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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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Author: Ara Chalvardjian, Robert E. Scully

Publish Year: 1973

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In this report, we describe the association between **sclerosing stromal tumor of the ovary**, **Meigs'**



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