevalence of autoimmune disease (29%) was found to be higher than in control group (4%, $P < 0.001$). Moreover, anti-nucleus antibody (ANA) positivity, a well-known marker of autoimmune setting, was present in the 46% of NCGS subjects, compared to the 2% of controls, and ANA positivity

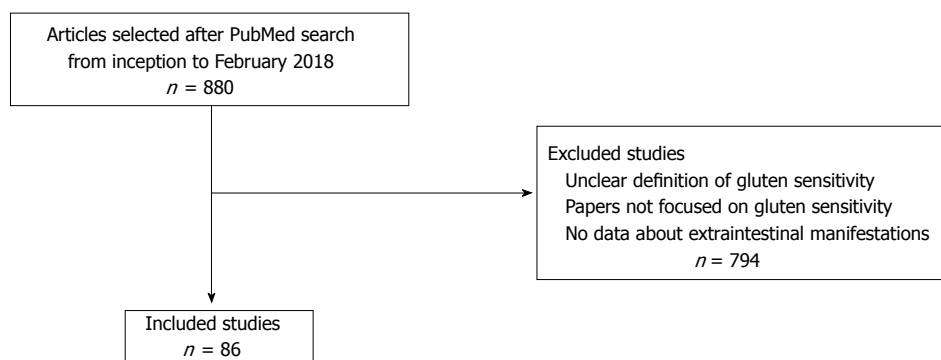


Figure 1 Flowchart summarizing the process of study selection.

correlated with DQ2/8 haplotypes. In detail, the most frequently reported NCGS-associated autoimmune disorder was Hashimoto thyroiditis (29 patients). Other diseases were psoriasis (4 cases), type 1 diabetes (4 cases), mixed connective tissue disease (1 case) and ankylosing spondylitis (1 case). The predominance of autoimmune thyroiditis represents an interesting finding, since it was indirectly confirmed by an Italian experience^[18], showing that autoimmune thyroiditis is a risk factor for the evolution towards NCGS in a group of patients with minimal duodenal inflammation^[19].

On these bases, an autoimmune stigma in NCGS is strongly supported; it could be a characteristic feature that could help the diagnosis and be simultaneously managed.

NEUROLOGIC AND PSYCHIATRIC MANIFESTATIONS

Recently, many studies explored the bond between the ingestion of gluten-containing food and the onset of neurologic and psychiatric disorders or symptoms such as ataxia, peripheral neuropathy, schizophrenia, autism, depression, anxiety, and hallucinations^[20].

In patients with CD, a neurological involvement could be the only clinical manifestation of the disease. The production of autoantibodies directed against the tissue transglutaminase isoform 6 (expressed selectively in brain tissue) has been found in up to the 85% of these patients^[21,22]. Anti-gliadin antibodies (AGA) frequently occur in such cases^[21,22]. It is unclear whether the production of these antibodies takes place in the brain or in the gut mucosa, but these antibodies are considered to be the etiologic agent of neurological manifestations of CD. Finally, an inflammatory infiltrate of T lymphocytes resembling IELs in the white matter or in perivascular cuff of nerves is an important finding suggesting a specific pathogenetic mechanism of gluten-induced neuropathies^[23].

Three main diseases have been described in the spectrum of gluten-related neurologic manifestations: gluten ataxia, gluten neuropathy and gluten encephalopathy^[23].

Gluten ataxia has the strongest relationship with gluten-related disorders. It encompasses about the 20% of all causes of ataxia. This is mainly characterized by pure cerebellar ataxia and, rarely, by ataxia combined with myoclonus, palatal tremor, opsoclonus, or chorea. Gaze-evoked nystagmus and other ocular marks of cerebellar dysfunction are observed in about the 80%. All subjects show gait ataxia and most of them have limb ataxia^[24]. A frequent finding at magnetic resonance imaging is cerebellar atrophy, secondary to necrosis of Purkinje cells^[25]. Less than 10% of patients with gluten ataxia complain of gastrointestinal symptoms. A gluten free diet is able to reverse symptoms, however an early diagnosis significantly improves the prognosis, since gluten free diet may stop the loss of Purkinje cells. Therefore, a late diagnosis may be associated with an irreversible damage^[26].

Gluten neuropathy is a form of peripheral neuronal damage, in which there is a serological evidence of CD positivity in the absence of alternative aetiologies. The most common type is a symmetrical sensorimotor axonal peripheral neuropathy, but other types have also been described (asymmetrical neuropathy, pure motor neuropathy or autonomic neuropathy)^[27]. Gluten neuropathy occurs in the sixth decade and slowly progresses with a 9 year mean latency time between the diagnosis of neuropathy and that of CD. A third of patients shows duodenal inflammation on biopsy, however the presence or absence of enteropathy does not influence the effect of a gluten-free diet^[28]. The most common histopathological feature of gluten neuropathy is lymphocyte infiltration of peri-neural vessels^[29].

Gluten encephalopathy is a central nervous system disease characterized by focal abnormalities of the white matter (usually area of low perfusion) in presence of AGA or anti-transglutaminase 2 antibodies^[30]. The most common symptom is migraine. It has been demonstrated that a gluten free diet improves the headaches and stops the progression of cerebral alterations detected at magnetic resonance imaging^[31].

Some reports about the direct relationship between the above cited diseases and NCGS have been

published in the last years. Hadjivassiliou *et al*^[32] have retrospectively evaluated 562 patients with gluten-related disorders (228 CD and 334 NCGS) and concomitant neurological involvement. In NCGS the most frequent disorder was peripheral neuropathy (54%) followed by ataxia (46%) and encephalopathy, while in CD, ataxia was the most frequent one (41%). In all cases a deep linkage with AGA positivity was recorded. Additionally, the severity of ataxia was similar in both conditions (CD and NCGS), while patients with CD exhibited more frequently severe forms of neuropathy. Rodrigo *et al*^[33] found, in a cohort of 31 subjects with gluten ataxia, AGA positivity rate of 100%; this value was more similar to NCGS (89%) than CD (48%) and was associated to Marsh 1 duodenal histological picture. On the bases of such results, they concluded that gluten ataxia shows a strict affinity to NCGS more than CD.

Headache is a very frequent finding in NCGS. However, no study has so far analyzed in depth the nature of this association. The available data relies mainly on observational studies aiming to elucidate the prevalence of this condition, which ranges around the 25%^[3-8,34,35]. However, the lack of case-control studies is a serious limitation to ascertain the reliability of the association. Moreover there are no studies investigating possible pathogenetic mechanisms.

