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Evidence for Alteration of the Vitamin D-Endocrine System in Obese Subjects

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Abstract

Serum immunoreactive parathyroid hormone (PTH) is increased in obese as compared with nonobese subjects and declines with weight loss. To determine whether alteration of the vitamin D-endocrine system occurs in obesity and whether ensuing secondary hyperparathyroidism is associated with a reduction in urinary calcium, a study was performed in 12 obese white individuals, five men and seven women, and 14 nonobese white subjects, eight men and six women, ranging in age from 20 to 35 yr. Body weight averaged 106 ± 6 kg in the obese and 68 ± 2 kg in the nonobese subjects ($P < 0.01$). Each of them were hospitalized on a metabolic ward and were given a constant daily diet containing 400 mg of calcium and 900 mg of phosphorus. Whereas mean serum calcium, serum ionized calcium, and serum phosphorus were the same in the two groups, mean serum immunoreactive PTH (518 ± 48 vs. 243 ± 33 pg/ml, $P < 0.001$), mean serum 1,25-dihydroxyvitamin D [$1,25(\text{OH})_2\text{D}$] (37 ± 2 vs. 29 ± 2 , $P < 0.01$), and mean serum Gla protein (33 ± 2 vs. 24 ± 2 ng/ml, $P < 0.02$) were significantly higher, and mean serum 25-hydroxyvitamin D (25-OHD) (8 ± 1 vs. 20 ± 2 ng/ml, $P < 0.001$) was significantly lower in the obese than in the nonobese men and women. Mean urinary phosphorus was the same in the two groups, whereas mean urinary calcium (115 ± 10 vs. 166 ± 13 mg/d, $P < 0.01$) was significantly lower, and mean urinary cyclic AMP (3.18 ± 0.43 vs. 1.84 ± 0.25 nM/dl GF, $P < 0.01$) and creatinine clearance (216 ± 13 vs. 173 ± 6 liter/d, $P < 0.01$) were significantly higher in the obese than in the nonobese individuals. There was a significant positive correlation between percentage of ideal body weight and urinary cyclic AMP ($r = 0.524$, $P < 0.01$) and between percentage of ideal body weight and serum immunoreactive PTH ($r = 0.717$, $P < 0.01$) in the two groups. The results provide evidence that alteration of the vitamin D-endocrine system in obese subjects is characterized by secondary hyperparathyroidism which is associated with enhanced renal tubular reabsorption of calcium and increased circulating $1,25(\text{OH})_2\text{D}$. The reduction of serum 25-OHD in them is attributed to feedback inhibition of hepatic synthesis of the precursor by the increased serum $1,25(\text{OH})_2\text{D}$.

Introduction

Available evidence indicates that serum immunoreactive parathyroid hormone (PTH)¹ is higher in obese than in nonobese

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young adults and declines with weight loss (1). In view of these observations, we carried out an investigation to determine whether obesity modifies the vitamin D-endocrine system and whether secondary hyperparathyroidism is associated with a reduction of urinary calcium in obese subjects.

Methods

26 normal white subjects were studied. There were 12 obese individuals (five men and seven women) and 14 nonobese subjects (eight men and six women) ranging in age from 20 to 35 yr. All of them were hospitalized on the General Clinical Research Center of the Medical University of South Carolina, Charleston, SC. They were given only distilled water to drink and a constant daily diet that was estimated to contain 400 mg of calcium, 900 mg of phosphorus, 18 meq of magnesium, 110 meq of sodium, and 65 meq of potassium. Fasting blood samples were collected for measurement of serum calcium, ionized calcium, phosphorus, magnesium, creatinine, Gla protein, 25-hydroxyvitamin D (25-OHD), 1,25-dihydroxyvitamin D [$1,25(\text{OH})_2\text{D}$], and immunoreactive PTH. 24-h urines were collected for measurement of calcium, phosphorus, sodium, potassium, magnesium, creatinine, and cyclic AMP.

Serum and urinary calcium (2), phosphorus (3), creatinine (4), and magnesium (5) were measured by automated colorimetric methods. Serum ionized calcium was measured with a solid state ion electrode. Urinary sodium and potassium were determined by flame photometer. Serum 25-OHD was measured in duplicate at two concentrations by competitive protein binding with vitamin D-deficient rat serum (6) after extraction with acetonitrile, washing with phosphate buffer, chromatography on C-18 Sep-Pak, and elution with acetonitrile (7). 25-OHD was separated from other vitamin D metabolites before the binding assay by chromatography on silica Sep-Pak and elution with hexane-propanol (94:6) (7). Serum $1,25(\text{OH})_2\text{D}$ was measured by the method of Reinhardt et al. (7). Serum immunoreactive PTH was measured by radioimmunoassay with a COOH-terminal specific antibody from chicken 77125 at a dilution of 1:10,000 (8). Serum Gla protein was determined by radioimmunoassay (9). Urinary cyclic AMP was measured by radioassay with a binding protein (10). Results are expressed as nM/dl glomerular filtrate (GF) (11).

