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Sclerosing stromal tumor of the ovary with masculinization, Meig's syndrome and CA125 elevation in an adolescent girl: A case report

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Abstract

BACKGROUND

Sclerosing stromal tumor (SST) is an extremely rare sex cord stromal tumor of the ovary. It was first reported and named by Chalvardjian *et al*[1] in 1973. These tumors typically present with pelvic/abdominal pain and tenderness, a mass, and/or abnormal menses, but rarely present with masculinity in children and adolescents. Only 2 cases of these tumors have been reported in premenarchal girls, who demonstrated hormonal activity, with a history of the development of a virilizing female due to

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A sclerosing stromal tumor of the ovary (SST) is an extremely rare benign solid ovarian **tumor**, which is derived from the ovarian stroma and is a distinct subtype of sex cord-**stromal tumor**.

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Cited by: 240 **Author:** Ara Chalvardjian, Robert E. Scully

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