The association with other neurologic diseases such as epilepsy^[36], miopathy^[37] and demyelinating disease^[38], is anecdotal or based on a non conventional diagnosis of NCGS, therefore it is not possible to draw solid conclusions.

Among the psychiatric diseases, depression and anxiety have been hypothesized as systemic manifestations of NCGS. In an Australian study^[39], a group of patients with established diagnosis of NCGS underwent a double blind crossover study with a placebo versus oral gluten supplementation after a gluten free diet. Results showed that gluten induced depression scale worsening when compared to placebo, while other symptoms (anxiety, curiosity and anger) were not influenced by the diet. However, the mechanism by which gluten may induce these changes is not yet clear. Depression is indeed a frequent finding in Western society, and it could be a distinctive mood tract of personality rather than an extra-intestinal manifestation of NCGS. However, in another study NCGS patients did not exhibit a tendency for general somatization. Additionally, personality and quality of life did not differ between NCGS and CD patients and were mostly similar to healthy controls^[40].

Some authors have invoked a role of gluten for some psychiatric diseases like schizophrenia or bipolar disorder^[41], but there are no studies exploring these entities in NCGS. On the other hand, some cases of "gluten psychosis" in patients with NCGS have been described^[42]. In these patients, hallucinations, crying spells, relevant confusion, ataxia, severe anxiety and paranoid delirium occurred shortly after gluten ingestion

and disappeared within one week of gluten free diet.

Finally, the relationship between autism and gluten is an hot topic. It has been shown that children with autism have more frequently IgG-AGA positivity than healthy children (24% vs 7%)^[43], but currently there are no studies in which a solid diagnosis of NCGS has been achieved in autistic subjects. A gluten free diet is often proposed to these children in an empiric setting, since it has been demonstrated that it improves behavioral scores^[44,45]. However, at present there are no evidence-based reasons to look for gluten sensitivity in autism and to advise an exclusion diet^[46].

SKIN MANIFESTATIONS

The association between CD and skin diseases, in particular dermatitis herpetiformis, is well known^[47]. Similarly to CD, the possibility of a skin involvement in the 18% of NCGS has been reported^[4]. In the published case series^[3-8], undefined dermatitis, rash and eczema were the most common skin manifestations in NCGS. The possibility of an association with skin autoimmune diseases such as psoriasis has been above mentioned^[17]. A case report has shown that even dermatitis herpetiformis may occur^[48].

Some reports have been mainly focused on the characteristics of skin lesions in NCGS from a dermatological point of view. In a series of 17 NCGS patients with skin lesions, the most common ones were very similar to dermatitis herpetiformis or subacute eczema (erythematous, excoriated papular-vesicular and extremely itchy)^[49]. Some patients had also hyperkeratotic scaly lesions resembling psoriasis. The most common skin location was the extensor surfaces of upper limbs, in the 94%, alike dermatitis herpetiformis. The histological analysis showed complement C3 deposits at dermoepidermal junction in the 82%. Finally, in all patients a gluten free diet was able to lead to lesions disappearance within one month, much faster than in dermatitis herpetiformis.

Some Authors have claimed that an allergic sensitivity to food allergens other than gluten could underlie NCGS^[50]. Indeed, an Italian study found that the 10% of NCGS patients suffered from nickel allergy with contact dermatitis and this prevalence was higher than in control group (5%, $P = 0.04$). However, NCGS subjects referred onset of dermatitis after wheat ingestion^[51].

RHEUMATOLOGIC MANIFESTATIONS

As we already mentioned, NCGS shows the tendency to cluster autoimmune diseases. Some reports about its coexistence with rheumatologic diseases are available. The first evidence demonstrated that in a group of 30 subjects with ankylosing spondylitis, 11 had AGA positivity, while no patient in a control group exhibited this finding^[52]. Isasi *et al*^[53] reported 4 cases of axial spondyloarthritis (2 ankylosing spondylitis and 1

Table 1 Studies reporting the prevalence of people avoiding gluten-containing foods

Ref.	Country	Population	Sample size	Avoidance rate of gluten-based products
Tanpowpong <i>et al</i> ^[60] , 2012	New Zealand	Pediatric	916	5.2%
Rubio-Tapia <i>et al</i> ^[61] , 2013	United States	Pediatric	7798	0.7%
DiGiacomo <i>et al</i> ^[62] , 2013	United States	National Health and Nutrition Examination Survey	7762	0.6%
Lis <i>et al</i> ^[63] , 2014	Australia	Adults	910	41.2%
Golley <i>et al</i> ^[64] , 2015	Australia	Adults	1184	10.6%
Mardini <i>et al</i> ^[65] , 2015	United States	Pediatric	14701	1%
Aziz <i>et al</i> ^[59] , 2014	United Kingdom	Adults	1002	3.7%
Van Gils <i>et al</i> ^[8] , 2016	The Netherlands	Adults	785	6.2%
Carroccio <i>et al</i> ^[7] , 2017	Italy	Adolescents	548	2.9%

psoriatic spondyloarthritis) with a microscopic enteritis picture at duodenal biopsy. They all underwent a gluten free diet, and in all cases an improvement or remission of back pain was reported, with a recrudescence after wheat challenge. The same result was recorded in another group of patients with systemic sclerosis, Raynaud's phenomenon, symmetric polyarthritis and Sjogren's syndrome^[53].

However, despite such reports, the evidence for NCGS/rheumatologic association is weak, since case reports represent only a low level of evidence and case-control studies are necessary.

FIBROMYALGIA AND OTHER FUNCTIONAL DISORDERS

Fibromyalgia is a disease characterized by widespread pain, often accompanied by fatigue, memory problems, sleep disturbances, depression or irritable bowel syndrome^[54]. In many case series, several NCGS patients complain of chronic muscle or joint pain, leg numbness, fatigue and headache^[3-8], therefore it is possible that an underlying undiagnosed fibromyalgia could be present. Indeed, starting from some case reports demonstrating this association^[55], further studies have analyzed in depth this relationship. In a Spanish series^[56] of 246 fibromyalgia patients undergoing gluten free diet, 90 showed clinical symptom improvement. Additionally, Authors described the features of 20 out of such 90 patients. They had a mean duration of fibromyalgia of 12 years, and 17 had also gastrointestinal symptoms. Eighteen had a DQ2/8 haplotype and all showed an increase in duodenal IELs. After a mean gluten free diet period of 16.4 mo, 15 of them (75%) experienced a full remission of pain and in 8 of them gluten challenge led to symptom re-appearance. In another trial, gluten free diet was able to induce a decrease in some scales evaluating fibromyalgia symptoms^[57]. On these bases, it is possible to hypothesize that the link between these two disorders is quite strong, but the role of microscopic enteritis in this setting should be tested in other controlled trials.

