

# Enamel defects and malocclusion in patients with celiac disease

*Sahar F. Abdul- Wahid B.D.S. .<sup>(1)</sup>*

*Lamia A. Al-Azawi B.D.S. M.Sc Ph.D. .<sup>(2)</sup>*

*Iman T. Ibrahim B.D.S M.Sc.<sup>(2)</sup>*

## ABSTRACT

**Background:** This study investigated the types and severity of enamel defects and malocclusion in celiac patients compared to matching controls.

**Materials and methods:** One hundred and two celiac patients were included in the study they were attending the Gastroenterology and Hepatology Teaching Hospital.

**Results:** Enamel defects in the study group showed a highly significant difference than found in the control group in both deciduous and permanent teeth. Concerning malocclusion study group, a higher percentage of severe anomalies (code 2) than control group the difference was statistically highly significant.

**Keywords:** Enamel defects, celiac disease, malabsorption (J Bagh Coll Dentistry 2005; 17(3): 98- 100)

## INTRODUCTION

Celiac disease (CD) also termed celiac sprue, and gluten sensitivity enteropathy is characterized by small intestinal malabsorption of nutrients after ingestion of wheat gluten or related proteins from rye and barley prompt clinical and histologic improvement after adherence to strict gluten free diet and clinical relapse when gluten is reintroduced.<sup>(1,2)</sup>

Enamel hypoplasia has been reported in relation to various gastro-intestinal disturbances of infancy and childhood.<sup>(3)</sup>

Because permanent tooth enamel defects strongly associated with CD both in children and adults, the study was conducted in order to find whether these defects could be used to screen for CD among healthy persons with mild or abdominal symptoms, however we have controversial findings from review of studies relevant to this study, prevalence of malocclusion was not reported in CD. This study showed that there was a significant difference between study and control groups.

## MATERIALS AND METHODS

The study included one hundred and two patients who attended the Gastroenterology and Hepatology Teaching Hospital compared to control group matching in age and sex. Age range 2-35 years old, these patients were diagnosed as having CD by GIT specialist and depending on clinical and histopathologic findings,

clinical examination included, questionnaire and oral examination which included

**Enamel opacities / hypoplasia** which were recorded according to Modified Developmental Defects of Enamel Index (DDE), (WHO1997).<sup>(4)</sup>

**Malocclusion** The clinical examination starts with overall look at the subject general appearance, symmetry. The examination should then proceed with assessment of malocclusion. Malocclusion was diagnosed according to the criteria of (WHO, 1987).<sup>(5)</sup>

## RESULTS

The results showed that normal enamel in deciduous teeth in study group was lower in percentage than control group (60.36%, 69.29).

Table (1) demonstrated that the percentage of enamel defects in deciduous teeth was higher in study group (71.43%) than that in control group (32.14%) the difference was statistically highly significant ( $z=2.94, p<0.01$ ).

Enamel defects in permanent teeth were demonstrated in table (2) the percentage of enamel defects in permanent teeth in study group was higher than in control group (89.66%, 29.99%) respectively and it was statistically significant ( $z= 8.03, p<0.001$ ).

Malocclusion: study group showed higher percentage of serious anomalies (code 2), 39.18% than that of control (12.16%) the difference was highly significant ( $z= 3.76, p<0.001$ ) the results are detailed in table (3).

(1) Assist. Lecturer, Depart. of Pedodontics and Preventive Dentistry, College of Dentistry, University of Baghdad

(2) Assist. Prof. Department of Pedodontics and Preventive Dentistry, College of Dentistry, University of Baghdad

## DISCUSSION

Different indices or classification had been used in examining the enamel defect in celiac disease in all previous studies compared to relevant studies<sup>(6,7)</sup>, no enamel defect were found in the deciduous teeth in children with CD, Smith and Miller described a case report of a 6 years old boy suffered from CD and had enamel hypoplasia in his deciduous molars, the controversy could be attributed also beside the difference in index system to the delay in diagnosis of the case as celiac condition.<sup>(3)</sup>

By comparing the results of this study to the study of Attavasso<sup>(7)</sup> who concluded that 78% of the study group had enamel defect and 96% in Aine<sup>(8)</sup> and several case reports studies which concluded that there is no statistical differences concerning enamel defect between celiac patients and control.<sup>(3,6,9-11)</sup>

In Merzec–koronczewska study,<sup>(12)</sup> occlusal abnormalities were noted in one third of 38 celiac children with mixed teeth and the dental age as delayed in relation to calendar age. Since children with CD often lost weight and have a recorded rate of growth their dental development can be expected to be related dietary treatment had on effect on both dental and skeletal maturity.<sup>(8)</sup>

