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Case Report

Retro-tracheal parathyroid adenoma: A rare location of a common pathology [☆]

Sekhar Iyer, MD^a, Michael Simon, MD^{a,*}, Christopher Tan, MD^a, Lyle Gesner, MD^b, Joseph Viggiano, MD^c, Shalini Chhabra, MD^d

^a Resident, Department of Radiology, RWJBH - Saint Barnabas Medical Center, 94 Old Short Hills Road, Livingston, NJ 07039, USA

^b Program Director, Chief of Neuroradiology, Department of Radiology, RWJBH - Saint Barnabas Medical Center, 94 Old Short Hills Road, Livingston, NJ 07039, USA

^c Chief of Nuclear Medicine, Department of Radiology, RWJBH - Saint Barnabas Medical Center, 94 Old Short Hills Road, Livingston, NJ 07039, USA

^d Associate Attending, Department of Radiology, RWJBH - Saint Barnabas Medical Center, 94 Old Short Hills Road, Livingston, NJ 07039, USA

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ABSTRACT

Ectopic parathyroid adenoma in the mediastinum has been reported in several publications; however, its location in the posterior mediastinum, especially a retro-tracheal location, has been rarely reported. We report a case of a 61-year-old patient who presented with clinical symptoms of malignant hypercalcemia due to a retro-tracheal mediastinal parathyroid adenoma. The surgical excision normalized the phosphocalcic balance with improvement in the patient's clinical symptoms. An ectopic hypersecreting parathyroid adenoma with life-threatening hypercalcemia should prompt radiological assessment and appropriate surgical management to prevent significant clinical complications.

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Introduction

Parathyroid adenomas in the setting of hyperparathyroidism are fairly common. The typical location of a parathyroid

adenoma is in the region of the inferior parathyroid glands. Here we present a rare case of a pathologically proven retro-tracheal parathyroid adenoma, a rare and atypical location.

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* Corresponding author.

E-mail address: michael.simon2@rwjbh.org (M. Simon).

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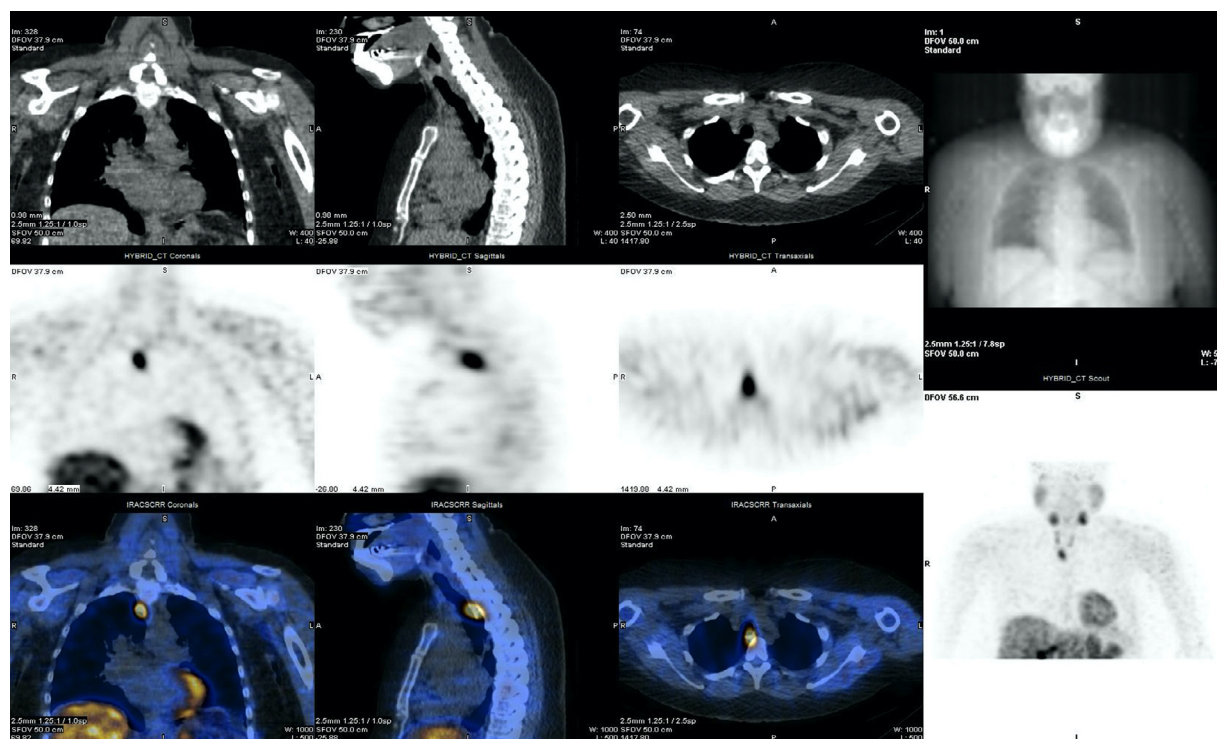


