Nocturnal calcium, phosphorus and parathyroid hormone in the diagnosis of concealed and subclinical hypoparathyroidism

Leah Even1,3, Tarif Bader1 and Ze'ev Hochberg2,3

1Department of Pediatrics, Nahariya Hospital, Nahariya, Israel, 2Division of Pediatric Endocrinology, Rambam Medical Center, Meyer Children’s Hospital, PO Box 9602, Haifa 31096, Israel and 3Faculty of Medicine, Technion, Israel Institute of Technology, Haifa, Israel

(Correspondence should be addressed to Z Hochberg who is now at Division of Pediatric Endocrinology, Rambam Medical Center, Meyer Children’s Hospital, PO Box 9602, Haifa 31096, Israel; Email: z_hochberg@rambam.health.gov.il)

Abstract

Context: Circadian rhythms of plasma parathyroid hormone (PTH) show peak values at night, whereas serum calcium levels peak in the evening and display a nadir at night.

Hypotheses: Subclinical hypoparathyroidism (HPT) can be detected by utilizing the knowledge of diurnal variations. Thalassemia major (TM) may provide a model system of subclinical HPT.

Design: Nocturnal plasma PTH and serum calcium values were determined in 13 TM patients with normal morning serum calcium levels as compared with the corresponding values in eight healthy control subjects.

Results: Six patients with TM presented a nadir serum calcium level of 8.3 mg/dl or lower (hypoCa TM) at 0200 h, whereas the remaining seven showed nadir levels of 8.4 mg/dl or higher (normoCa TM). Patients with hypoCa TM displayed a drop between peak and nadir of 1.2±0.5 mg/dl as compared with a considerably smaller fall of 0.3±0.7 mg/dl in control subjects (P<0.05). NormoCa TM patients experienced comparable nocturnal variation to that of control subjects. Patients from both the hypoCa and normoCa TM groups presented significantly lower nocturnal PTH levels than those of control subjects and lost the nocturnal PTH variation characteristic of healthy subjects. A plot of all serum calcium against plasma PTH levels provides a clear distinction of the three groups.

Conclusions: All 13 daytime normocalcemic TM patients presented a certain degree of HPT. The hypoCa TM group displayed a concealed HPT detected in all, except the morning sampling, whereas normoCa TM patients experienced subclinical HPT observed in the absence of nocturnal HPT variation. Nocturnal measurements of serum minerals thus enhance the sensitivity of HPT diagnosis.
Patients and methods

The Nahariya Hospital Internal Review Board approved the study protocol, and both patients and parents signed informed consents. Thirteen TM patients, age 17 ± 3.3 (mean ± s.d.) years, and Tanner stage 1–4, were recruited for this study. All of them presented with normal morning levels of serum calcium, as defined by the adolescence lowest normal value of 8.4 mg/dl (2.1 mmol/l) (7). Eight sex- and Tanner stage-matched healthy subjects served as controls. Tanner stage matching was employed as an additional measure because mineral metabolism correlates better with pubertal development than age (7, 8). All TM patients were regularly treated by transfusion of packed red cells every three weeks (at a pre-transfusion hemoglobin level of 7.8–10.2 g/dl), and received daily chelating therapy with s.c. desferrioxamine mesylate (Desferal, Teva, Israel). Patients did not receive calcium or vitamin D supplements.

Subjects were admitted at afternoon hours, 3 weeks after the last blood transfusion, to eliminate the effect of citrate on calcium level. After a light non-dairy supper at 1800 h, serum calcium, phosphorus, creatinine and plasma PTH were taken at four time points (2000, 0000 (midnight), 0200, and 0800 h). 25 OH-Vit D, 1,25 dihydroxyvitamin D, ferititin and hemoglobin levels were measured by the Aeroset system and the Architect c8000 system (Abbott Laboratories, Abbott Park, IL, USA). Serum calcium levels of 8.4–10.2 mg/dl (2.1–2.3 mmol/l), and were designated hypocalcemic (hypoCa) TM, the remaining seven patients showed nadir calcium levels over 8.3 mg/dl (8.4–9.2 mg/dl, 2.1–2.3 mmol/l), and were designated normocalcemic TM (normoCa TM) patients.

The nocturnal serum phosphorus levels of the control group showed a mirror image of serum calcium levels, with rising values from 2000 h through 0200 h, receding by 0800 h (Fig. 2). The hypoCa TM group presented higher phosphorus levels at all time points (P < 0.05), whereas the NormoCa TM group showed normal phosphorus levels, as compared with control subjects. The nocturnal variations did not reach statistical significance for any of the groups.

