

Revista Latinoamericana de Hipertensión ISSN: 1856-4550 latinoamericanadehipertension@gmail.com Sociedad Latinoamericana de Hipertensión Venezuela

# Correlation of IDA severity with endoscopic, histopathological and serological findings in CD patients at AL-Anbar district, [west of Iraq]: A Comparative cross-sectional study

#### Hamad Majeed, Yasin; Ismael Ghazzay, Hazim; Jassim, Maher A

Correlation of IDA severity with endoscopic, histopathological and serological findings in CD patients at AL-Anbar district, [west of Iraq]: A Comparative cross-sectional study

Revista Latinoamericana de Hipertensión, vol. 15, núm. 4, 2020

Sociedad Latinoamericana de Hipertensión, Venezuela

Disponible en: https://www.redalyc.org/articulo.oa?id=170269715003

**DOI:** https://doi.org/10.5281/zenodo.4441343

Queda prohibida la reproducción total o parcial de todo el material contenido en la revista sin el consentimiento por escrito del editor en jefe. Copias de los artículos: Todo pedido de separatas deberá ser gestionado directamente con el editor en jefe, quien gestionará dicha solicitud ante la editorial encargada de la publicación



Esta obra está bajo una Licencia Creative Commons Atribución-SinDerivar 4.0 Internacional.



Artículos

# Correlation of IDA severity with endoscopic, histopathological and serological findings in CD patients at AL-Anbar district, [west of Iraq]: A Comparative cross-sectional study

Correlación de la gravedad de la IDA con hallazgos endoscópicos, histopatológicos y serológicos en pacientes con EC en el distrito de AL-Anbar, [oeste de Irak]: una comparación estudio transversal

Yasin Hamad Majeed University of Anbar, Arabia Saudita Yasin\_hamad@yahoo.com

https://orcid.org/0000-0002-9574-3258

Hazim Ismael Ghazzay University of Anbar,, Arabia Saudita

https://orcid.org/0000-0001-9937-3349

Maher A Jassim University of Anbar, Arabia Saudita

https://orcid.org/0000-0003-4932-1078

DOI: https://doi.org/10.5281/zenodo.4441343 Redalyc: https://www.redalyc.org/articulo.oa? id=170269715003

> Recepción: 12 Agosto 2020 Aprobación: 15 Septiembre 2020 Publicación: 20 Octubre 2020

## **ABSTRACT:**

Celiac disease (CD) is an autoimmune disorder described as a chronic inflammatory reaction due to gluten ingestion, which induces a particular degree of villous atrophy & possible -nutrient -malabsorption. Multi etiological factor of Anemia in CD patients; at the same time, iron deficiency anemia (IDA) may also be the most frequently recorded characteristic in children and adults with CD perhaps it is a unique finding existing. No previous study regarding the correlation of endoscopic, histopathological, and serological features of CD patients with IDA at Al-Anbar province. To find the correlation between the severity of IDA with endoscopic, histopathological, and serological features of CD patients. Hospital-based comparatives crosssectional study was conducted in the Gastroenterology Clinic of AL-Ramadi teaching hospital attached to Anbar Medical College, Ramadi, Iraq from August 2017 through June. 2020. A total of 108 consecutive new CD cases with iron-deficiency anemia, 70 of them were females, and 38 were males. CBC, blood films, ferritin level, endoscopic findings had been registered for all CD patients. In every CD case, gastro¬intestinal biopsies in addition to lgA (AntitTG levels) had been recorded. Histo¬pathology outcomes had been separated through the use of "taxonomy of modified¬ Marsh". A proper suitable statistical analysis had observed results. A total of 108 CD patients with iron deficiency anemia the mean age was 28.86 with a range from 15-66 years; the female was 64.8% and male 35.2%. There was a weak negative significant correlation Corr. Coef (r) (R=-0.003, p=0.000& R=-0.0180.161, p=0.000) amongst endoscopic and histopathological findings with IDA severity (HB gr%) respectively of CD patients using spearman's correlation coefficients (r). Negatively non-significant correlation of Anti- tTG level with IDA severity of CD (HB gr%) patients as detected through the use of Pearson's correlation (R=-149; P values 0.123). A weak negative significant correlation was found between the severity of IDA with endoscopic, histopathological finding, and but there was a negatively non-significant correlation between anti- tTG IgA level and IDA severity in CD.

