1. Introduction

- Sacrococcygeal teratomas [SCT] are extremely rare in adults. We present a 65-year-old male with a possible SCT with de-differentiation into thyroid cancer treated with high level ablation with 202 mCi I-131 NaI.
- This is the first documented case of ablation therapy for a SCT, and the second documented case of malignant thyroid tissue arising from a SCT. Extensive literature review including information by Monteiro (1) Head et al(2), Ahmed and Pollock (3), Ng et al (4), and Bull et al (5) reveals 120 cases of adult SCTs with only one of which de-differentiated into thyroid tissue.
- This case signifies the importance of considering SCT, which is often misdiagnosed, in the differential diagnosis for a pelvic mass presenting with neurologic and/or obstructive symptoms.

2. Methods

- Extensive review of the literature was performed, including Ovid, Medline and Pub Med databases, searching for adult SCTs and their treatment.

3. Patient Description

- 65 year old male with a 2-3 year history of right hip pain radiating to right leg, not relieved by conservative measures.
- Initial work up revealed thyroglobulin quantity of >3000.
- At 3 month post ablation, patient still has mild leg pain controlled with OTC medications, and no evidence of recurrence of the thyroid tissue.
- Initial work up revealed thyroglobulin quantity of >3000. Subsequently, an CT of the pelvis and lumbar spine (image 1) and 1-131 whole body scan followed (see image 2).
- The location of the SCT was very difficult to excise surgically so we decided to perform high level ablation using 202mCi-131 NaI. The thyroid was removed due to fears of swelling following NaI therapy leading to airway compromise. Pathological examination of thyroid revealed no evidence of neoplasm.
- At 3 month post ablation, patient still has mild leg pain controlled with OTC medications, and no evidence of recurrence of the thyroid tissue.

4. Discussion

- 65 year old male with a 2-3 year history of right hip pain radiating to right leg, not relieved by conservative measures.
- SCTs are the most common type of germ cell tumors diagnosed in neonates, infants, and children younger than 4 years. SCTs occur at a rate of 1 in 35,000-45,000, and more often in girls than in boys; ratios of 3:1 to 4:1 have been reported (2).
- SCTs are thought to originate from multipotential cells in Henson's node, which migrates caudally to rest in the coccyx.
- Unlike infant teratomas which usually are externally visible, adult teratoma are usually confined to intrapelvic areas.
- Teratoma can present with complaints of constipation, rectal obstruction, recurrent urinary tract infections, neurologic symptoms such as lower extremity weakness and/or numbness, and pain (1).

5. Conclusions

- SCTs should be included in the differential diagnosis, whatever the nature of the lesion is (2-5).
- Reported cases of SCTs in adults are rare. Reviews of the literature demonstrates fewer than 120 SCTs, with only 20 cases described as malignant. Only 1 previously reported case suggests thyroid carcinoma arising from SCT. In all adult SCT, surgical excision was the treatment of choice (4-6).
- Newer protocols suggest that cisplatin, bleomycin, vinblastine, and radiotherapy, in addition to surgical excision, may improve the prognosis of malignant teratomas (1,10).

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6. Work Cited