Statistical analyses were performed with nonpaired *t* test and correlation coefficient by standard methods. The percentage of ideal body weight was determined from tables of the Metropolitan Life Insurance Company, New York.

Results

Weights of the obese and nonobese subjects averaged 106 ± 6 and 68 ± 2 kg, respectively ($P < 0.01$). The mean age was 26 ± 1 yr for the obese and 24 ± 1 yr for the nonobese individuals.

The results are summarized in Tables I and II and in Fig. 1. Mean serum Gla protein, mean serum immunoreactive

1. Abbreviations used in this paper: GF, glomerular filtrate; PTH, parathyroid hormone.

Table I. Serum Values in Obese and Nonobese White Subjects

Subjects	Serum calcium	Serum Ca ²⁺	Serum phosphorus	Serum magnesium	Serum Gla protein*	Serum iPTH‡	Serum 25-OHD	Serum 1,25(OH) ₂ D
	mg/dl	mg/dl	mg/dl	meq/liter	ng/ml	pg/ml	ng/ml	pg/ml
Obese (12)	9.0±0.1	4.8±0.1	4.0±0.1	1.88±0.02	33±2	518±48	8±1	37±2
Nonobese (14)	9.0±0.1	4.8±0.1	3.9±0.2	1.85±0.04	24±3	243±33	20±2	29±2
P value	NS	NS	NS	NS	<0.02	<0.001	<0.001	<0.01

Results are given as mean±SE. Figures in parentheses are the number of subjects. * Serum Gla protein was measured in 11 obese and 14 nonobese subjects. ‡ Serum immunoreactive PTH (iPTH) was measured in 12 obese and 13 nonobese subjects.

PTH, and mean serum 1,25(OH)₂D were significantly higher in the obese as compared to the nonobese men and women (Table I). Mean serum calcium, serum ionized calcium, serum phosphorus, and serum magnesium were the same in the two groups. Mean serum 25-OHD was significantly lower in the obese than in the nonobese individuals. During the 2 d on the constant diet, mean urinary calcium was significantly lower in the obese than in the nonobese subjects (Table II). Mean urinary cyclic AMP and creatinine clearance were significantly higher in the obese subjects, and mean urinary phosphorus, potassium, and magnesium were the same in the two groups. As shown in Fig. 1, in all subjects there was a significant positive correlation between percentage of ideal body weight and urinary cyclic AMP ($r = 0.524, P < 0.01$). There was also a significant positive correlation between percentage of ideal body weight and serum immunoreactive PTH ($r = 0.717, P < 0.01$).

Discussion

Bone mass is increased (12–18) and urinary calcium is diminished (19) in black as compared with white individuals. We previously demonstrated increases in mean serum immunoreactive PTH, serum 1,25(OH)₂D, and urinary cyclic AMP, and confirmed that mean urinary calcium is reduced in normal nonobese blacks (20). Further, the demonstration that the blacks excreted an intravenous calcium load (15 mg calcium/kg body weight infused over 8 h) as efficiently as whites provided evidence that the reduction in urinary calcium in blacks was caused by increases in circulating PTH and not by primary enhancement of tubular reabsorption of the cation. Mean serum Gla protein, an index of PTH status (21), was

lower in blacks than in whites, despite the higher circulating PTH in the blacks. We interpreted these results to indicate that modification of the vitamin D-endocrine system, with enhanced renal tubular reabsorption of calcium and increased circulating 1,25(OH)₂D caused by increases in serum PTH, may contribute to the greater bone mass in blacks (20). The larger bone mass in them is attributed to increased muscle mass (13).

The present findings in obese white subjects are similar to the ones obtained by us in nonobese blacks. Thus, obese white individuals show increases in mean serum immunoreactive PTH (1), serum 1,25(OH)₂D, and urinary cyclic AMP, and decreases in urinary calcium as compared with nonobese white men and women. The reduction of urinary calcium in the obese individuals is all the more evident since it occurred despite an average increase of 24.8% in creatinine clearance. Obese postmenopausal women were found to have a lower urinary calcium-to-creatinine ratio as compared with nonobese postmenopausal women (22). One difference for which we have no explanation is the reduction of serum Gla protein in blacks in our previous study and the increase of serum Gla protein in the obese as compared with nonobese white subjects, which was seen in the present investigation. We interpret our results to indicate that alteration of the vitamin D-endocrine system also occurs in obesity, and is characterized by secondary increases in circulating PTH with consequent enhanced tubular reabsorption in calcium and increased renal production of 1,25(OH)₂D.