KEYWORDS: Iron deficiency anemia, celiac disease, Endoscopy, histopathology, lgA anti tTG, Cross-sectional study, Ramadi district, Iraq.

#### RESUMEN:

La enfermedad celíaca (EC) es un trastorno autoinmune descrito como una reacción inflamatoria crónica debida a la ingestión de gluten, que induce un grado particular de atrofia vellositaria y posible malabsorción de nutrientes. Factor etiológico múltiple de la anemia en pacientes con EC; Al mismo tiempo, la anemia ferropénica (AIF) también puede ser la característica más frecuentemente registrada en niños y adultos con EC. Quizás sea un hallazgo único existente. No hay ningún estudio previo sobre la correlación



de las características endoscópicas, histopatológicas y serológicas de los pacientes con EC con AF en la provincia de Al-Anbar. Encontrar la correlación entre la gravedad de la AF con las características endoscópicas, histopatológicas y serológicas de los pacientes con EC. Se realizó un estudio transversal comparativo basado en hospitales en la Clínica de Gastroenterología del hospital universitario AL-Ramadi adjunto al Colegio Médico de Anbar, Ramadi, Irak, desde agosto de 2017 hasta junio. 2020. Un total de 108 nuevos casos consecutivos de EC con anemia ferropénica, 70 de ellos mujeres y 38 hombres. Se había registrado hemograma completo, frotis de sangre, nivel de ferritina y hallazgos endoscópicos para todos los pacientes con EC. En todos los casos de EC, se registraron biopsias gastrointestinales además de IgA (niveles de AntitTG). Los resultados histopatológicos se habían separado mediante el uso de "taxonomía de pantanos modificados". Los resultados se habían observado mediante un análisis estadístico adecuado. De un total de 108 pacientes con EC con anemia ferropénica, la edad media fue de 28,86 años con un rango de 15 a 66 años, la mujer 64,8% y el hombre 35,2%. Hubo una correlación significativa negativa débil Corr. Coef (r) (R=-0.003, p=0.000 & R=-0.0180.161, p=0.000) entre los hallazgos endoscópicos e histopatológicos con severidad IDA (HB gr%) respectivamente de pacientes con EC usando los coeficientes de correlación de Spearman (r). Correlación negativamente no significativa del nivel de Anti-tTG con la gravedad de la IDA de los pacientes con EC (HB gr%) detectada mediante el uso de la correlación de Pearson (R=-149; valores de P 0,123). Se encontró una correlación negativa significativa débil entre la gravedad de la IDA con el hallazgo endoscópico e histopatológico, pero hubo una correlación negativa no significativa entre el nivel de IgA anti-tTG y la gravedad de la IDA en la EC.

PALABRAS CLAVE: Anemia por deficiencia de hierro, enfermedad celíaca, endoscopia, histopatología, lgA anti tTG, estudio transversal, distrito de Ramadi, Irak.

#### Introduction:

Celiac disease (CD) is an autoimmune disorder described as a chronic inflammatory reaction due to gluten ingestion, which induces a particular degree of villi atrophy & possible -nutrient -malabsorption  $^1$ .

Its identification depending on the existence of duodenal villous  $\neg$ atrophy that appear histologically, in addition to +ve ABs against specific target antigens, especially tissue "transglutaminase" with "gliadin" or endomysium and also biopsies examination  $^2$ . Villous  $\neg$ atrophy findings on superior Endoscopy may appear for example "scalloped folds" or some time a mosaic pattern  $^3$ . It has a wide variety of clinically silent cases from overt malabsorption or standard gastrointestinal symptoms  $^4$ . IDA is a predominant sign of CD extra intestinally & was defined as the sole- sign of the CD disease  $^{5, 6}$ .

Multi etiological factor of Anemia in CD patients; at the same time, iron deficiency anemia (IDA) may also be the most frequently recorded characteristic in children and adults with CD perhaps it is a unique finding existing  $^7$ . Stimulatingly, in the CD, IDA is not only linked to damage of gluten driven in the intestinal mucosa, because IDA has also been described in serology positive CD patients earlier the duodenal atrophy progress  $^8$ .

