

# EVALUATION OF BONE MINERAL DENSITY AND BONE METABOLİSM MARKERS İN CHİLDREN DİAGNOSED AS CELİAC DİSEASE

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## INTRODUCTION AND AIM

Metabolic bone disorders due to calsium and vitamin D deficiency are one of the most frequent extraintestinal symptoms in Celiac disease.

In this study it is aimed to evaluate bone mineral density in patients with Celiac disease during diagnose

and evaluate the factors related to bone mineral metabolism.

# MATERIAL AND METHOD

The study included 43 children (34 girl) diagnosed as Celiac disease. Mean age was 9.9±4.8 (2.5-17.7) years.

	Clinical	Calsiyum (Ca) Phosphor (P)	Lumbal (L1-L4) bone mineral density (BMD) levels measured via DEXA (Dual Energy X-Ray Absorptiometry) were evaluated	Z scores due to cronological age (Z-CA)	2015 Decemt
	Patological	Parathormon (PTH) 250H vitamin D levels		Z scores due to height age (Z-HA)	

# RESULTS

Table 1: Clinical features of the cases						
	Mean±SD	min-max				
Age (year)	9.9±4.8	2.5-17.7				
Weight (Kg)	29.9±14.6	11.4-65.5				
Weight SD	-1.1±1.1	-3.5-0.8				
Haight (cm)	120 0+27 0	96 5 179 3				

Table 1: Clinical features of the cases			Table 2: Laboratory features of cases			of cases				
	Mean±SD	min-ma×		Mean	±SD	min-max	There were no difference in BMD Z scores due to			
Age (year)	9.9±4.8	2.5-17.7	<b>Ca</b> (mg/dl <b>)</b>	9.7±	0.4	8.9-10.9	chronological and height ages (p=0.150, p=0.225, respectively).			
Weight (Kg)	29.9±14.6	11.4-65.5	P (mg/dl)	4.7±	0.6	2.7-6.0	There was positive correlation between BMD Z scores due to			
Weight SD	-1.1±1.1	-3.5-0.8	ALP (U/L) PTH	203.0±	100.1 21.5	100.0-466.0	chronological age and body weight, height and BMI Z scores (n<0.001 n=0.005 n=0.015 respectively)		Z scores	
Height (cm)	130.0±27.0	86.5-178.3	(pg/ml) 25 OH vit D	13.5	±7.7	4.6-35.1				
Height SD	-0.9±1.3	-4.2-1.9	(ng/ml) BMD	0.6±0.2 0.3-1.0   -0.8±1.1 -3.6-1.6   -0.2±1.1 -3.6-1.9		0.3-1.0	No relation	1 between BMD Z scores and plasma vitamin D, Ca. P. ALP and PTH levels (n>0.050)		
BMI (kg/m²)	16.4±2.1	12.9-22.4	(gr/cm <sup>2</sup> ) <b>Z -CA</b>			-3.6-1.6				
BMI SD	-0.8±1,1	-3.1-1.0	Z-HA			No relation between BMD Z scores and Celiac type and Marsh histopathologic stage (p>0.050)				
Stage of nuberty 1 1-5					BMD Z scores due to chronological age					
(median)	_				Marsh stage	Age group	<-2	-1 to -2	>-1	
0-6 years 30.2% (n=13)		Celiac typ Typical 16.7%	7pe Type-1 9.3% (n=4) % (n=7) Type-2 4 7% (n=2)		-1 9.3% (n=4) -2 4 7% (n=2)	0-6 years	0% (n=0)	53.8% (n=7)	46.2% (n=6)	
7-11 years 30.2% (n=13)		Atypical 64.3%	% (n=28) Type-3a 32.5%(n=14)		7-11 years	0% (n=0)	46.2% (n=6)	53.8% (n=7)		
>11 years 39.5% (n=1/)		Silent 19%	(n=8)	Type-:	3b 25.6%(n=11)	>11 years	35.3% (n=6)	11.8% (n=2)	52.9% (n=9)	
46.5% (n=20) pubertal				Type-	3c 27.9%(n=12)	Total	14 % (n=6)	34.9% (n=15)	51.2% (n=22)	

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#### DISCUSSION

Rickets, osteopenia, osteoporosis and bone deformities may develop due to impaired calcium and vitamin D absorption in Celiac patients. In the

literature, it is suggested to perform BMD measurement in case of newly diagnosed celiac disease.

It has been suggested that at the time of diagnosis the Marsh histopathologic stage can predict the formation of low BMD with a risk of developing osteoporosis. On the other hand, there are also studies suggesting that BMD of Celiac patients have no association with Marsh histopathologic stage at

the time of diagnosis and that there are no clinical and laboratory markers for low BMD in children with Celiac disease.

An important point about Celiac disease-related osteoporosis is that full improvement in BMD are possible in children by gluten-free diet therapy,

which is not true for adults. This requires early diagnosis and treatment of patients.

It has been reported in the literature that in children with Celiac disease, the BMD Z-scores decrease while age increases and that bone mineral loss increases proportionally with age.

## CONCLUSION

Diagnose in higher ages effects bone mineral density negatively in Celiac disease. Diagnose in early ages decreases bone mineral leak and decreases

morbidity in patients with osteopeni and osteoporosis via treatment posibilities.



Bone, growth plate and mineral metabolism