Fibromyalgia is frequently recognized as a functional disease. In this regard, NCGS has a tight bond with

irritable bowel syndrome (IBS)^[58]. Many patients with IBS often identify some foods that they believe to be more offending, and wheat is often invoked. Furthermore, a certain symptom overlap between NCGS and IBS-type symptoms exists^[4,59]. For this reason, many patients tend to exclude gluten from their diet on their own, without medical advice, as summarized in Table 1^[7,8,59-65]. The basic difference between the two conditions is that patients with NCGS assert that symptoms take place when they eat wheat so that they believe to have identified gluten as the culprit. Some experimental investigations have shown that gliadin can alter the integrity of the small intestinal mucosa, as shown by the appearance of epithelial leaks/gaps and widened inter-villous spaces detected by using confocal laser endomicroscopy^[66]. Based on these assumptions, some clinical trials have demonstrated that a gluten free diet may lead to improvement of gastrointestinal symptoms in IBS, as reported in Table 2^[5,67-73]. However it is not clear whether gluten is really the responsible for such symptoms. Indeed wheat contains FODMAPs as well, which are considered as a possible trigger for IBS itself, and FODMAP restriction demonstrated an improvement in IBS symptoms in up to the 74%^[74]. Additionally, one trial underlined that subjects with self-reported NCGS (and IBS-like symptoms) had benefits by a low FODMAP diet despite they were still consuming a gluten free diet^[75]. Based on these evidences, the link between IBS and NCGS seems to be strict even if quite nebulous. Is it possible that IBS and NCGS should be considered as the two sides of the same coin? Such fascinating question needs to be answered by well designed studies for this purpose.

NUTRITIONAL IMPAIRMENT IN NCGS

CD is often disclosed by nutritional impairments, such as vitamin D or iron deficiency, anemia or alterations in bone mineralization^[76,77].

Anemia prevalence value ranges between 15% and 23% in NCGS^[3,4]. Nevertheless, studies enclosing a control group are lacking, therefore it is not possible to establish which is the real relationship between anemia and NCGS. Additionally, folate deficiency has

Table 2 Main studies exploring the effect of gluten free diet in irritable bowel syndrome

Ref.	Country	Population	Outcome
Wahnschaffe <i>et al</i> ^[67] , 2001	Germany	102 IBS-D	Stool frequency/bowel movement improved in DQ2-8 positive subjects
Aziz <i>et al</i> ^[68] , 2016	United Kingdom	40 IBS-D	A 6-wk GFD reduced symptoms in 70%
Vazquez-Roque <i>et al</i> ^[69] , 2013	United States	45 IBS-D	Stool frequency/bowel movement reduced in patients under GFD
Di Sabatino <i>et al</i> ^[5] , 2015	Italy	59 IBS with self-diagnosis of NCGS	A challenge with 4 g/d of gluten worsened symptoms compared to placebo
Shahbazkhani <i>et al</i> ^[70] , 2015	Iran	72 IBS	Worsening of intestinal symptoms with gluten compared to placebo
Zanwar <i>et al</i> ^[71] , 2016	India	60 IBS	A 4-wk GFD improved a visual-analogue scale of symptoms
Elli <i>et al</i> ^[72] , 2016	Italy	140 IBS with self-diagnosis of NCGS	Only the 14% showed a response to GFD as well as challenge test
Barmeyer <i>et al</i> ^[73] , 2017	Germany	34 IBS	The 34% showed clinical improvement to GFD and continued for one year

GFD: Gluten free diet; IBS-D: Irritable bowel syndrome, diarrhea subtype; NCGS: Non celiac gluten sensitivity.

Table 3 Main extra-intestinal manifestations of non-celiac gluten sensitivity and associated disorders

Manifestations	Extra-intestinal manifestations	Level of evidence	Associated disorders	Level of evidence
General symptoms	Tiredness	4	Aphthous stomatitis	4
	Lack of wellbeing	4		
	Foggy mind	4		
	Joint or muscle pain	4		
	Arm/leg numbness	4		
Neurologic manifestations			Ataxia	3b
			Neuropathy	3b
			Encephalopathy	3b
			Epilepsy	4
			Miopathy	4
			Myelopathy	4
			Demyelinating disease	4
Psychiatric manifestations	Depression	1c	Bipolar disorder	4
	Anxiety	1c	Gluten psychosis	4
			Autism	2b
			Schizophrenia	4
Other autoimmune diseases and rheumatologic diseases			Psoriasis	2b
			Autoimmune thyroiditis	2b
			Rheumatoid arthritis	4
			Scleroderma	4
			Sjogren syndrome	4
			Raynaud phenomenon	4
Skin diseases			Dermatitis herpetiformis	2b
			Contact dermatitis	2b
			Rash and undetermined dermatitis	2b
Functional disorders			Fibromyalgia	1c
			Irritable bowel syndrome	1c
Nutritional imbalance	Anemia	4		
	Osteoporosis	2b		
Other			Interstitial cystitis	4
			Ingrown hairs	4
			Rhinitis, asthma	4
			Postural tachycardia syndrome	2b
			Oligo- or polymenorrhea	4

The level of evidence was expressed according to the Oxford consensus^[85]

been reported in NCGS with solid evidence and it has been even described as a predictive factor for its development^[18].

An Italian study illustrated that NCGS carries a risk of osteopenia similar to CD^[78]. Low bone mineral density

measured by Dual-energy X-ray absorptiometry was found in 28% of NCGS subjects, vs 6% of IBS as well as an influence of body mass index on mineralization was observed. This result has been explained by a lower calcium dietary intake (only 615 mg/d, while

recommended dose is 1000 mg/d).

This last observation may suggest that NCGS patients could experience an alteration in macro- and micronutrients intake due to dietary self-restrictions. Indeed Zingone *et al.*^[79] evaluated diet habits of 29 NCGS subjects and discovered that they ingested lower mean amounts of carbohydrates, proteins, fiber, and polyunsaturated fatty acids. Patients with NCGS reported avoiding fruit, vegetables, milk, and dairy products as well as snacks and mixed spices when compared to a control population.

NCGS is characterized by absent or minimal duodenal inflammation and, therefore, cannot be associated to nutrient deficiencies linked to malabsorption. However, an inflammatory status of duodenal mucosa, witnessed by increased expression of interferon gamma, may not be overlooked^[33,80-82]. Finally, alterations in dietary pattern should not be underestimated. Gluten free diet itself can lead to an inadequate balance in macronutrients assumption^[83-85].