This confirms that there is a need for IDA testing in patients with CD, with the recommendation and in the early stages to give a gluten-free diet to these patients who display the so-called "celiac trait" (with manifestations outside the intestine)  $^9$ .

IDA might also be the most frequently recorded characteristic in children & adults with CD; perhaps it is unique finding existing  $^{10-12}$ .

The chief mechanism for IDA in CD is linked to malabsorption because the iron absorption location in the proximal duodenum is virtually permanently involved  $^{13}$ .

No previous study regarding the correlation of endoscopic, histopathological, and serological features of CD patients with IDA at Al-Anbar province.



# PATIENTS AND METHODS:

Of 108 consecutive fresh patients aged over 15 years identified in the Gastroenterology Clinical department in ALRamadi teaching hospital during a period from August 2017 to June 2020 had been involved through using a retrospective cross-sectional epidemiological study. Seventy of them were females & other males. ELISA registered CBC, blood films, ferritin levels, lgA Anti tTG levels.

During the present study, gastrointestinal tract bleeding with patients with thalassemia had been excluded. Basically, IgA anti tTG <  $10 \, IU/ml$  measured as a negative, whereas IgA anti tTG 10.1- $20 \, IU/ml$  measured as suspicious even in the clinical- symptoms nonappearance CD patients. Whoever, IgA anti tTG >  $20 \, IU/ml$  ml were considered measured as a positive as described by the manufacturer & it had a calibration- a variety of  $3 \, to \, 100 \, IU/ml$ .

Endoscopy of the upper gastrointestinal tract with four duodenum biopsies from D2 was carried out for all patients with the estimate of endoscopic markers of the duodenum that include, "reduction in duodenal folds", scalloping of duodenal folds, the presence of mucosal fissures, and nodularity of the mucosa. Results of histopathology were categorized according to the "modified¬Marsh arrangement" <sup>14, 15</sup>.

Two expert pathologists have observed the specimen. Gender had been identified as a CD using "Gastroenterology ¬WHO¬ criteria" <sup>16</sup>. CD identification frequently depending on "serological EMA¬IgA¬antibodies" as well as "anti-tTG antibodies" positivity<sup>17</sup>, &/or <sup>18</sup>, plus anti¬tTG specific Abs targeting intestinal- mucosa histopathological antigens <sup>19</sup>.

Severity of IDA was classified according to Hb gr% level in non-pregnant female (15 years of age and above) [normal >=12 gr% mild anemia 11-10.9gr%, moderate 10.9-8 gr%, sever <8gr%.and in male (15 years of age and above) normal Hb >=13gr%, mild 12.9-11 gr%, moderate 10.9-8gr%, sever <8 gr%<sup>20,21</sup>.

CD patients with missing medical data have been excluded from the current study. Family recorded consent was taken from a total of patients. Ethical approval of the current study was taken from the Anbar medical faculty ethics approval Committee, Iraq.

**Statistical- Examination:** Statistical- investigation of our data was through using- the SPSS/software 24.0. Correlation of endoscopic, histopathological, and serological features of CD patients with IDA was calculated according to Pearson and spearman correlation. The variance in the concordance output was verified using the chi-square test.

## RESULTS:

Of 108 CD examinations patients during a year of study, 35.2% of them were male, and 64.8% were females giving a male/female 1:1.8. The range of age was from 15 years to 66 years. The mean age was (28.86±12.597 years). Most of the patients with CD presented in Hb% group 8-10.9 (48; 44.4%) followed by 11 and over (45;13.9%), while the remaining 15 [13.9%], was with Hb < 8gr shown in Table 1. A statistically significant between- different Hb % groups of CD patients (P-Value 0.000) (Table 1).

Regarding the Endoscopy, the study showed that 23 (21.3), 58 (53.7), 4 (3.7) and 1 (0.9) of CD patients were with a reduction in duodenal folds, scalloping, mucosal fissures, and atrophy with visible vessels respectively while the other 22 (20.4) of CD patients were normal Endoscopy. There was statistically significant between different endoscopic features of CD patients (P-Value 0.000) (Table 1)

Regarding the histopathological features, the study showed that 41(38.0%), 33(30.6%), 26(24.1%), 5 (4.6%) and 1 (0.9%) of CD patients were with G3A, G3B, G3C, G2, and G1respectively while the other 2(1.9%) of CD patients were normal histopathological features. There was a statistically significant between different histopathological features of CD patients (P-Value 0.000) (Table 1).