An alternative explanation for our findings in obese men and women is impaired intestinal absorption of calcium. This possibility appears unlikely, since malabsorption of calcium, if prolonged, would lead to a decline in bone mass. In this

Table II. Urinary Values in Obese and Nonobese White Subjects

Subjects	Urinary calcium	Urinary phosphorus	Urinary sodium	Urinary potassium	Urinary magnesium	Urinary cyclic AMP	Creatinine clearance
	mg/d	mg/d	meq/d	meq/d	meq/d	nM/dl GF	liter/d
Obese (12)	115±10	1,043±50	104±8	62±3	9.3±0.9	3.18±0.43	216±17
Nonobese (14)	166±13	938±37	124±8	62±4	9.2±0.4	1.84±0.25	173±6
P value	<0.01	NS	NS	NS	NS	<0.01	<0.01

Results are given as mean±SE of the average of two consecutive 24-h urine values in each subject. Figures in parentheses are the number of subjects.

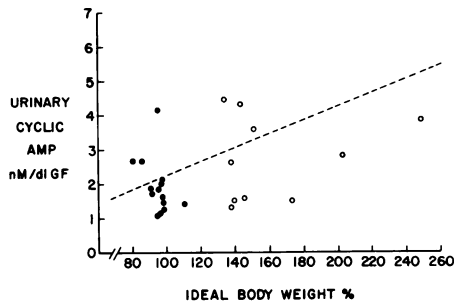


Figure 1. Relationship between urinary cyclic AMP and percentage of ideal body weight in obese and nonobese white subjects. In all of them, there was a significant positive correlation between the two determinations ($r = 0.524$, $P < 0.01$). ●, nonobese; ○, obese.

regard, measurement of bone mass in obese subjects has yielded conflicting results. Radiographic measurements of metacarpal cortical area showed that skeletal mass was greater in obese than in age-matched nonobese subjects (23). On the other hand, bone mass of the forearm determined by single-photon absorptiometry demonstrated values that were within the normal range (24). Urinary calcium-to-creatinine ratio in obese postmenopausal women, which was reduced as compared with values in nonobese postmenopausal women, correlated negatively with serum estrogen, and serum estrogens were shown to correlate with body weight (22). It is possible, therefore, that increases in circulating estrogen that occur in obesity because of increased peripheral conversion from androgens, could modify the skeletal response to PTH in obese women (22). It is unlikely, however, that this phenomenon accounts for changes in the vitamin D-endocrine system observed in obese men.

As noted already, increased bone mass in blacks is thought to result from an increased muscle mass (13). If greater strain on the skeleton produced by the increased body weight diminishes the skeletal response to PTH in obesity, the elevated serum immunoreactive PTH should decline with weight loss. Atkinson et al. (1) followed 27 massively obese subjects for periods of up to 1 yr after jejunal-ileal intestinal bypass, and noted that serum immunoreactive PTH decreased with weight loss. Indeed, the reductions in body weight and serum immunoreactive PTH paralleled each other. Thus, at 6 mo after surgery there was a significant correlation between the amount of weight loss, which averaged 44 kg, and the decrease in serum immunoreactive PTH (1).

Mean serum 25-OHD was lower in the obese than in the nonobese subjects. A reduction in mean serum 25-OHD was also found in two other groups of obese individuals, one of which had undergone jejunal-ileal bypass (24, 25). In these subjects, there was a progressive increase in serum 25-OHD after surgery to values that were in the normal range. In view of the present findings, and the previously described decline in elevation of serum immunoreactive PTH in association with weight loss after intestinal bypass (1), we attribute the low serum 25-OHD in obese subjects to feedback inhibition of hepatic synthesis of the metabolite by increased circulating 1,25(OH)₂D (26). Reversal of obesity and elevated serum immunoreactive PTH would lead to a decline in serum 1,25(OH)₂D and allow serum 25-OHD to return to values in the normal range. This explanation would also account for the

impaired production of 25-OHD in response to vitamin D challenge described in the obese individuals (24). Low values for serum 25-OHD were also observed in normal nonobese black men and women in whom mean serum 1,25(OH)₂D was increased (20). It is likely that reduction of serum 25-OHD in them results from impaired dermal production of vitamin D because of increased skin pigment (27).

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