CONCLUSION

Data from literature about extra-intestinal manifestations of NCGS strongly suggests that this condition could have a systemic involvement, similarly to CD. However, the novelty of this topic has generated an expansion of literature data with the unavoidable consequence that some reports are often based on low levels of evidence, as summarized in Table 3, with a grading of evidence according to the Oxford classification^[9]. Therefore, only studies performed on large samples with the addition of control groups will be able to clearly establish whether the large information from the literature regarding extra-intestinal NCGS manifestations could be supported by evidence-based agreements.

REFERENCES

- 1 **Catassi C**, Bai JC, Bonaz B, Bouma G, Calabrò A, Carroccio A, Castillejo G, Ciacci C, Cristofori F, Dolinsek J, Francavilla R, Elli L, Green P, Holtmeier W, Koehler P, Koletzko S, Meinhold C, Sanders D, Schumann M, Schuppan D, Ullrich R, Vécsei A, Volta U, Zevallos V, Sapone A, Fasano A. Non-Celiac Gluten sensitivity: the new frontier of gluten related disorders. *Nutrients* 2013; **5**: 3839-3853 [PMID: 24077239 DOI: 10.3390/nu5103839]
- 2 **Sapone A**, Bai JC, Ciacci C, Dolinsek J, Green PH, Hadjivassiliou M, Kaukinen K, Rostami K, Sanders DS, Schumann M, Ullrich R, Villalta D, Volta U, Catassi C, Fasano A. Spectrum of gluten-related disorders: consensus on new nomenclature and classification. *BMC Med* 2012; **10**: 13 [PMID: 22313950 DOI: 10.1186/1741-7015-10-13]
- 3 **Volta U**, Tovoli F, Cicola R, Parisi C, Fabbri A, Piscaglia M, Fiorini E, Caio G. Serological tests in gluten sensitivity (nonceliac gluten intolerance). *J Clin Gastroenterol* 2012; **46**: 680-685 [PMID: 22138844 DOI: 10.1097/MCG.0b013e3182372541]
- 4 **Volta U**, Bardella MT, Calabrò A, Troncone R, Corazza GR; Study Group for Non-Celiac Gluten Sensitivity. An Italian prospective multicenter survey on patients suspected of having non-celiac gluten sensitivity. *BMC Med* 2014; **12**: 85 [PMID: 24885375 DOI: 10.1186/1741-7015-12-85]
- 5 **Di Sabatino A**, Volta U, Salvatore C, Biancheri P, Caio G, De Giorgio R, Di Stefano M, Corazza GR. Small Amounts of Gluten in Subjects With Suspected Nonceliac Gluten Sensitivity: A Randomized, Double-Blind, Placebo-Controlled, Cross-Over Trial. *Clin Gastroenterol Hepatol* 2015; **13**: 1604-1612.e3 [PMID: 25701700 DOI: 10.1016/j.cgh.2015.01.029]
- 6 **Carroccio A**, Mansueto P, Iacono G, Soresi M, D'Alcamo A, Cavataio F, Brusca I, Florena AM, Ambrosiano G, Seidita A, Pirrone G, Rini GB. Non-celiac wheat sensitivity diagnosed by double-blind placebo-controlled challenge: exploring a new clinical entity. *Am J Gastroenterol* 2012; **107**: 1898-1906; quiz 1907 [PMID: 22825366 DOI: 10.1038/ajg.2012.236]
- 7 **Carroccio A**, Giambalvo O, Blasca F, Iacobucci R, D'Alcamo A, Mansueto P. Self-Reported Non-Celiac Wheat Sensitivity in High School Students: Demographic and Clinical Characteristics. *Nutrients* 2017; **9**: pii: E771 [PMID: 28753927 DOI: 10.3390/nu9070771]
- 8 **van Gils T**, Nijeboer P, IJssennagger CE, Sanders DS, Mulder CJ, Bouma G. Prevalence and Characterization of Self-Reported Gluten Sensitivity in The Netherlands. *Nutrients* 2016; **8**: pii: E714 [PMID: 27834802 DOI: 10.3390/nu8110714]
- 9 **Centre for Evidence-Based Medicine**. Oxford Centre for Evidence-based Medicine – Levels of Evidence (March 2009). Accessed March 17, 2018 Available from: URL: <https://www.cebm.net/2009/06/oxford-centre-evidence-based-medicine-levels-evidence-march-2009/>
- 10 **Caio G**, Riegler G, Patturelli M, Facchiano A, DE Magistris L, Sapone A. Pathophysiology of non-celiac gluten sensitivity: where are we now? *Minerva Gastroenterol Dietol* 2017; **63**: 16-21 [PMID: 27808487 DOI: 10.23736/S1121-421X.16.02346-1]
- 11 **Sapone A**, Lammers KM, Casolaro V, Cammarota M, Giuliano MT, De Rosa M, Stefanile R, Mazzarella G, Tolone C, Russo MI, Esposito P, Ferraraccio F, Carteni M, Riegler G, de Magistris L, Fasano A. Divergence of gut permeability and mucosal immune gene expression in two gluten-associated conditions: celiac disease and gluten sensitivity. *BMC Med* 2011; **9**: 23 [PMID: 21392369 DOI: 10.1186/1741-7015-9-23]
- 12 **Zevallos VF**, Raker V, Tenzer S, Jimenez-Calvente C, Ashfaq-Khan M, Rüssel N, Pickert G, Schild H, Steinbrink K, Schuppan D. Nutritional Wheat Amylase-Trypsin Inhibitors Promote Intestinal Inflammation via Activation of Myeloid Cells. *Gastroenterology* 2017; **152**: 1100-1113.e12 [PMID: 27993525 DOI: 10.1053/j.gastro.2016.12.006]
- 13 **Skodje GI**, Sarna VK, Minelle IH, Rolfsen KL, Muir JG, Gibson PR, Veierød MB, Henriksen C, Lundin KEA. Fructan, Rather Than Gluten, Induces Symptoms in Patients With Self-Reported Non-Celiac Gluten Sensitivity. *Gastroenterology* 2018; **154**: 529-539.e2 [PMID: 29102613 DOI: 10.1053/j.gastro.2017.10.040]
- 14 **De Giorgio R**, Volta U, Gibson PR. Sensitivity to wheat, gluten and FODMAPs in IBS: facts or fiction? *Gut* 2016; **65**: 169-178 [PMID: 26078292 DOI: 10.1136/gutjnl-2015-309757]
- 15 **Carroccio A**, Rini G, Mansueto P. Non-celiac wheat sensitivity is a more appropriate label than non-celiac gluten sensitivity. *Gastroenterology* 2014; **146**: 320-321 [PMID: 24275240 DOI: 10.1053/j.gastro.2013.08.061]
- 16 **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]
- 17 **Carroccio A**, D'Alcamo A, Cavataio F, Soresi M, Seidita A, Sciumè C, Geraci G, Iacono G, Mansueto P. High Proportions of People With Nonceliac Wheat Sensitivity Have Autoimmune Disease or Antinuclear Antibodies. *Gastroenterology* 2015; **149**: 596-603.e1 [PMID: 26026392 DOI: 10.1053/j.gastro.2015.05.040]
- 18 **Losurdo G**, Piscitelli D, Giangaspero A, Principi M, Buffelli F, Giorgio F, Montenegro L, Sorrentino C, Amoroso A, Ierardi E, Di Leo A. Evolution of nonspecific duodenal lymphocytosis over 2 years of follow-up. *World J Gastroenterol* 2015; **21**: 7545-7552 [PMID: 26140001 DOI: 10.3748/wjg.v21.i24.7545]
- 19 **Ierardi E**, Losurdo G, Iannone A, Piscitelli D, Amoroso A, Barone M, Principi M, Pisani A, Di Leo A. Lymphocytic duodenitis or microscopic enteritis and gluten-related conditions: what needs