Table 1. General characteristic of endoscopic, histopathology and Hb % groups findings of CD patients



Negatively correlation (Corr. Coef (r)) of endoscopic finding with IDA severity of CD (HB gr %) of celiac disease patients as measured by using a spearman's correlation (R-0.003; P values 0.000) (Table 2, Figure, 1B). There was a weak correlation of endoscopic finding with the IDA severity of CD (HB gr %) the determined the grade of anemia of celiac disease patients as measured by using "spearman's correlation" (R 0.014; p-values 0.000) as shown in Table 2 and Figure 1

No.(%)	Hb gr%	No.(%)	Histopathology	No.(%)	Endoscopic finding	
15(13.9%)	< 8gr	41(38.0%)	GBA	23 (21.3)	Reduction in duodenal folds	
48(44.4%)	8-10.9	33(30.6%)	G3B	58 (53.7)	Scalloping	
45(41.7%).	>=11	26 (24.1%)	G3C	22 (20.4)	Normal	
		5 (4.6%)	G2	4 (3.7)	Mucosal fissures	
		1 (0.9%)	G1	1 (0.9)	Atrophy with visible vessels	
		2 (1.9%)	Normal			
	p.value=0.000		p.value=0.000		p.value 0.000	

Table 2: Correlation between Hb count (gr%) and endoscopical finding in CD patients (n = 108)

TABLE 2

Correlation between Hb count gr% and endoscopical finding in CD patients n 108

Spearman's Correlation				Mean of HB gr % (No. of patients)			Endoscopic finding
Endoscopic finding with HB gr%	Endoscopic finding with HB gr% group	P-value	Mean of Hb	> 11	8-10.9	<8	HB gr% group
-0.003	0.014	0.000	9.874 (23)	10.773 (11)	9.644%	7.267% (3)	Reduction in duodenal folds
			9.505 (58)	10.805% (20)	9.255% (29)	7.422% (9)	Scalloping
			10.182 (22)	10.779% (14)	9.671% (7)	5.400% (1)	Normal
			8.525 (4)	0.0% (0)	8.867% (3)	7.500% (1)	Mucosal fissures
			7.300 (1)	0.0% (0)	0.0% (0)	8.000% (1)	Atrophy with visible vessels
		9.671	·	10.789%	9.365%	7.300%	Total



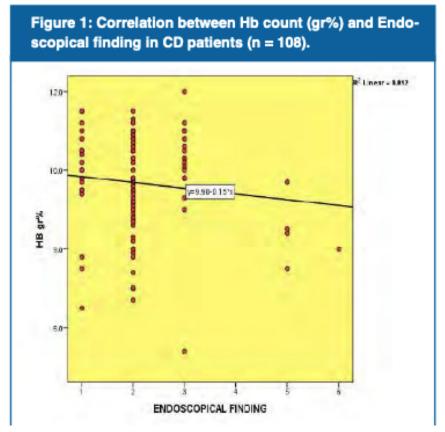


FIGURE 1: Correlation between Hb count (gr%) and Endoscopical finding in CD patients (n = 108).

Negatively Correlation (Corr. Coef (r)) of histopathological finding with IDA severity (HB gr %) of CD patients as measured by using "spearman's correlation" (R-0.018; p values 0.000) (Table 3, Figure, 2). There was a weak correlation of histopathological findings with the IDA severity (HB gr %) of CD that determined the grade of anemia of celiac disease patients as measured by using "spearman's Correlation" (R 0.030; P values 0.000) as described in Table 2, Figure 1.