Control subjects showed predictably nocturnal peak levels of plasma PTH (Fig. 3). The hypoCa and normoCa TM groups presented significantly lower PTH levels than those of control subjects at all time points. Importantly, patients from both groups lost the nocturnal variation, characteristic of control subjects (ANOVA P < 0.05).

Dot plot of all serum calcium levels as a function of time, with significant differences (multivariate test P < 0.001).

Results

The nocturnal serum calcium levels in the control group corresponded to previous reports of peak levels at 2000 h decreasing to nadir at 0200 h and recovered at 0800 h (Fig. 1). Among patients with TM, five showed 0200 h nadirs of 8.3 mg/dl or lower (6.8–8.3 mg/dl, 1.7–2.0 mmol/l), and were designated hypocalcemic (hypoCa) TM. The remaining seven patients showed nadir calcium levels over 8.3 mg/dl (8.4–9.2 mg/dl, 2.1–2.3 mmol/l), and were designated normocalcemic TM (Table 1). The former group was older (mean 19.4 years) than the latter (16.3 years). HypoCa TM patients presented a decrease between peak and nadir of 1.2 ± 0.5 mg/dl when compared with a decline of 0.3 ± 0.7 mg/dl in control subjects (P < 0.05). NormoCa TM patients displayed comparable nocturnal variation to that observed in controls.

Control subjects showed predictably nocturnal peak levels of plasma PTH (Fig. 3). The hypoCa and normoCa TM groups presented significantly lower PTH levels than those of control subjects at all time points. Importantly, patients from both groups lost the nocturnal variation, characteristic of control subjects (ANOVA P < 0.05).

Patients and methods

The Nahariya Hospital Internal Review Board approved the study protocol, and both patients and parents signed informed consents. Thirteen TM patients, age 17 ± 3.3 (mean ± s.d.) years, and Tanner stage 1–4, were recruited for this study. All of them presented with normal morning levels of serum calcium, as defined by the adolescence lowest normal value of 8.4 mg/dl (2.1 mmol/l) (7). Eight sex- and Tanner stage-matched healthy subjects served as controls. Tanner stage matching was employed as an additional measure because mineral metabolism correlates better with pubertal development than age (7, 8). All TM patients were regularly treated by transfusion of packed red cells every three weeks (at a pre-transfusion hemoglobin level of 7.8–10.2 g/dl), and received daily chelating therapy with s.c. desferrioxamine mesylate (Desferal, Teva, Israel). Patients did not receive calcium or vitamin D supplements.

Subjects were admitted at afternoon hours, 3 weeks after the last blood transfusion, to eliminate the effect of citrate on calcium level. After a light non-dairy supper at 1800 h, serum calcium, phosphorus, creatinine and plasma PTH were taken at four time points (2000, 0000 (midnight), 0200, and 0800 h). 25 OH-Vit D, 1,25 dihydroxyvitamin D, ferititin and hemoglobin levels were measured by the Aeroset system and the Architect c8000 system (Abbott Laboratories, Abbott Park, IL, USA). Serum calcium levels of 8.4–10.2 mg/dl (2.1–2.3 mmol/l), and were designated hypocalcemic (hypoCa) TM, the remaining seven patients showed nadir calcium levels over 8.3 mg/dl (8.4–9.2 mg/dl, 2.1–2.3 mmol/l), and were designated normocalcemic TM (normoCa TM) patients.

The nocturnal serum phosphorus levels of the control group showed a mirror image of serum calcium levels, with rising values from 2000 h through 0200 h, receding by 0800 h (Fig. 2). The hypoCa TM group presented higher phosphorus levels at all time points (P < 0.05), whereas the NormoCa TM group showed normal phosphorus levels, as compared with control subjects. The nocturnal variations did not reach statistical significance for any of the groups.

Control subjects showed predictably nocturnal peak levels of plasma PTH (Fig. 3). The hypoCa and normoCa TM groups presented significantly lower PTH levels than those of control subjects at all time points. Importantly, patients from both groups lost the nocturnal variation, characteristic of control subjects (ANOVA P < 0.05).

Dot plot of all serum calcium levels as a function of time, with significant differences (multivariate test P < 0.001).

Results

The nocturnal serum calcium levels in the control group corresponded to previous reports of peak levels at 2000 h decreasing to nadir at 0200 h and recovered at 0800 h (Fig. 1). Among patients with TM, five showed 0200 h nadirs of 8.3 mg/dl or lower (6.8–8.3 mg/dl, 1.7–2.0 mmol/l), and were designated hypocalcemic (hypoCa) TM. The remaining seven patients showed nadir calcium levels over 8.3 mg/dl (8.4–9.2 mg/dl, 2.1–2.3 mmol/l), and were designated normocalcemic TM (Table 1). The former group was older (mean 19.4 years) than the latter (16.3 years). HypoCa TM patients presented a decrease between peak and nadir of 1.2 ± 0.5 mg/dl when compared with a decline of 0.3 ± 0.7 mg/dl in control subjects (P < 0.05). NormoCa TM patients displayed comparable nocturnal variation to that observed in controls.