Fig. 1 – SPECT-CT following the administration of 16.0 mCi of ^{99m}Tc -Sestamibi with reconstruction in the three standard orthogonal planes demonstrates focal sestamibi uptake in the midline retrotracheal region, separate from the right lateral aspect of the esophagus.

Case report

A 61-year-old female with a history of hyperparathyroidism and hypercalcemia presented to the hospital with complaints of generalized body weakness and multiple episodes of vomiting. The patient was seen by endocrinology in 2018 but was lost to follow up. At the time of the patient's visit to the ED, serum calcium was 11.8 mg/dl, albumin 3.6 g/dl and PTH was 208.5 pg/ml. Urine creatinine and calcium were ordered and were 765.0 mg/24hr and 97.5 mg/24hr respectively. The patient was assessed with a ^{99m}Tc -Sestamibi scan as part of her workup. Focal uptake in the retro-tracheal region along the right lateral aspect of the esophagus was identified (Fig. 1). A CT scan of the neck was subsequently performed, which revealed a 2.4 × 1.3 cm enhancing prevertebral paratracheal/paraesophageal mass that corresponded to the focus of increased ^{99m}Tc -Sestamibi uptake. The patient underwent surgical exploration, and a retro-tracheal parathyroid adenoma measuring 2.5 × 1.5 × 0.5 cm was confirmed and removed. The patient tolerated the procedure well and was discharged without any complications for continued outpatient follow up. On follow up the patient's calcium level was noted to be 9.1 mg/dl and her weakness improved.

Discussion

The incidence of parathyroid adenoma in patients treated for primary hyperparathyroidism is close to 80% [1]. Parathyroid adenomas typically occur in the inferior parathyroid glands. Although ectopic localization within the mediastinum has been reported, a retro-tracheal location remains rare [2,3].

Embryologically, the parathyroid glands originate from the third and fourth branchial pouch prior to migrating to the mediastinum [4]. The superior parathyroid glands originate in the thyroid near its upper pole, directly behind or at the cricothyroid junction. The inferior parathyroids originate within the thymus and descend in a posterolateral direction to the lower thyroid pole. This longer pathway can give rise to ectopic parathyroid tissue [5].

As the tissue migrates, cells can be left behind along its migration path and can subsequently develop into functioning parathyroid tissue. This is more common than some might think. In a case series of 231 patients, 16% were found to have ectopic parathyroid tissue [6]. As the migration path is in the anterior mediastinum, it is extremely rare to find tissue outside of this area [7].

Nuclear Medicine Scintigraphy is currently considered the technique of choice for suspicion of parathyroid adenomas due to its high sensitivity (95%) and better image quality than CT with a lower radiation dose delivered to the patient [8,9]. In the Mitchell Report, scintigraphy identified 16/17 cases of adenomas and 19/21 cases of glandular hyperplasia [10].

Complete resection of the tumor or ectopic tissue is the primary therapeutic procedure to alleviate the patient's symptoms. Surgery is recommended for both symptomatic and asymptomatic patients due to the significant morbidity associated with the disease [11–14]. In symptomatic patients, parathyroidectomy leads to normalization of phosphocalcic metabolism and an increase in bone density [15].

The most common postoperative complication is hypocalcemia. The absence of hypocalcemia suggests incomplete resection of the ectopic tissue. Persistent hypocalcemia more than 4 to 5 days after the surgery is often due to an avidity of bone for calcium, requiring phosphocalcic supplementation and vitamin D therapy [16].

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