Table 3: Correlation between Hb count (gr%) and Histopathology in CD patients (n = 108)

TABLE 3
Correlation between Hb count gr% and Histopathology in CD patients n 108

Spearman's Correlation				Mean of HB			
histopathology with Hb gr%	histopathology with Hb gr% group	P-value	Mean Hb	Hb >11gr%	Hb 8-10.9 gr%	Hb<8 gr	Hb gr% group
-0.018	0.030		9.644	10.65%	9.375%	6.675%	GBA
		0.000	9.767	10.81%	9.527	7.40 %	G3B
			9.227	10.820%	9.213%	7.68%	G3C
			11.020	11.02%	0.00%	0.0%	G2
			11.000	11.00%	0.0%	0.0%	G1
			10.400	11.000%	9.800%	0.0%	Normal
			9.671	10.789%	9.365%	7.30%	Total



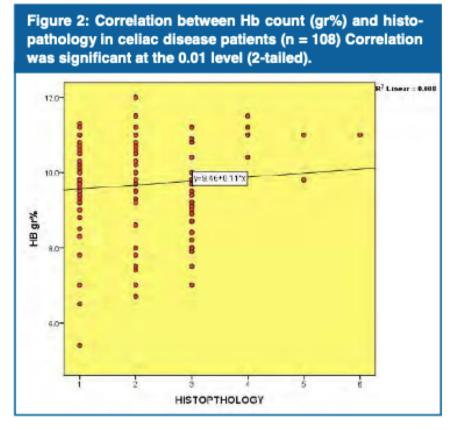


FIGURE 2: Correlation between Hb count (gr%) and histopathology in celiac disease patients (n = 108) Correlation was significant at the 0.01 level (2-tailed).

Negatively Correlation, (Corr. Coef (r)) of Anti- tTG level with IDA severity (HB gr %) of CD patients as measured by using a "Pearson's correlation" (R-.149-; p values 0.123) as described in Table 3, Figure 2. **Table 4:** Correlation between Hb count (gr%) with anti-tTG U/ml in celiac disease patients (n = 108)

TABLE 4
Correlation between Hb count gr% with anti tTG Uml in celiac disease patients n 108

Pearson Correlation	P-value	Mean Hb	Hb gr% >11	Hb gr% 8-10.9	Hb gr% <8	Anti- tTG level
149-	0.123	9.713	40 (43.0%)	41 (44.1%)	12 (12.9%)	10.1-20
		9.413	5 (33.3%)	7 (46.7%)	3 (20.0%)	>20
			45 (41.7%)	48 (44.4%)	15 (13.9%)	Total



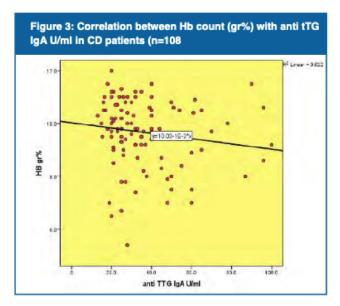


FIGURE 3: Correlation between Hb count (gr%) with anti tTG IgA U/ml in CD patients (n=108  $\,$ 

#### **Discussion:**

Our study revealed female predominance which was identical to the US, Europe, & Middle East as overall showed a female –predominance of CD patients  $^{22-24}$ , the causes are unknown but might due to the autoimmune-mediated illnesses generally more predominant in females  $^{25}$ .

The study showed a weak negative correlation between endoscopic finding and anemia severity this finding is consistent with the research done by Mauri no et al.  $^{26}$  and Kalhan S, et al.  $^{27}$  and inconsistent with the study done by Oxentenko et al.  $^{28}$ .

20.4% of our CD patients with the iron-deficiency anemia had routine Endoscopy. "Endoscopic markers" nonappearance of villous atrophy doesn't reject the identification & duodenal biopsies must be complete in cases of clinical & o, CD laboratory suspicion <sup>29, 30</sup>.

Our study showed a weak negative correlation with the severity of anemia and duodenal histopathology this finding is consistent with Zamani et al.  $^{31}$  and inconsistent with Ganji et al.  $^{32}$ .

Our study revealed 2 cases of positive serology and normal Endoscopy and histopathology; these are considered as latent celiac <sup>33</sup>.

Our study revealed that there is a negative non-significant correlation between anti-tTG lgA level and severity of anemia, this result was inconsistent with the result reported by Ertekin et al. who identified that Hb levels of CD patients negatively correlated with tTG titers <sup>34</sup>.