The nocturnal serum phosphorus levels of the control group showed a mirror image of serum calcium levels, with rising values from 2000 h through 0200 h, receding by 0800 h (Fig. 2). The hypoCa TM group presented higher phosphorus levels at all time points (P < 0.05), whereas the NormoCa TM group showed normal phosphorus levels, as compared with control subjects. The nocturnal variations did not reach statistical significance for any of the groups.

Control subjects showed predictably nocturnal peak levels of plasma PTH (Fig. 3). The hypoCa and normoCa TM groups presented significantly lower PTH levels than those of control subjects at all time points. Importantly, patients from both groups lost the nocturnal variation, characteristic of control subjects (ANOVA P < 0.05).

Dot plot of all serum calcium levels as a function of time, with significant differences (multivariate test P < 0.001).

Table 1 Characteristics of the study subjects.

<table>
<thead>
<tr>
<th>Age (year)</th>
<th>N</th>
<th>Sex F/M</th>
<th>Hb (mg%)</th>
<th>Ferritin (ng/ml)</th>
<th>25OH Vit D (pg/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>19.4 ± 2.3</td>
<td>8</td>
<td>3/5</td>
<td>12.6 ± 1.8</td>
<td>34 ± 13.9</td>
<td>6200 ± 1010</td>
</tr>
<tr>
<td>16.3 ± 4.7</td>
<td>6</td>
<td>3/2</td>
<td>12.0 ± 0.5</td>
<td>8.0 ± 0.3</td>
<td>4600 ± 800</td>
</tr>
<tr>
<td>10.0 ± 0.7</td>
<td>7</td>
<td>3/4</td>
<td>6.8 ± 0.5</td>
<td>Normal (10–40)</td>
<td>Normal (16–42)</td>
</tr>
</tbody>
</table>

HypoCa TM, thalassemia major patients with calcium levels < 8.4 mg/dl; NormoCa TM, thalassemia major patients with calcium levels ≥ 8.4 mg/dl; F, female; M, male; Hb, hemoglobin.
Discussion

Healthy subjects exhibit a diurnal variation of serum calcium and plasma PTH (1). Our study shows that measuring serum calcium levels at a single time point in the morning yields unreliable diagnosis of HPT, as previously suggested for other reasons (9). Nocturnal sampling, however, reveals both concealed and subclinical cases of hypoparathyroidism.

Given the well-documented delayed puberty in TM (6), one had to select either age or puberty for matching of the study and control groups. With regard to the known effect of puberty on mineral metabolism (8), we elected the latter option, with an obvious outcome of younger controls.

Modeling the hypothesis in TM, where the odds for developing HPT are high (11), enabled us to track HPT at its inception. A surprising finding was that every single one of our normocalcemic TM patients presented a certain degree of HPT, exceedingly more than the reported 4.5–30% (2–6). The possibility cannot be excluded that our patients are also less compliant with their chelating therapy, as evident from their abnormally high ferritin levels. TM patients of the present report also presented vitamin D insufficiency, as compared with the recommended pubertal levels (10). It is recommended that these very sick children, whose outdoors activities may be limited, should receive vitamin D supplementation.

Our data show two levels of potentially missed diagnosis, which we designate as concealed and subclinical HPT. The former is characterized by nocturnal hypocalcemia and the latter becomes apparent when plasma levels of PTH are plotted as a function of serum calcium levels, showing PTH insufficiency. TM patients with concealed HPT were older than those with a subclinical disease, indicating the progressive nature of HPT.

On a plot scale of all serum calcium against plasma PTH levels, the concept of concealed and subclinical HPT becomes evident. HPT shows decreases that are masked by the routine morning sampling, and may be unraveled by nocturnal sampling. Patients from the latter group show subclinical HPT with normocalcemia and relative PTH insufficiency. We recommend the periodic use of such measurements in all TM, and suggest that these tools may be applicable to other cases of suspected HPT.

References


9 Ahmad AM, Hopkins MT, Fraser WD, Ooi CG, Durham BH & Vora J P. Parathyroid hormone secretory pattern, circulating activity and effect on bone turnover in adult growth hormone deficiency. *Bone* 2003 **32** 170–179.


Received 6 July 2006
Accepted 11 October 2006