CLINICAL STUDY

Patients with primary hyperparathyroidism have lower circulating sclerostin levels than euparathyroid controls

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Abstract

Objective: *In vitro* and *in vivo* studies in animal models have shown that parathyroid hormone (PTH) inhibits the expression of the *SOST* gene, which encodes sclerostin, an osteocyte-derived negative regulator of bone formation. We tested the hypothesis that chronic PTH excess decreases circulating sclerostin in humans.

Design: We studied 25 patients with elevated serum PTH concentrations due to primary hyperparathyroidism (PHPT) and 49 patients cured from PHPT after successful parathyroidectomy (PTx; euparathyroid controls (EuPTH)).

Methods: We measured plasma PTH and serum sclerostin levels and the serum markers of bone turnover alkaline phosphatase, P1NP, and β -CTX.

Results: As expected by the design of the study, mean plasma PTH was significantly higher ($P < 0.001$) in PHPT patients (15.3 pmol/l; 95% confidence interval (CI): 11.1–19.5) compared with that of EuPTH controls (4.1 pmol/l; 95% CI: 3.6–4.5). PHPT patients had significantly lower serum sclerostin values compared with those in EuPTH subjects (30.5 pg/ml; 95% CI: 26.0–35.1 vs 45.4 pg/ml; 95% CI: 40.5–50.2; $P < 0.001$) and healthy controls (40.0 pg/ml; 95% CI: 37.1–42.9; $P = 0.01$). Plasma PTH concentrations were negatively correlated with serum sclerostin values ($r = -0.44$; $P < 0.001$). Bone turnover markers were significantly correlated with PTH, but not with sclerostin.

Conclusion: Patients with PHPT have significantly lower serum sclerostin values compared with PTx controls with normal PTH concentrations. The negative correlation between PTH and sclerostin suggests that *SOST* is downregulated by PTH in humans.

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Introduction

Parathyroid hormone (PTH) exerts its calciotropic action by acting directly on bone and kidney and indirectly on the intestine to increase the transport of calcium to the circulation. The skeletal effect of PTH is to increase the rate of remodeling (1), and it is generally believed that this effect is achieved by the binding of PTH to its specific receptor (PTH1R) on stromal/osteoblastic cells of the bone marrow. This in turn stimulates the production of RANK ligand and decreases that of its decoy receptor osteoprotegerin (2–5). However, recent *in vitro* and animal studies suggest that some of the effects of PTH on bone are also exerted by specific binding of the hormone to PTH1R in osteocytes, resulting in inhibition of the expression of the *SOST* gene (6–9). This gene encodes sclerostin, a protein exclusively expressed in osteocytes in the skeleton (10), which decreases bone formation by binding to LRP5/6, resulting in inhibition of the Wnt signaling pathway in osteoblasts (11, 12).

Whether chronic PTH excess has similar effects on sclerostin secretion in humans as in animal models has

not so far been investigated. In this study, we tested the hypothesis that chronic hypersecretion of PTH, as seen in patients with hyperparathyroidism, may decrease sclerostin secretion, and that PTH may thus represent a potential regulator of sclerostin production in humans. To this effect, we measured sclerostin in serum of patients with untreated primary hyperparathyroidism (PHPT) and in a control group of patients with PHPT after establishment of cure following parathyroidectomy (PTx).

Subjects and methods

Patients

Thirty-four consecutive patients with PHPT that was untreated, persistent, or recurrent following PTx, and 54 patients cured after successful PTx (euparathyroid controls, EuPTH) were studied. Inclusion criteria included willingness to participate in the study, no impairment in renal function (serum creatinine levels $< 120 \mu\text{mol/l}$), adequate vitamin D status (25-hydroxy

vitamin D (25-OHD) levels > 50 nmol/l), and no use of bone and mineral metabolism modifying agents such as bisphosphonates, calcimimetics, or glucocorticoids.

We defined PHPT as plasma PTH concentrations above the upper limit of the normal laboratory reference range (> 8 pmol/l) in the presence of increased serum calcium concentrations (> 2.55 mmol/l).

Patients with EuPTH were included when cure was confirmed by postoperative normalization of serum PTH and calcium concentrations, which was sustained for at least 6 months after PTx.

As per inclusion criteria, nine patients were excluded from the PHPT group, three because of impaired renal function and six because of the use of bisphosphonates, and six were excluded from the EuPTH group because of the use of bisphosphonates.

The study was approved by the Medical Ethics Committee of the Leiden University Medical Center, and informed consent was obtained from all patients.

Methods

Serum biochemistry Serum calcium was adjusted for albumin binding; phosphate and creatinine were measured by semi-automated techniques. Serum alkaline phosphatase (ALP) activity was measured using a fully automated P800 modulator system (Roche BV). P1NP (a marker of bone formation) and β -CTX (a marker of bone resorption) were determined using the E-170 system (Roche BV). Plasma PTH was measured using the Immulite 2500 (Siemens Diagnostics, Breda, The Netherlands) and serum 25-OHD was measured using the LIAISON 25-OHD TOTAL assay (DiaSorin S.A./N.V., Bruxelles, Belgium).