#### Conclusion:

Our study identified a weak negative significant correlation was registered between the severity of IDA with endoscopic, histopathological findings. A negatively non-significant correlation between anti- tTG IgA level and IDA severity in CD



# REFERENCES

- 1.Ludvigsson JF, Leffler DA, Bai JC, Biagi F, Fasano A, Green PH, Hadjivassiliou M, Kaukinen K, Kelly CP, Leonard JN, Lundin KE. The Oslo definitions for coeliac disease and related terms. Gut. 2013 Jan 1;62(1):43-52.
- 2. Vives-Pi M, Takasawa S, Pujol-Autonell I, Planas R, Cabre E, Ojanguren I, Montraveta M, Santos AL, Ruiz-Ortiz E. Biomarkers for diagnosis and monitoring of celiac disease. Journal of clinical gastroenterology. 2013 Apr 1;47(4):308-13.
- 3. Mauriño E, Capizzano H, Niveloni S, Kogan Z, Valero J, Boerr L, Bai JC. Value of endoscopic markers in celiac disease. Digestive diseases and sciences. 1993 Nov 1;38(11):2028-33.
- 4.Chellat H, Salihoun M, Kabbaj N, Amrani L, Serraj I, Chaoui Z, Acherki M, Amrani N. Diagnostic yield of routine duodenal biopsies in iron deficiency anemia for celiac disease diagnosis. International Scholarly Research Notices. 2013;2013.
- 5.Corazza GR, Valentini RA, Andreani ML, D'anchino M, Leva MT, Ginaldi L, De Feudis L, Quaglino D, Gasbarrini G. Subclinical coeliac disease is a frequent cause of iron-deficiency anaemia. Scandinavian journal of gastroenterology. 1995 Jan 1;30(2):153-6.
- 6.Schmitz U, Ko Y, Seewald S, Düsing R, Vetter H. Iron-deficiency anemia as the sole manifestation of celiac disease. The clinical investigator. 1994 Jul 1;72(7):519-21.
- 7. Berry N, Basha J, Varma N, Varma S, Prasad KK, Vaiphei K, Dhaka N, Sinha SK, Kochhar R. Anemia in celiac disease is multifactorial in etiology: A prospective study from India. JGH Open. 2018 Oct;2(5):196-200.
- 8.Repo M, Lindfors K, Mäki M, Huhtala H, Laurila K, Lähdeaho ML, Saavalainen P, Kaukinen K, Kurppa K. Anemia and iron deficiency in children with potential celiac disease. Journal of pediatric gastroenterology and nutrition. 2017 Jan 1;64(1):56-62.
- 9.Popp A, Mäki M. Gluten-Induced Extra-Intestinal Manifestations in Potential Celiac Disease—Celiac Trait. Nutrients. 2019 Feb;11(2):320.
- 10.Freeman HJ. Iron deficiency anemia in celiac disease. World Journal of Gastroenterology: WJG. 2015 Aug 21;21(31):9233.
- 11.Unsworth DJ, Lock FJ, Harvey RF. Iron-deficiency anemia in premenopausal women. The Lancet. 1999 Mar 27;353(9158):1100.
- 12.Bottaro G, Cataldo F, Rotolo N, Spina M, Corazza GR. The clinical pattern of subclinical/silent celiac disease: an analysis of 1026 consecutive cases. The American journal of gastroenterology. 1999 Mar 1;94(3):691-6.
- 13.Baydoun A, Maakaron JE, Halawi H, Abou Rahal J, Taher AT. Hematological manifestations of celiac disease. Scandinavian journal of gastroenterology. 2012 Dec 1;47(12):1401-11.
- 14.Marsh MN. Gluten, major histocompatibility complex, and the small intestine: a molecular and immunologic approach to the spectrum of gluten sensitivity ('celiac sprue'). Gastroenterology. 1992 Jan 1;102(1):330-54.
- 15.Oberhuber G, Granditsch G, Vogelsang H. The histopathology of coeliac disease: time for a standardized report scheme for pathologists. European journal of gastroenterology & hepatology. 1999 Oct 1;11(10):1185.
- 16.Fasano A. Novel therapeutic/integrative approaches for celiac disease and dermatitis herpetiformis. Clinical and Developmental Immunology. 2012 Oct;2012.
- 17.Bonatto MW, Kotze L, Orlandoski M, Tsuchiya R, de Carvalho CA, Lima D, Kurachi G, Orso IR, Kotze L. Endoscopic evaluation of celiac disease severity and its correlation with histopathological aspects of the duodenal mucosa. Endoscopy international open. 2016 Jul;4(7): E767.
- 18. Volta U, Molinaro N, De Franceschi L, Fratangelo D, Bianchi FB. IgA anti-endomysial antibodies on human umbilical cord tissue for celiac disease screening. Digestive diseases and sciences. 1995 Sep 1;40(9):1902-5.
- 19.Dieterich W, Ehnis T, Bauer M, Donner P, Volta U, Riecken EO, Schuppan D. Identification of tissue transglutaminase as the autoantigen of celiac disease. Nature medicine. 1997 Jul;3(7):797-801.
- 20. World Health Organization. World declaration and plan of action for nutrition. Food and Agriculture Organization of the United Nations; 1992.



