

PRIMARY HYPERPARATHYROIDISM PRESENTED AS HUMERAL BROWN TUMOR AND MULTIPLE BONE FRACTURES



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INTRODUCTION

Primary hyperparathyroidism (PHPT) is frequently diagnosed as an incidental finding of hypercalcemia. Overt bone disease (osteitis fibrosa cystica or brown tumor), with history of pathologic fractures is a rare presentation.

CASE PRESENTATION

48-year-old woman

Medical history

- multiple bone fractures in the last six years left patella and both wrists
- Surgery, four years before, to lytic lesion of the right humerus, assumed as aneurysmal bone cyst -> last follow-up CT, revealed recurrence of lytic lesion and she was propose to shoulder arthroplasty

Admitted with

- pain in the right thigh after a trivial fall
- generalized weakness and lethargy over the last 2 years



X-rays revealed:

right femur

iliac bones

• pathologic fracture of the

multiple osteolytic lesions

(right humerus, right femur,

both shoulder blades and

subperiosteal erosions in

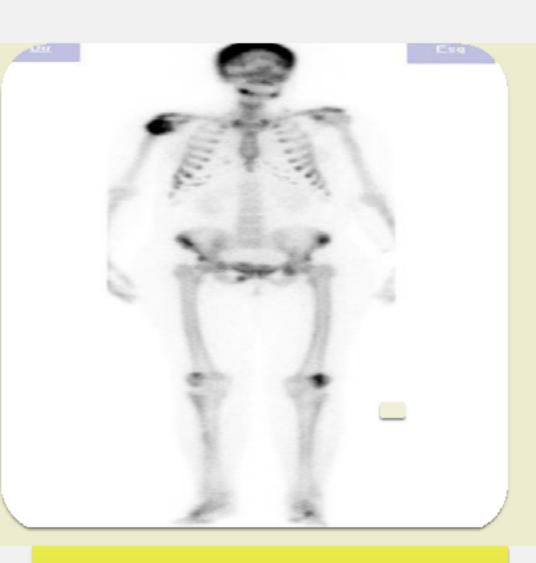
the distal phalanges

old fractures in the ribs





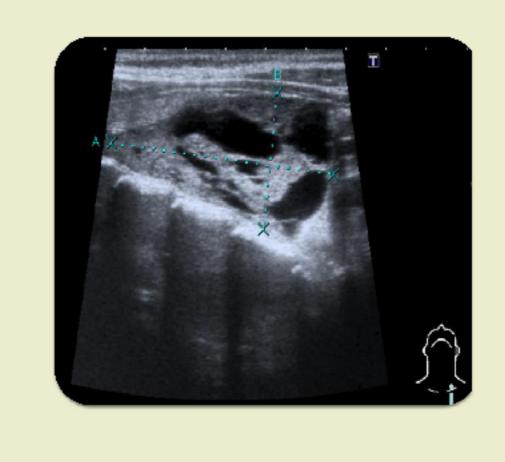
Femur fracture was managed with osteosynthesis and humeral biopsy documented multiple giant cells consistent with brown tumor of PHPT.



Bone scan showed increased uptakes over right humerus, right femur and tibias.

Analysis revealed severe PHPT Serum calcium: 3 49mmol/l

3,49mmol/L
Phosphorus: 0,53mmol/L
PTH: 1435pg/mL



Ultrasoundmixed texture

lesion in the left superior parathyroid (5,8x3,9x2,4cm)



Sestamibi scanning confirmed a left parathyroid adenoma, with a necrotic centre and bone uptakes coincident with previous bone

scan.

- Considering patient's age and high level of PTH, metastatic disease and hereditary cancer syndromes were excluded.
- Treatment was initiated with hydration and zolendronic acid and parathyroidectomy was perform. Hungry bone syndrome developed following surgery and was necessary treatment with calcium carbonate and alfacalcidol.
- Histopathology confirmed the diagnosis of parathyroid adenoma.

CONCLUSION

Pathological fractures in young adults should always be investigated. A high index of suspicion is necessary to diagnose this unusual presentation of PHPT.

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