Sclerostin measurement Sclerostin was measured in serum by an electrochemiluminescence assay (MSD 96-well MULTI-ARRAY Human Sclerostin Assay, Gaithersburg, MD, USA) in which two polyclonal antibodies were raised against the whole sclerostin molecule. The sclerostin standard for the assay was produced in an NS0-derived myeloma cell line, and the purity was checked by SDS-PAGE gel with silver stain. In our hands, the intra- and inter-assay coefficient of variation values were 4.9 and 7% respectively, the detection limit was ± 1 pg/ml, and the detection range was 1–10 000 pg/ml.

Sclerostin was measured in the serum of 77 healthy subjects (30 males and 47 females, aged 20–77 years). All had normal serum calcium concentration, renal function, and bone turnover and none were using bisphosphonates, the calcimimetic cinacalcet, or glucocorticoids. Sclerostin was detected in the serum of all healthy subjects: mean 40.0 pg/ml (95% CI = 37.1–42.9 pg/ml), range 12.4–68.19 pg/ml, while it was undetectable in the serum of three patients with sclerosteosis.

Statistical analysis

Data were analyzed using SPSS 16.0 (SPSS Inc., Chicago, IL, USA). Between-group differences in baseline characteristics and serum biochemistry were assessed by Student's *t*-test. Pearson correlation coefficients were calculated to assess correlations between PTH (after logarithmic transformation), sclerostin, and biochemical markers of bone turnover. A probability level of random difference of 0.05 was considered significant.

Results

Baseline characteristics

There were no differences in age, gender, weight, or body mass index between patients with PHPT and EuPTH controls (Table 1).

As expected by the inclusion criteria, mean serum calcium and PTH concentrations were significantly higher and those of phosphate significantly lower in the PHPT group compared with the EuPTH group in which PTH concentrations were detectable in all patients. There were no differences in serum 25-OHD or creatinine concentrations between the two groups.

Patients with PHPT had significantly higher levels of biochemical markers of bone formation (P1NP) and bone resorption (β -CTX) compared with EuPTH controls. Combining all patients, there was a significant positive correlation between plasma PTH concentrations and the concentrations of all three measured biochemical markers of bone turnover (ALP: $r = 0.23$, $P = 0.047$; P1NP: $r = 0.45$, $P < 0.001$; β -CTX: $r = 0.54$, $P < 0.001$). There was also a significant correlation between PTH and P1NP ($r = 0.51$, $P = 0.009$) in the PHPT group, but not between PTH and ALP ($r = 0.35$, $P = 0.085$), or PTH and β -CTX ($r = 0.31$, $P = 0.13$). In the EuPTH group alone, PTH was not correlated with any of the biochemical markers of bone turnover (Table 2).

Serum sclerostin

Mean serum sclerostin level of patients with PHPT (30.5 pg/ml, 95% CI: 26.0–35.1) was significantly lower than that of patients with EuPTH and healthy controls (45.4 pg/ml, 95% CI: 40.5–50.2; $P < 0.001$,

Table 1 Subject characteristics. Values are given as mean $\pm$ s.d.

	PHPT	EuPTH	<i>P</i> value ^a
Male:female	10:15	13:36	0.41
Age (years)	59.6 $\pm$ 16.7	62.4 $\pm$ 10.9	0.44
Weight (kg)	80.0 $\pm$ 18.2	81.1 $\pm$ 15.3	0.78
BMI (kg/m ²)	27.0 $\pm$ 6.0	28.1 $\pm$ 4.6	0.42

^aPHPT versus EuPTH.

Table 2 Biochemical measurements. Values are given as mean $\pm$ s.d.

	PHPT	EuPTH	Reference range	P value ^a
Calcium homeostasis				
PTH (pmol/l)	15.3 $\pm$ 10.7	4.1 $\pm$ 1.6	1.5–8.0	<0.001
Calcium (mmol/l)	2.61 $\pm$ 0.13	2.26 $\pm$ 0.12	2.15–2.55	<0.001
Phosphate (mmol/l)	0.92 $\pm$ 0.14	1.14 $\pm$ 0.24	0.90–1.50	<0.001
25-OHD (nmol/l)	53.1 $\pm$ 34.3	53.9 $\pm$ 20.2	30–120	0.90
Creatinine (μ mol/l)	80.2 $\pm$ 18.1	76.8 $\pm$ 14.7	44–80	0.39
Bone turnover				
ALP (U/l)	87.0 $\pm$ 23.0	75.7 $\pm$ 23.6	40–20	0.055
P1NP (ng/ml)	45.9 $\pm$ 16.9	34.0 $\pm$ 15.5	16–80	0.004
β -CTX (ng/ml)	0.32 $\pm$ 0.15	0.17 $\pm$ 0.11	0.01–0.66	<0.001

^aPHPT versus EuPTH.

and 40.0 pg/ml, 95% CI: 37.1–42.9; $P=0.01$ respectively; Fig. 1). There was no significant difference in mean sclerostin values between EuPTH and healthy subjects ($P=0.13$).

