

Bone mineral density in postmenopausal women with high levels of parathyroid hormone

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Objectives

- One of the main problems encountered with increased levels of serum PTH is decrease in bone mineral density in the long term.
- Decrease in bone mineral density is seen not only in hypercalcemic patients with primary hyperparathyroidism and also seen in normocalcemic patients with primary hyperparathyroidism.
- Secondary hyperparathyroidism due to vitamin D deficiency leads to mineralization disorders of the bones, low bone mineral density, osteoporosis and ultimately an increased risk of bone fracture in adults.
- In this study, we aimed to determine the reasons of secondary hyperparathyroidism for bone mineral density.

Methods

- One hundred and thirteen postmenopausal women with high serum PTH levels and 43 ones with normal PTH were enrolled to the study. Serum PTH level higher than 68.3 pg/ml was considered as hyperparathyroidism.
- Premenopausal women aged <50 years, postmenopausal women aged ≥ 75 years, women with parathyroid gland disease, chronic renal failure, malabsorption, biphosphonate, thiazid or corticosteroid drug use and acute infection were excluded.
- Bone mineral density measurements in all participants were detected by dual-energy x-ray absorptiometry (DEXA) method using with Hologic QDR-4500A S/N 45130 analyzer (Hologic Inc. Bedford, MA, USA). The measurements of the lumbar spine and the left femur were recorded. The mineral density (g/cm²) of the second lumbar (L) vertebra and the average mineral density of the L 1-4 vertebrae were compared statistically. Also the mineral density (g/cm²) of the left femoral neck and the average mineral density of the femoral neck, trochanteric and intertrochanteric regions were compared.

Results

The average age was 56.9 \pm 0.5 (50-74, n=113) in patients with hyperparathyroidism and 57.9 \pm 0.7 (50-72, n=43) in the control group. There were no significant differences between the groups in terms of menopausal age and body mass index (Table 1).

There was no significant difference between groups in terms of serum calcium levels but serum phosphorus levels were significantly lower in the study group. 24-hour urinary calcium excretion was reduced in the study group and 24-hour urinary phosphorus excretion was not different from the control group. Serum 25-OH vitamin D levels were low in both groups but there were no statistically significant differences between groups (Table 2).

The study group and the control group were compared in terms of bone mineral density (Table 3). The mineral density of the second lumbar vertebra and the average mineral density of the L 1-4 vertebrae were compared statistically. Also the mineral density of the left femoral neck and the average mineral density of the femoral neck, wards triangle and trochanteric region were compared. There was no significant difference in both regions in terms of bone mineral density. However, when all cases were evaluated together, the average mineral density of the L 1-4 vertebrae showed a reduction in parallel with the increase in serum PTH levels. It was statistically significant (r = -0.175; p = 0.029) but the average mineral density of the femoral neck, wards triangle and trochanteric region did not show any increase or decrease in parallel with the increase in serum PTH levels.

	Table-1: Some clinical and laboratory characteristics of the people							
		Study Group	Control Group	All Cas	es			
		(n=113)	(n=43)	(n=156	j)			
	Age (year)	56.9±0.5	57.9±0.7	57.2±0	.4			
	Menopausal age (year)	10.2±0.6	9.7±0.9	10.1±0	.5			
	BMI (kg/m²)	30.2±0.5	29.4±0.5	30.0±0	.3			
	Creatinine (mg/dl)	0.7±0.0	0.6±0.1	0.7±0.	0			
	Chlorine (mmol/l)	104.2±0.2	104.2±0.3	104.0±0	.2			
	ALP (IU/I)	83.6 ±2.3	81.3±3.1	83.0±1.	.8			
	Data are given as mean±SE (sta	ta are given as mean±SE (standart error). ALP: alkaline phosphates, BMI: body mass index						
	Table-3: Comparison of the stu	3: Comparison of the study group and the control group in terms of bone mineral density						
		Study Group (n=113) Control C	Froup (n=43)	P			
	L2 (g/cm²)	0.90 ± 0.12	0.89	± 0.12	0.84			
	L1-L4 total (g/cm²)	0.90 ± 0.12	0.92	± 0.02	0.36			
	Femoral neck (g/cm²)	0.75 ± 0.11	0.75	± 0.01	0.68			
	Femur total (g/cm²)	0.95 ± 0.01	0.95	± 0.01	0.97			
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	Study Group (n=113)	Control Group (n=43)	P
Calcium (mg/dl)	9.5 ± 0.04	9.5 ± 0.05	0.588
Adjusted calcium (mg/dl)	9.6 ± 0.04	9.6 ± 0.05	0.355
Phosphorus (mg/dl)	3.3 ± 0.04	3.6 ± 0.7	<0.001
Creatinine clearance (ml/min)	110.6 ± 2.2	108 ± 3.5	0.53
24hUCa (mg/24s)	132.3 ± 7.3	168.5 ± 11.8	0.006
24hUp (mg /24s)	637.6 ± 23.4	653.4 ±42.7	0.956
TRp (%)	86.0 ± 0.4	86.7 ± 0.6	0.478
25- OH vitamin D (μg/l)	10.7 ± 0.6	12.1 ± 1.3	0.439
Uca /Ucr	0.009 ± 0.004	0.012 ± 0.004	0.001

Conclusions

- Decrease in bone mineral density was seen not only in patients with primary hyperparathyroidism and also seen in patients with secondary hyperparathyroidism due to vitamin D deficiency. Nakamura et al. (12) showed a significant decrease in bone mineral density in patients with hyperparathyroidism but they also showed a decrease in bone mineral density in patients with vitamin D deficiency independently from PTH levels. There are no adequate data about what levels of increase in serum PTH may adversely affect bone mineral density (13). Albettazzi et al. (5) compared the cases with elevated serum PTH levels and with normal serum PTH levels in their study but they could not detect a significant relationship between serum PTH levels and the bone mineral density of the lumbar spine and the femoral neck. Similarly, Kaya et al. (14) found no association between bone mineral density and elevated serum PTH levels in their study. But Silverberg et al. (15) demonstrated slightly decrease in bone mineral density with elevated PTH levels in their study. In our study, there is no significant difference between the study and control group in terms of bone mineral density and it may be related to the duration of hyperparathyroidism.
- Our study was a cross-sectional study and we did not know how long the patients had hyperparathyroidism. Probably, due to short-term exposure of elevated PTH levels and the small number of the study group, a significant difference could not be identified between the two groups in terms of bone mineral density. In addition, the levels of vitamin D deficiency in both groups were not very different. Therefore, it is considered that the levels of secondary hyperparathyroidism, the main factor that causes bone resorption, were not at the levels of negative effect on bone mineral density.

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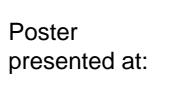
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