Spontaneous bilateral femoral neck fracture in a man with end-stage renal disease and severe secondary hyperparathyroidism

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Abstract
Medical treatment is the primary recommended method for the management of severe secondary hyperparathyroidism (SHPT) in pre-dialysis individuals. This paper reports a case of end-stage renal disease (ESRD) and severe SHPT with pathologic bilateral hip fractures. A 61-year-old man, as a known case of ESRD, with radial osteoporosis and persistent SHPT (intact PTH >2000 pg/mL) and hypercalcemia with appropriate level of phosphorus and 25(OH) D who did not respond to medical treatment referred to hospital with multiple fractures. The patient referred for subtotal parathyroidectomy. One month after parathyroidectomy, parathyroid hormone (PTH) level significantly diminished from its preoperative values. Serum calcium and phosphorus returned to normal values and muscle weakness progressively improved until the patient became able to walk again. Parathyroidectomy in patients with severe secondary hyperparathyroidism (persistent serum levels of intact PTH> 800 pg/mL) is suggested, particularly when is associated with severe hypercalcemia and/or hyperphosphatemia, and progressive and debilitating hyperparathyroidism bone disease, that are refractory and resistant to medical therapy.

Keywords: Secondary hyperparathyroidism, Chronic kidney disease, End-stage renal disease, Hypercalcemia, Cardiovascular disease, Tertiary hyperparathyroidism, Parathyroidectomy

Case Presentation
A 61-year-old man, as a known case of ESRD for 11 years with unknown etiology who was under hemodialysis three times a week, referred to the hospital with multiple fractures in right knee and fifth, sixth, seventh anterior ribs in the left side. Patient was under treatment of calcitriol 0.25 mg two times daily, tablets of calcium carbonate 500 mg/four times a day, and sub-cutaneous erythropoietin three times a week.

In physical examination, he had ash-colored dry skin with scratch marks, hyper-pigmented papules over the scapular area, thoracic kyphosis, globular abdomen, arteriole-venous (A-V) shunt on the left arm and decreased range of motion in right shoulder, painful active and passive motion of knees, and also a bipedal edema.

Received: 8 January 2017, Accepted: 25 April 2017, ePublished: 12 May 2017

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Implication for health policy/practice/research/medical education

Parathyroidectomy is suggested in patients with severe secondary hyperparathyroidism (persistent serum levels of intact PTH> 800 pg/mL) is suggested particularly when it is associated with severe hypercalcemia and/or hyperphosphatemia with progressive and debilitating hyperparathyroidism bone disease, that is usually refractory and resistant to medical therapy.

To investigate the causes of multiple pathologic fractures, various para-clinical analysis were conducted. According to laboratory data, serum calcium was 10.8 mg/dL (8.5-10.5), phosphorus was 4.5 mg/dL, and albumin value was 3.6 g/dL (3.5-5). Additionally, serum alkaline phosphate was 2495 IU/L, serum 25(OH)D3 was 24 ng/mL (30-50 ng/mL), and finally intact parathyroid hormone (iPTH) was 2281 pg/mL (normal values: 15-65 pg/mL). Since hypercalcemia was mild, calcium carbonate (CaCo3) was discontinued. Several weeks after discontinuing CaCo3, the laboratory test was repeated and serum calcium level was 10.5 mg/dL, serum phosphorus level was 4 mg/dL, and serum alkaline phosphate level was 2514 IU/L and also serum iPTH level was 2500 pg/mL. Around one month after hip replacement, a bone mineral densitometry (BMD) was conducted. It detected low density in cortical bone due to SHPT. The results of T-scores were as follows: (Lumbar Spine) T-score = -1.4 , Z-score= -0.7 (Fore Arm) T-score = -6.7, Z-score = -5.8

In imaging study, renal sonography showed several stones (5-15 mm) in both kidneys. In echocardiography, patient had left ventricular ejection fraction of 50%. Additionally, mild MR and TR were observed. The patient had several lytic lesions with decreased bone density in pelvic X-ray, rotator cuff calcific tendonitis in shoulder X-ray, and rugger jersey spine in thoracolumbar X-ray (Figure 1A, B).

According to the persistence of severe SHPT and hypercalcemia and while the patient did not respond to medical treatment after several months, he referred for subtotal-parathyroidectomy. Around 3/5 of parathyroid glands were removed. Nodular parathyroid hyperplasia was the result of pathology report (Figure 2).

One month after parathyroidectomy, the laboratory tests showed a significant reduction in parathormone, as compared with preoperative values. Serum calcium returned to normal level (Ca; 8.2 mg/dL, P; 4 mg/dL, Alp; 1910 IU/L, iPTH; 190 pg/ml) too. Additionally, bone pain symptoms and muscle weakness progressively improved until the patient had became able to walk again.

