ABSTRACT POSTER

NOME PRIMO AUTORE: Elena

COGNOME PRIMO AUTORE: Castellano

Età: 27 anni

SEDE: S.C. Endocrinologia e Metabolismo, A.S.O. S. Croce e Carle, Cuneo

NOME COAUTORE: Sara

COGNOME COAUTORE: Cassibba

SEDE: S.C. Endocrinologia e Metabolismo, A.S.O. S. Croce e Carle, Cuneo

NOME COAUTORE: Flora

COGNOME COAUTORE: Cesario

SEDE: S.C. Endocrinologia e Metabolismo, A.S.O. S. Croce e Carle, Cuneo

NOME COAUTORE: Fabrizio

COGNOME COAUTORE: Giordano

SEDE: S.C. Anatomia Patologica, A.S.O. S. Croce e Carle, Cuneo

NOME COAUTORE: Laura

COGNOME COAUTORE: Gianotti

SEDE: S.C. Endocrinologia e Metabolismo, A.S.O. S. Croce e Carle, Cuneo

NOME COAUTORE: Claudia

COGNOME COAUTORE: Baffoni

SEDE: S.C. Endocrinologia e Metabolismo, A.S.O. S. Croce e Carle, Cuneo

NOME COAUTORE: Giorgio

COGNOME COAUTORE: Borretta

SEDE: S.C. Endocrinologia e Metabolismo, A.S.O. S. Croce e Carle, Cuneo

TIPOLOGIA: POSTER

ARGOMENTO: Caso clinico

TITOLO: Sclerosing mucoepidermoid carcinoma with eosinophilia of the thyroid: a case report and review of the literature.

INTRODUZIONE: Sclerosing mucoepidermoid carcinoma with eosinophilia of the thyroid (SMECE) was first described as a low-grade carcinoma, generally occurring in a background of Hashimoto's thyroiditis. So far, less than 30 cases have been described in literature. Most cases manifested an indolent clinical course despite contiguous adenopathy and soft-tissue extension; metastasis has been described as an unusual manifestation. However some patients give a history of recent rapid enlargement and, rarely, may present with symptoms resulting from extrathyroidal extention. We report a case of SMECE in a woman with Hashimoto's thyroiditis and a thyroid node known for over 10 years.

METODI: Clinical and biochemical data of the patient are presented and the pertinent literature is reviewed.

RISULTATI: A 70years old woman comes to our center for evaluation of severe osteoporosis in DMT1 with microangiopathy and Hashimoto's thyroiditis. She had hypothyroidism and a thyroid nodule known at least since 2000,reported of stable size at US follow-up. She had not history of neck irradiation and her family did not indicated history of thyroid cancer. In 2012, at US evaluation, the left lobar node appeared hypoechoic, heterogeneous, non-vascularised, with irregular margins,37mm in diameter, stable compared to previous evaluations. Plasmatic calcitonin was normal. A fine-needle aspiration on this node resulted Thy3. She underwent left thyroidectomy with a histological diagnosis of SMECE. The proliferation showed foci of perithyroidal extension reaching the margins of surgical excision. A subsequent right lobectomy and recurrent lymphonode dissection was performed. TNM: pT3NO. One month after surgery the total body CT-PET showed a tracer accumulation (14mm, suv/max 3.3) in the neck, confirmed on MRI. The patient was addressed to nuclear medicine, where ablative radio metabolic therapy with I131 has been programmed.

CONCLUSIONI: SMECE is a rare malignant neoplasm that often develops in the context of an autoimmune thyroiditis. The differential diagnosis with undifferentiated or squamous carcinoma, mucoepidermoid carcinoma and nodular tumor-like squamous metaplasia could be difficult. It's an aggressive cancer that can have indolent clinical course. Optimal treatment and follow-up are yet to be determined.