- to be explored? *Ann Gastroenterol* 2017; **30**: 380-392 [PMID: 28655974 DOI: 10.20524/aog.2017.0165]
- 20 **Jackson JR**, Eaton WW, Cascella NG, Fasano A, Kelly DL. Neurologic and psychiatric manifestations of celiac disease and gluten sensitivity. *Psychiatr Q* 2012; **83**: 91-102 [PMID: 21877216 DOI: 10.1007/s11126-011-9186-y]
- 21 **Hadjivassiliou M**, Aeschlimann P, Strigun A, Sanders DS, Woodroffe N, Aeschlimann D. Autoantibodies in gluten ataxia recognize a novel neuronal transglutaminase. *Ann Neurol* 2008; **64**: 332-343 [PMID: 18825674 DOI: 10.1002/ana.21450]
- 22 **Zis P**, Rao DG, Sarrigiannis PG, Aeschlimann P, Aeschlimann DP, Sanders D, Grünewald RA, Hadjivassiliou M. Transglutaminase 6 antibodies in gluten neuropathy. *Dig Liver Dis* 2017; **49**: 1196-1200 [PMID: 28886934 DOI: 10.1016/j.dld.2017.08.019]
- 23 **Hadjivassiliou M**, Sanders DS, Grünewald RA, Woodroffe N, Boscolo S, Aeschlimann D. Gluten sensitivity: from gut to brain. *Lancet Neurol* 2010; **9**: 318-330 [PMID: 20170845 DOI: 10.1016/S1474-4422(09)70290-X]
- 24 **Hadjivassiliou M**, Sanders DS, Woodroffe N, Williamson C, Grünewald RA. Gluten ataxia. *Cerebellum* 2008; **7**: 494-498 [PMID: 18787912 DOI: 10.1007/s12311-008-0052-x]
- 25 **Wilkinson ID**, Hadjivassiliou M, Dickson JM, Wallis L, Grünewald RA, Coley SC, Widjaja E, Griffiths PD. Cerebellar abnormalities on proton MR spectroscopy in gluten ataxia. *J Neurol Neurosurg Psychiatry* 2005; **76**: 1011-1013 [PMID: 15965215]
- 26 **Pellecchia MT**, Scala R, Perretti A, De Michele G, Santoro L, Filla A, Ciacci C, Barone P. Cerebellar ataxia associated with subclinical celiac disease responding to gluten-free diet. *Neurology* 1999; **53**: 1606-1608 [PMID: 10534283]
- 27 **Chin RL**, Sander HW, Brannagan TH, Green PH, Hays AP, Alaedini A, Latov N. Celiac neuropathy. *Neurology* 2003; **60**: 1581-1585 [PMID: 12771245]
- 28 **Hadjivassiliou M**, Kandler RH, Chattopadhyay AK, Davies-Jones AG, Jarratt JA, Sanders DS, Sharrack B, Grünewald RA. Dietary treatment of gluten neuropathy. *Muscle Nerve* 2006; **34**: 762-766 [PMID: 17013890]
- 29 **Hadjivassiliou M**, Grünewald RA, Kandler RH, Chattopadhyay AK, Jarratt JA, Sanders DS, Sharrack B, Wharton SB, Davies-Jones GA. Neuropathy associated with gluten sensitivity. *J Neurol Neurosurg Psychiatry* 2006; **77**: 1262-1266 [PMID: 16835287]
- 30 **Hadjivassiliou M**, Grünewald RA, Lawden M, Davies-Jones GA, Powell T, Smith CM. Headache and CNS white matter abnormalities associated with gluten sensitivity. *Neurology* 2001; **56**: 385-388 [PMID: 11171906]
- 31 **Serratrice J**, Disdier P, de Roux C, Christides C, Weiller PJ. Migraine and coeliac disease. *Headache* 1998; **38**: 627-628 [PMID: 11398309]
- 32 **Hadjivassiliou M**, Rao DG, Grünewald RA, Aeschlimann DP, Sarrigiannis PG, Hoggard N, Aeschlimann P, Mooney PD, Sanders DS. Neurological Dysfunction in Coeliac Disease and Non-Coeliac Gluten Sensitivity. *Am J Gastroenterol* 2016; **111**: 561-567 [PMID: 26832652 DOI: 10.1038/ajg.2015.434]
- 33 **Rodrigo L**, Hernández-Lahoz C, Lauret E, Rodríguez-Peláez M, Soucek M, Ciccocioppo R, Kruzliak P. Gluten ataxia is better classified as non-celiac gluten sensitivity than as celiac disease: a comparative clinical study. *Immunol Res* 2016; **64**: 558-564 [PMID: 26676361 DOI: 10.1007/s12026-015-8750-1]
- 34 **Losurdo G**, Giorgio F, Piscitelli D, Montenegro L, Covelli C, Fiore MG, Giangaspero A, Iannone A, Principi M, Amoroso A, Barone M, Di Leo A, Ierardi E. May the assessment of baseline mucosal molecular pattern predict the development of gluten related disorders among microscopic enteritis? *World J Gastroenterol* 2016; **22**: 8017-8025 [PMID: 27672296 DOI: 10.3748/wjg.v22.i35.8017]
- 35 **Losurdo G**, Piscitelli D, Pezzuto F, Fortarezza F, Covelli C, Marra A, Iannone A, Amoroso A, Principi M, Ierardi E, Di Leo A. T Helper Lymphocyte and Mast Cell Immunohistochemical Pattern in Nonceliac Gluten Sensitivity. *Gastroenterol Res Pract* 2017; **2017**: 5023680 [PMID: 29362561 DOI: 10.1155/2017/5023680]