Discussion

Initial treatment for managing SHPT in pre-dialysis individuals is the management of hyperphosphatemia, first initiated with limiting phosphorus diet. Although optimal limit of phosphate is unclear, 900 mg/d is suggested. If hyperphosphatemia does not improve by diet restriction, then calcium carbonate or non-calcium phosphate binders (based on serum Ca) should be administrated. Then, vitamin D deficiency should be treated if 25(OH)D is lower than 30 ng/mL. If elevated PTH remains persistent over a 6-month interval, we can propose administering a low dose active oral vitamin D analog. In patients who are resistant to treatment regimens containing vitamin D analogues and calcium supplements, and phosphate binders, it might be useful to administer cinacalcet (4).

Although cinacalcet therapy of subjects on dialysis with SHPT improves clinical outcomes, especially reduces the risk of parathyroidectomy, fractures, and cardiovascular hospitalizations, there is no statistically significant reduction in the rate of mortality (2,7).

Current guidelines for hemodialysis patients recommend; maintaining iPTH levels in a range which is approximately two to nine times higher than the normal limit (6). In elevated serum PTH, administration of calcitriol, vitamin D analogs, or calcimimetics, or a combination of these drugs will reduce serum PTH level. However, in observed patients, if hypercalcemia with serum Ca>10.2 mg/dL and hyperphosphatemia with P>6 mg/dL occurs, calcitriol or other vitamin D analogs should be promptly reduced or discontinued.

Due to the presence of hypercalcemia in our patient, we stopped calcium supplementation therapy and calcitriol. Despite discontinuing calcium supplementation and calcitriol after several months, serum calcium remained...

Figure 1. A; Rugger jersey spine in thoracolumbar X-ray. B; rotator cuff (calcific tendonitis) in shoulder X-ray.

Figure 2. Nodular parathyroid hyperplasia in secondary hyperparathyroidism in ESRD patient.
above 10 mg/dL and iPTH level did not reduce. In patients with severe hyperparathyroidism (persistent serum levels of intact PTH >800 pg/mL) associated with hypercalcemia and/or hyperphosphatemia that are refractory and resistant to medical therapy NKF recommended parathyroidectomy (8). Recently a study in Japan has suggested surgical indication, when there is a high PTH level above 500 pg/mL. A persistent high intact parathormone level (>500 pg/mL), when accompanied by hypercalcemia (>10.2 mg/dL), and/or presence of high serum phosphorus levels (>6.0 mg/dL) which cannot be corrected by medical therapy, is also considered as the surgical indication for parathyroidectomy. Likewise, parathyroidectomy is indicated when parathyroid glands are expanded (volume of the biggest parathyroid gland is >500 mm³ in ultrasonography) (5). Furthermore, individuals with high bone turnover, presence of osteitis fibrosa on X-ray, severe symptoms of secondary HPT, development of ectopic calcifications, calcification, progression of bone loss, and anemia resistant to erythropoietin therapy should definitely undergo parathyroidectomy (5). Since, there are no long-term follow up controlled studies on patients with persistent moderate hyperparathyroidism (iPTH 300-800 pg/mL), such patients are usually managed medically (5).

Our patient had persistent elevated intact PTH levels (>2000 pg/mL), accompanied by hypercalcemia (>10.2 mg/dL), that did not respond to appropriate medical therapy. Serum PTH alone is not a sufficient evidence for parathyroidectomy in asymptomatic patients, however, our case had several symptoms with extra-skeletal calcification and debilitating hyper-parathyroid bone disease with multiple non-traumatic fractures that indicated parathyroidectomy was a useful modality.

**Conclusion**

The most important factors for stimulating parathyroid gland function in patients with kidney insufficiency are hypocalcaemia, diminished 1,25-dihydroxyvitamin D levels, and hyperphosphatemia. It seems that calcitriol therapy is effectual for managing parathormone levels (by monitoring serum calcium and phosphorus levels). While there is no sufficient evidence to justify parathyroidectomy in ESRD subjects (since there is no recent studies and randomized controlled trials in this field), we can recommend parathyroidectomy for patients with severe SHPT (persistent serum levels of intact PTH >800 pg/mL), associated with severe hypercalcemia and/or hyperphosphatemia, progressive and debilitating hyperparathyroidism bone disease, progressive extra-skeletal calcification or calciphylaxis, that are refractory and resistant to medical therapy.

**Authors’ contribution**

SD and MF contributed equally to the manuscript.

**Conflicts of interest**

The authors report no conflicts of interest. The authors alone are responsible for the content and writing of the article.

**Ethical considerations**

Ethical issues (including plagiarism, data fabrication, double publication) have been completely observed by the authors.

**Funding/Support**

None.

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