There was no significant correlation between PTH and sclerostin concentrations within each individual group of patients but there was a significant negative correlation between sclerostin and PTH when all patients were pooled together ($r=-0.44$, $P<0.001$; Fig. 2).

There was no significant relationship between serum sclerostin and biochemical markers of bone turnover in patients with PHPT or in all patients combined.

Discussion

Data from our study demonstrates that in humans, chronic PTH excess, as observed in patients with PHPT, is associated with a significant decrease in circulating sclerostin and that there is a significant negative correlation between PTH and serum sclerostin levels. Taken together, these results suggest that, similar to animal models (6–9), PTH has a regulatory role in sclerostin production in humans also.

Sclerostin, a glycoprotein expressed by osteocytes in bone and encoded by the *SOST* gene, has emerged in recent years as an important regulator of bone formation in humans as well as in animals (13, 14). Inactivating mutations of the *SOST* gene leading to sclerostin deficiency have been shown to be associated with the rare skeletal disorder sclerosteosis, which is characterized by a marked increase in bone mass (13). Deletion of the *Sost* gene in mice has also been shown to increase bone formation, bone mass, and bone strength (14) demonstrating an evolutionary conservation of the function of sclerostin in the regulation of bone formation. Moreover, inhibition of sclerostin secretion by an MAB to sclerostin has been shown to increase bone formation and bone mass in rodents, primates, and humans (14–16). Conversely, transgenic mice overexpressing *Sost* have low bone mass and impaired biomechanical competence (17). The mechanism of action of sclerostin to decrease bone

formation involves inhibition of the Wnt signaling pathway (11, 12), although its precise molecular mechanism and factors controlling its secretion are yet to be determined. Recent animal studies have shown that mechanical loading and high PTH levels down-regulate the expression of *Sost* in osteocytes and decrease the production of sclerostin resulting in stimulation of bone formation (8, 18).

Human studies of sclerostin regulation have lagged behind due to lack of noninvasive techniques to determine sclerostin production. A number of assays have been recently developed for the measurement of sclerostin in blood. In our study, we used a sclerostin assay that proved to have excellent performance characteristics in our hands with a wide detection range. Sclerostin was detectable in serum in all healthy subjects studied, suggesting that the protein is secreted

Figure 1 Serum sclerostin levels in PHPT, EuPTH, and healthy subjects. *** $P<0.001$, ** $P=0.01$ (Student's *t*-test).

Figure 2 Relationship between circulating sclerostin and PTH levels. $r = -0.44$, $P < 0.001$ (Pearson's correlation). PTH levels were log transformed because of skewness.

and enters the circulation, while it was undetectable in three patients with sclerosteosis in whom it was measured. The difference in absolute values with other commercial assays is not clear and may be related to different methodologies. We chose to study patients with PHPT in order to mimic as closely as possible the effect of chronic PTH excess on sclerostin, as previously studied in animals (6, 8). In addition, we chose to use as controls, patients with PHPT cured after PTx to exclude potential confounding factors, other than PTH excess.

We show here that serum sclerostin levels are significantly decreased in patients with chronic PTH excess due to PHPT compared with EuPTH and healthy subjects although there was an overlap of values between the groups. PTH has been shown to decrease *SOST* transcription *in vitro* (6, 7), and continuous and intermittent chronic administration of PTH to rodents is associated with decreased *Sost* mRNA and sclerostin expression in osteocytes (6, 7, 9). Moreover, transgenic mice expressing a constitutively active PTH receptor in osteocytes exhibit decreased expression of sclerostin and increased Wnt signaling associated with increased bone mass (8). Additional evidence for an interaction between PTH and *Sost*/sclerostin was recently provided by a study showing that the anabolic actions of PTH on bone was blunted in *Sost*-overexpressing mice (19). In keeping with the notion of a regulatory role of PTH for sclerostin production, our data show a significant correlation between circulating PTH and sclerostin. These data extend those of Mirza *et al.* (20), who recently reported a negative relationship between serum PTH and sclerostin in healthy postmenopausal women,

and those of Drake *et al.* (21), who showed that intermittent PTH treatment decreased serum sclerostin levels in postmenopausal women. Put together, these data suggest that *SOST* regulation by PTH is also conserved between humans and rodents.

As expected in the presence of chronic PTH excess, patients with PHPT had increased bone turnover as indicated by increased biochemical parameters of bone formation and resorption. There was a significant relationship between circulating PTH concentrations and serum P1NP and β -CTX. However, we did not find a significant relationship between biochemical parameters of bone turnover and serum sclerostin, either in patients with PHPT or the combined group of PHPT and EuPTH subjects. This lack of correlation between sclerostin and bone turnover markers was previously reported in healthy postmenopausal women (20) but not in elderly women (22). The functional significance of circulating sclerostin is as yet to be established.

The actions of PTH on bone are complex and involve a variety of signaling pathways in bone marrow stroma cells, osteoblasts, and osteocytes (23, 24). Despite the significant progress in our understanding of the actions of PTH on bone, it should be appreciated that the cellular and molecular actions of PTH, which determine the action of the hormone on bone remodeling and bone balance, have only been partially unraveled and studies are needed to further elucidate these actions.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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