- 36 **Bruni O**, Dosi C, Luchetti A, Della Corte M, Riccioni A, Battaglia D, Ferri R. An unusual case of drug-resistant epilepsy in a child with non-celiac gluten sensitivity. *Seizure* 2014; **23**: 674-676 [PMID: 24813861 DOI: 10.1016/j.seizure.2014.04.005]
- 37 **Hadjivassiliou M**, Chattopadhyay AK, Grünewald RA, Jarratt JA, Kandler RH, Rao DG, Sanders DS, Wharton SB, Davies-Jones GA. Myopathy associated with gluten sensitivity. *Muscle Nerve* 2007; **35**: 443-450 [PMID: 17143894]
- 38 **Jacob S**, Zarei M, Kenton A, Allroggen H. Gluten sensitivity and neuromyelitis optica: two case reports. *J Neurol Neurosurg Psychiatry* 2005; **76**: 1028-1030 [PMID: 15965221]
- 39 **Peters SL**, Biesiekierski JR, Yelland GW, Muir JG, Gibson PR. Randomised clinical trial: gluten may cause depression in subjects with non-coeliac gluten sensitivity - an exploratory clinical study. *Aliment Pharmacol Ther* 2014; **39**: 1104-1112 [PMID: 24689456 DOI: 10.1111/apt.12730]
- 40 **Brottveit M**, Vandvik PO, Wojnusz S, Løvik A, Lundin KE, Boye B. Absence of somatization in non-coeliac gluten sensitivity. *Scand J Gastroenterol* 2012; **47**: 770-777 [PMID: 22519894 DOI: 10.3109/00365521.2012.679685]
- 41 **Casella G**, Pozzi R, Cigognetti M, Banchetti F, Torti G, Cadei M, Villanacci V, Baldini V, Bassotti G. Mood disorders and non-celiac gluten sensitivity. *Minerva Gastroenterol Dietol* 2017; **63**: 32-37 [PMID: 27647538 DOI: 10.23736/S1121-421X.16.02325-4]
- 42 **Lionetti E**, Leonardi S, Franzonello C, Mancardi M, Ruggieri M, Catassi C. Gluten Psychosis: Confirmation of a New Clinical Entity. *Nutrients* 2015; **7**: 5532-5539 [PMID: 26184290 DOI: 10.3390/nu7075235]
- 43 **Lau NM**, Green PH, Taylor AK, Hellberg D, Ajamian M, Tan CZ, Kosofsky BE, Higgins JJ, Rajadhyaksha AM, Alaedini A. Markers of Celiac Disease and Gluten Sensitivity in Children with Autism. *PLoS One* 2013; **8**: e66155 [PMID: 23823064 DOI: 10.1371/journal.pone.0066155]
- 44 **Ghalichi F**, Ghaemmaghami J, Malek A, Ostadrahimi A. Effect of gluten free diet on gastrointestinal and behavioral indices for children with autism spectrum disorders: a randomized clinical trial. *World J Pediatr* 2016; **12**: 436-442 [PMID: 27286693 DOI: 10.1007/s12519-016-0040-z]
- 45 **El-Rashidy O**, El-Baz F, El-Gendy Y, Khalaf R, Reda D, Saad K. Ketogenic diet versus gluten free casein free diet in autistic children: a case-control study. *Metab Brain Dis* 2017; **32**: 1935-1941 [PMID: 28808808 DOI: 10.1007/s11011-017-0088-z]
- 46 **Cruchet S**, Lucero Y, Cornejo V. Truths, Myths and Needs of Special Diets: Attention-Deficit/Hyperactivity Disorder, Autism, Non-Celiac Gluten Sensitivity, and Vegetarianism. *Ann Nutr Metab* 2016; **68** Suppl 1: 43-50 [PMID: 27356007 DOI: 10.1159/000445393]
- 47 **Collin P**, Salmi TT, Hervonen K, Kaukinen K, Reunala T. Dermatitis herpetiformis: a cutaneous manifestation of coeliac disease. *Ann Med* 2017; **49**: 23-31 [PMID: 27499257 DOI: 10.1080/07853890.2016.1222450]
- 48 **Philips C**, Paramaguru R, Indiran DA, Augustine P. Dermatitis Herpetiformis as the Initial Presentation of Primary Biliary Cholangitis in a Male with Gluten Sensitivity. *Cureus* 2017; **9**: e1247 [PMID: 28630805 DOI: 10.7759/cureus.1247]
- 49 **Bonciolini V**, Bianchi B, Del Bianco E, Verdelli A, Caproni M. Cutaneous Manifestations of Non-Celiac Gluten Sensitivity: Clinical Histological and Immunopathological Features. *Nutrients* 2015; **7**: 7798-7805 [PMID: 26389946 DOI: 10.3390/nu7095368]
- 50 **Mansueto P**, D'Alcamo A, Seidita A, Carroccio A. Food allergy in irritable bowel syndrome: The case of non-celiac wheat sensitivity. *World J Gastroenterol* 2015; **21**: 7089-7109 [PMID: 26109796 DOI: 10.3748/wjg.v21.i23.7089]
- 51 **D'Alcamo A**, Mansueto P, Soresi M, Iacobucci R, Blasca F, Geraci G, Cavataio F, Fayer F, Arini A, Di Stefano L, Iacono G, Bosco L, Carroccio A. Contact Dermatitis Due to Nickel Allergy in Patients Suffering from Non-Celiac Wheat Sensitivity. *Nutrients* 2017; **9**: pii: E103 [PMID: 28157173 DOI: 10.3390/nu9020103]
- 52 **Toğrol RE**, Nalbant S, Solmazgöl E, Ozyurt M, Kaplan M, Kiralp MZ, Dinçer U, Sahan B. The significance of coeliac disease antibodies in patients with ankylosing spondylitis: a case-controlled