## REFERENCES

1. Trier JS. Celiac sprue and refractory sprue in sleisenger and fordtrans gastrointestinal liver disease pathophysiology, diagnosis, management 6<sup>th</sup> St. Louis C.V Mosby, 1998: pp 1557- 73.
2. Farrel RJ, Kelly CD. Celiac sprue and refractory sprue in sleisenger fordtrans gastrointestinal and liver disease 7 th Ed. c.v Mosby 2002: 1817– 41.
3. Smith D, Miller J. Gastro-enteritis celiac disease and enamel hypoplasia. Br Dent J 1979; 147, 91- 5.
4. World Health Organization, Oral Health Surveys Basic Methods, 4<sup>th</sup> Ed, Geneva, Switzerland, 1997.
5. World Health Organization, Oral Health Surveys Basic Methods: 3<sup>rd</sup> Ed, Geneva, Switzerland, 1987.
6. Rasmussen P, Espelid I. Celiac disease and dental malformation. J Dent Child 1980; 47: 190-2.
7. Attanassov N, Targova S, Kovaceva J. Enamel hypoplasia in children with celiac disease. Stomatogüa 1983; 65: 77-81 (midline abstract).
8. Aine L. Dental enamel defects and dental maturity in children and adolescents with celiac disease. Proce Finn Dent Soc 1986; 83(suppl 13): 1-71.
9. Aine L. Permanent tooth dental enamel defects leading to the diagnosis of celiac disease. Br Dent J 1994; 177: 253-4.
10. Bertoldi C, Balldi F, Tanza D. Expermantation and clinical analysis of the interrelation.,Dermatology 1995; 200: 340.
11. Rassmusson CG, Eriksson MA. Celiac disease and mineralization disturbances of permanent teeth. Int J Pediat Dent 1993; 11(3): 179-83.
12. Merzec – Koronczewskaz MK. The condition of the stomatognathic system in children with gluten dependent celiac disease. Gzas Stomatol 1992; April 3(4): 207 -12.

**Table 1: Enamel defects in deciduous teeth in both study and control groups by age and gender**

Age	Gender	Study		Control		Sig
		No.	%	No	%	
2-5	M	5	71.43	1	14.29	S*
	F	7	87.50	3	37.50	S*
	T	12	80.00	4	26.67	S**
6-10	M	4	57.14	2	28.57	N.S
	F	4	66.67	3	50.00	N.S
	T	8	61.54	5	38.46	N.S
Total	M	9	64.29	3	21.43	S*
	F	11	78.57	6	42.86	N.S
	T	20	71.43	9	32.14	S**

**Table 2: Enamel defects in permanent teeth in both study and control groups by age and gender.**

Age	Gender	Study		Control		Sig
		No.	%	No.	%	N.S.
6-10	M	3	42.86	1	14.	N.S.
	F	4	66.67	3	50.00	N.S.
	T	7	53.85	4	30.77	S**
11-15	M	9	100.00	3	33.33	S***
	F	16	100.00	6	37.50	S***
	T	25	100.00	9	36.00	S***
16-21	M	8	100.00	1	12.50	S***
	F	5	100.00	0	.00	S**
	T	13	100.00	1	8.69	S***
22-30	M	6	85.71	1	14.29	S**
	F	12	100.00	5	41.67	S**
	T	18	94.74	6	31.58	S***
31-35	M	9	90.00	2	20.00	S**
	F	6	85.71	4	57.14	N.S.
	T	15	88.24	6	35.29	S**
Total	M	35	85.37	8	19.51	S***
	F	43	93.48	18	39.13	S***
	T	78	89.66	26	29.89	S***

**Table 3: Malocclusion in study and control groups according to age and gender.**

Age	Gender	Study						Control						Sig.		
		0		1		2		0		1		2		0	1	2
		No.	%	No.	%	No.	%	No.	%	No.	%	No.	%			
11-5	Male	2	22.2	3	33.3	4	44.4	5	55.5	4	44.4	0	0	N.S.	N.S.	S*
	Female	3	18.75	4	25.0	9	56.25	9	56.25	2	12.5	5	31.25	S*	N.S.	N.S.
	Total	5	20.0	7	28.0	13	52.0	14	56.0	6	24.0	5	20.0	S**	N.S.	S*
16-21	Male	1	12.5	1	12.5	6	75.0	4	50.0	3	37.5	1	12.5	N.S.	N.S.	S*
	Female	1	20.0	3	60.0	1	20.0	4	80.0	0	0	1	20.0	N.S.	N.S.	N.S.
	Total	2	15.4	4	53.8	7	53.8	8	61.5	3	23.1	2	15.4	S*	S*	S*
22-30	Male	3	42.8	2	28.6	2	28.6	4	57.1	3	42.9	0	0	N.S.	N.S.	N.S.
	Female	4	33.3	5	41.7	3	25.0	7	58.3	3	25.0	2	16.7	N.S.	N.S.	N.S.
	Total	7	36.8	7	36.8	5	26.3	11	57.9	6	31.6	2	10.5	N.S.	N.S.	N.S.
31-35	Male	5	40.0	3	30.0	3	30.0	6	60.0	4	40.0	0	0	N.S.	N.S.	N.S.
	Female	4	71.4	1	14.3	1	14.3	4	57.1	3	42.9	0	0	N.S.	N.S.	N.S.
	Total	9	52.9	4	23.5	4	23.5	10	58.2	7	41.2	0	0	N.S.	N.S.	S*
Total	Male	10	29.4	9	26.5	15	44.1	19	55.9	14	41.2	1	2.9	S*	N.S.	S***
	Female	13	32.5	13	32.5	14	35.0	24	60.0	8	20.0	8	20.0	S*	N.S.	N.S.
	Total	23	31.1	22	26.7	29	39.2	43	58.1	22	29.7	9	12.2	S***	N.S.	S***