- 21.WHO U. UNU. Iron deficiency anemia: assessment, prevention, and control. A guide for program managers, Geneva, WHO. 2001.
- 22.Green PH, Stavropoulos SN, Panagi SG, Goldstein SL, McMahon DJ, Absan H, Neugut AI. Characteristics of adult celiac disease in the USA: results of a national survey. The American journal of gastroenterology. 2001 Jan 1;96(1):126-31.
- 23. Ciacci C, Cirillo M, Sollazzo R, Savino G, Sabbatini F, Mazzacca G. Gender and clinical presentation in adult celiac disease. Scandinavian journal of gastroenterology. 1995 Jan 1;30(11):1077-81.
- 24.Nejad¹ MR, Rostami K, Pourhoseingholi¹ MA, Mojarad EN. Atypical presentation is dominant and typical for coeliac. J Gastrointestin Liver Dis. 2009 Sep;18(3):285-91.
- 25. Tajuddin T, Razif S, Thorne J, Murray FE. Clinical presentation of adult coeliac disease.
- 26. Mauriño E, Capizzano H, Niveloni S, Kogan Z, Valero J, Boerr L, Bai JC. Value of endoscopic markers in celiac disease. Digestive diseases and sciences. 1993 Nov 1;38(11):2028-33.
- 27.Kalhan S, Joseph P, Sharma S, Dubey S, Dudani S, Dixit M. Comparative study of histopathological Marsh grading with clinical and serological parameters in the celiac iceberg of north India. Indian Journal of Pathology and Microbiology. 2011 Apr 1;54(2):279.
- 28.Oxentenko AS, Grisolano SW, Murray JA, Burgart LJ, Dierkhising RA, Alexander JA. The insensitivity of endoscopic markers in celiac disease. The American journal of gastroenterology. 2002 Apr 1;97(4):933-8.
- 29. Cammarota G, Fedeli P, Gasbarrini A. Emerging technologies in upper gastrointestinal Endoscopy and celiac disease. Nature Clinical Practice Gastroenterology & Hepatology. 2009 Jan;6(1):47-56.
- 30.Dickey W. Endoscopic markers for celiac disease. Nature clinical practice Gastroenterology & hepatology. 2006 Oct;3(10):546-51.
- 31.Zamani F, Mohamadnejad M, Shakeri R, Amiri A, Najafi S, Alimohamadi SM, Tavangar SM, Ghavamzadeh A, Malekzadeh R. Gluten-sensitive enteropathy in patients with iron deficiency anemia of unknown origin. World Journal of Gastroenterology: WJG. 2008 Dec 28;14(48):7381.
- 32. Ganji A, Esmaielzadeh A, Aghayee MA, Goshayeshi L, Ghaffarzadegan K. The clinical presentation of celiac disease: experiences from northeastern Iran. Middle East Journal of digestive diseases. 2014 Apr;6(2):93.
- 33. James SP. National Institutes of Health consensus development conference statement on celiac disease, June 28–30, 2004.
- 34.Freeman HJ. Iron deficiency anemia in celiac disease. World Journal of Gastroenterology: WJG. 2015 Aug 21;21(31):9233.