- study. *J Int Med Res* 2009; **37**: 220-226 [PMID: 19215694 DOI: 10.1177/147323000903700127]
- 53 **Isasi C**, Tejerina E, Morán LM. Non-celiac gluten sensitivity and rheumatic diseases. *Reumatol Clin* 2016; **12**: 4-10 [PMID: 25956352 DOI: 10.1016/j.reuma.2015.03.001]
- 54 **Clauw DJ**. Fibromyalgia: a clinical review. *JAMA* 2014; **311**: 1547-1555 [PMID: 24737367 DOI: 10.1001/jama.2014.3266]
- 55 **Isasi C**, Tejerina E, Fernandez-Puga N, Serrano-Vela JI. Fibromyalgia and chronic fatigue syndrome caused by non-celiac gluten sensitivity. *Reumatol Clin* 2015; **11**: 56-57 [PMID: 25047419 DOI: 10.1016/j.reuma.2014.06.005]
- 56 **Isasi C**, Colmenero I, Casco F, Tejerina E, Fernandez N, Serrano-Vela JI, Castro MJ, Villa LF. Fibromyalgia and non-celiac gluten sensitivity: a description with remission of fibromyalgia. *Rheumatol Int* 2014; **34**: 1607-1612 [PMID: 247280274 DOI: 10.1007/s00296-014-2990-6]
- 57 **Slim M**, Calandre EP, Garcia-Leiva JM, Rico-Villademoros F, Molina-Barea R, Rodriguez-Lopez CM, Morillas-Arques P. The Effects of a Gluten-free Diet Versus a Hypocaloric Diet Among Patients With Fibromyalgia Experiencing Gluten Sensitivity-like Symptoms: A Pilot, Open-Label Randomized Clinical Trial. *J Clin Gastroenterol* 2017; **51**: 500-507 [PMID: 27548732 DOI: 10.1097/MCG.0000000000000651]
- 58 **Verdu EF**, Armstrong D, Murray JA. Between celiac disease and irritable bowel syndrome: the “no man’s land” of gluten sensitivity. *Am J Gastroenterol* 2009; **104**: 1587-1594 [PMID: 19455131 DOI: 10.1038/ajg.2009.188]
- 59 **Aziz I**, Lewis NR, Hadjivassiliou M, Winfield SN, Rugg N, Kelsall A, Newrick L, Sanders DS. A UK study assessing the population prevalence of self-reported gluten sensitivity and referral characteristics to secondary care. *Eur J Gastroenterol Hepatol* 2014; **26**: 33-39 [PMID: 24216570 DOI: 10.1097/01.meg.0000435546.87251.f7]
- 60 **Tanpowpong P**, Ingham TR, Lampshire PK, Kirchberg FF, Epton MJ, Crane J, Camargo CA Jr; New Zealand Asthma and Allergy Cohort Study Group. Coeliac disease and gluten avoidance in New Zealand children. *Arch Dis Child* 2012; **97**: 12-16 [PMID: 22075107 DOI: 10.1136/archdischild-2011-300248]
- 61 **Rubio-Tapia A**, Ludvigsson JF, Brantner TL, Murray JA, Everhart JE. The prevalence of celiac disease in the United States. *Am J Gastroenterol* 2012; **107**: 1538-44; quiz 1537, 1545 [PMID: 22850429 DOI: 10.1038/ajg.2012.219]
- 62 **DiGiacomo DV**, Tennyson CA, Green PH, Demmer RT. Prevalence of gluten-free diet adherence among individuals without celiac disease in the USA: results from the Continuous National Health and Nutrition Examination Survey 2009-2010. *Scand J Gastroenterol* 2013; **48**: 921-925 [PMID: 23834276 DOI: 10.3109/00365521.2013.809598]
- 63 **Lis DM**, Stellingwerff T, Shing CM, Ahuja KD, Fell JW. Exploring the popularity, experiences, and beliefs surrounding gluten-free diets in nonceliac athletes. *Int J Sport Nutr Exerc Metab* 2015; **25**: 37-45 [PMID: 24901744 DOI: 10.1123/ijsnem.2013-0247]
- 64 **Golley S**, Corsini N, Topping D, Morell M, Mohr P. Motivations for avoiding wheat consumption in Australia: results from a population survey. *Public Health Nutr* 2015; **18**: 490-499 [PMID: 24739252 DOI: 10.1017/S1368980014000652]
- 65 **Mardini HE**, Westgate P, Grigorian AY. Racial Differences in the Prevalence of Celiac Disease in the US Population: National Health and Nutrition Examination Survey (NHANES) 2009-2012. *Dig Dis Sci* 2015; **60**: 1738-1742 [PMID: 25577269 DOI: 10.1007/s10620-014-3514-7]
- 66 **Fritscher-Ravens A**, Schuppan D, Ellrichmann M, Schoch S, Röcken C, Brasch J, Bethge J, Böttner M, Klose J, Milla PJ. Confocal endomicroscopy shows food-associated changes in the intestinal mucosa of patients with irritable bowel syndrome. *Gastroenterology* 2014; **147**: 1012-1020.e4 [PMID: 25083606 DOI: 10.1053/j.gastro.2014.07.046]
- 67 **Wahnschaffe U**, Ullrich R, Riecken EO, Schulzke JD. Celiac disease-like abnormalities in a subgroup of patients with irritable bowel syndrome. *Gastroenterology* 2001; **121**: 1329-1338 [PMID: 11729112]
- 68 **Aziz I**, Trott N, Briggs R, North JR, Hadjivassiliou M, Sanders DS. Efficacy of a Gluten-Free Diet in Subjects With Irritable Bowel Syndrome-Diarrhea Unaware of Their HLA-DQ2/8 Genotype. *Clin Gastroenterol Hepatol* 2016; **14**: 696-703.e1 [PMID: 26748221 DOI: 10.1016/j.cgh.2015.12.031]
- 69 **Vazquez-Roque MI**, Camilleri M, Smyrk T, Murray JA, Marietta E, O’Neill J, Carlson P, Lamsam J, Janzow D, Eckert D, Burton D, Zinsmeister AR. A controlled trial of gluten-free diet in patients with irritable bowel syndrome-diarrhea: effects on bowel frequency and intestinal function. *Gastroenterology* 2013; **144**: 903-911.e3 [PMID: 23357715 DOI: 10.1053/j.gastro.2013.01.049]
- 70 **Shahbazzkhani B**, Sadeghi A, Malekzadeh R, Khatavi F, Etemadi M, Kalantri E, Rostami-Nejad M, Rostami K. Non-Celiac Gluten Sensitivity Has Narrowed the Spectrum of Irritable Bowel Syndrome: A Double-Blind Randomized Placebo-Controlled Trial. *Nutrients* 2015; **7**: 4542-4554 [PMID: 26056920 DOI: 10.3390/nu7064542]
- 71 **Zanwar VG**, Pawar SV, Gambhire PA, Jain SS, Surude RG, Shah VB, Contractor QQ, Rathi PM. Symptomatic improvement with gluten restriction in irritable bowel syndrome: a prospective, randomized, double blinded placebo controlled trial. *Intest Res* 2016; **14**: 343-350 [PMID: 27799885]
- 72 **Elli L**, Tomba C, Branchi F, Roncoroni L, Lombardo V, Bardella MT, Ferretti F, Conte D, Valiante F, Fini L, Forti E, Cannizzaro R, Maiero S, Londoni C, Lauri A, Fornaciari G, Lenoci N, Spagnuolo R, Basilisco G, Somalvico F, Borgatta B, Leandro G, Segato S, Barisani D, Morreale G, Buscarini E. Evidence for the Presence of Non-Celiac Gluten Sensitivity in Patients with Functional Gastrointestinal Symptoms: Results from a Multicenter Randomized Double-Blind Placebo-Controlled Gluten Challenge. *Nutrients* 2016; **8**: 84 [PMID: 26867199 DOI: 10.3390/nu8020084]
- 73 **Barmeyer C**, Schumann M, Meyer T, Zielinski C, Zuberbier T, Siegmund B, Schulzke JD, Daum S, Ullrich R. Long-term response to gluten-free diet as evidence for non-celiac wheat sensitivity in one third of patients with diarrhea-dominant and mixed-type irritable bowel syndrome. *Int J Colorectal Dis* 2017; **32**: 29-39 [PMID: 27695975 DOI: 10.1007/s00384-016-2663-x]
- 74 **Shepherd SJ**, Gibson PR. Fructose malabsorption and symptoms of irritable bowel syndrome: guidelines for effective dietary management. *J Am Diet Assoc* 2006; **106**: 1631-1639 [PMID: 17000196 DOI: 10.1016/j.jada.2006.07.010]
- 75 **Biesiekierski JR**, Peters SL, Newnham ED, Rosella O, Muir JG, Gibson PR. No effects of gluten in patients with self-reported non-celiac gluten sensitivity after dietary reduction of fermentable, poorly absorbed, short-chain carbohydrates. *Gastroenterology* 2013; **145**: 320-8.e1-3 [PMID: 23648697 DOI: 10.1053/j.gastro.2013.04.051]
- 76 **Abenavoli L**, Delibasic M, Peta V, Turkulov V, De Lorenzo A, Medić-Stojanoska M. Nutritional profile of adult patients with celiac disease. *Eur Rev Med Pharmacol Sci* 2015; **19**: 4285-4292 [PMID: 26636515]
- 77 **García-Manzanares A**, Lucendo AJ. Nutritional and dietary aspects of celiac disease. *Nutr Clin Pract* 2011; **26**: 163-173 [PMID: 21447770 DOI: 10.1177/0884533611399773]
- 78 **Carroccio A**, Soresi M, D’Alcamo A, Sciumè C, Iacono G, Geraci G, Brusca I, Seidita A, Adragna F, Carta M, Mansueto P. Risk of low bone mineral density and low body mass index in patients with non-celiac wheat-sensitivity: a prospective observation study. *BMC Med* 2014; **12**: 230 [PMID: 25430806 DOI: 10.1186/s12916-014-0230-2]
- 79 **Zingone F**, Bartalini C, Siniscalchi M, Ruotolo M, Bucci C, Morra I, Iovino P, Ciacci C. Alterations in Diets of Patients With Nonceliac Gluten Sensitivity Compared With Healthy Individuals. *Clin Gastroenterol Hepatol* 2017; **15**: 63-68.e2 [PMID: 27552857 DOI: 10.1016/j.cgh.2016.08.017]
- 80 **Ierardi E**, Amoroso A, Giorgio F, Principi M, Losurdo G, Piscitelli D, Buffelli F, Fiore MG, Mongelli A, Castellaneta NM, Giangaspero A, De Francesco V, Montenegro L, Di Leo A. Mucosal molecular pattern of tissue transglutaminase and interferon gamma in suspected seronegative celiac disease at marsh

- 1 and 0 stages. *Saudi J Gastroenterol* 2015; **21**: 379-385 [PMID: 26655133 DOI: 10.4103/1319-3767.167189]
- 81 **Dinu M**, Macchia D, Pagliai G, Gori AM, Cesari F, Marcucci R, Sofi F, Casini A. Symptomatic efficacy of buckwheat products in Non-Celiac Gluten Sensitivity (NCGS). *Asia Pac J Clin Nutr* 2017; **26**: 630-636 [PMID: 28582812 DOI: 10.6133/apjcn.072016.07]
- 82 **Brottveit M**, Beitnes AC, Tollefsen S, Bratlie JE, Jahnsen FL, Johansen FE, Sollid LM, Lundin KE. Mucosal cytokine response after short-term gluten challenge in celiac disease and non-celiac gluten sensitivity. *Am J Gastroenterol* 2013; **108**: 842-850 [PMID: 23588237 DOI: 10.1038/ajg.2013.91]
- 83 **Vici G**, Belli L, Biondi M, Polzonetti V. Gluten free diet and nutrient deficiencies: A review. *Clin Nutr* 2016; **35**: 1236-1241 [PMID: 27211234 DOI: 10.1016/j.clnu.2016.05.002]
- 84 **Barone M**, Della Valle N, Rosania R, Facciorusso A, Trotta A, Cantatore FP, Falco S, Pignatiello S, Viggiani MT, Amoruso A, De Filippis R, Di Leo A, Francavilla R. A comparison of the nutritional status between adult celiac patients on a long-term, strictly gluten-free diet and healthy subjects. *Eur J Clin Nutr* 2016; **70**: 23-27 [PMID: 26173865 DOI: 10.1038/ejcn.2015.114]
- 85 **Shepherd SJ**, Gibson PR. Nutritional inadequacies of the gluten-free diet in both recently-diagnosed and long-term patients with coeliac disease. *J Hum Nutr Diet* 2013; **26**: 349-358 [PMID: 23198728 DOI: 10.1111/jhn.12018]

P- Reviewer: Jadallah KA, Rostami-Nejad M, Sergi CM
S- Editor: Gong ZM **L- Editor:** A **E- Editor:** Huang Y





Published by **Baishideng Publishing Group Inc**
7901 Stoneridge Drive, Suite 501, Pleasanton, CA 94588, USA
Telephone: +1-925-223-8242
Fax: +1-925-223-8243
E-mail: bpgoffice@wjgnet.com
Help Desk: <http://www.f6publishing.com/helpdesk>
<http://www.wjgnet.com>



ISSN 1007-9327

