Introduction

Ovarian teratomas, also known as dermoid cysts, are the most common ovarian germ cell tumors. Teratomas can be divided into two histological subgroups: Mature and immature teratomas. Immature Ovarian Teratomas (IOTs) are a histological subtype that represent less than 1% of all ovarian cancers [1-3] and account for 10% to 20% of ovarian cancers in women under 20 years of age [3].

Growing Teratoma Syndrome (GTS) is a rare clinical entity that entails the appearance of large intraabdominal or thoracic masses of mature teratoma cells after systemic chemotherapy for the treatment of germinal cell tumors with immature components. We present the case of a 15-year-old patient who had an immature teratoma in the left ovary treated with surgery and chemotherapy and posterior mature teratomas in the lungs and contralateral ovary.

Case report

A 15-year-old patient with no personal or family medical history of interest presented to the emergency department complaining of gastrointestinal symptoms with pain in the left...
An exploratory laparotomy was conducted which found a large mass consisting of solid parts mixed with tumoral cysts of a whitish-yellow color and soft consistency. The tumor was removed completely by means of a left adnexectomy. The histopathological analysis revealed an ovarian mixed germ cell tumor with 90% immature components and 10% vitelline yolk sac tumor. The patient was referred to the fertility unit in order to preserve fertility. At that time, fertility preservation was dismissed due to the malignant histology of the tumor and the need to start chemotherapy urgently.

The patient received four cycles of a Bleomycin, Etoposide, and Cisplatin (BEP) chemotherapy regimen. On a PET-CT scan five months after completing treatment, pulmonary nodules were detected. With a suspicion of metastasis, the nodules were removed via a thorascoscopy. Microscopically, the nodules from the left upper lobe were found to be mature teratoma. On a transvaginal ultrasound two months later, a new cyst in the right ovary was observed. The image showed a unilocular cystic mass measuring 24 x 19 mm with a solid papillary mass inside that measured 10 x 9 mm, Doppler score 1, with healthy surrounding ovarian tissue. An MRI test revealed a cystic formation in the right ovary measuring 44 x 33 x 36 mm that contained a solid papillary mass inside. No macroscopic fat component was observed (Figure 2).

The patient was again referred to the fertility unit to plan a fertility preservation strategy due to the risk of losing the right ovary. Even though growing teratoma syndrome was suspected and there was a high probability that this new cyst was also a mature teratoma, it was still possible that it was a metastasis of the original tumor. The patient was in amenorrhea secondary to the chemotherapy received and the remaining ovarian tissue, the patient was offered the possibility of hormonal stimulation to vitrify oocytes in order to cryopreserve healthy ovarian tissue. In the end, a cystectomy was possible and the histopathological analysis found it to be a mature teratoma.

Six months after the first surgery, an ovarian reserve study was conducted. An antral follicle count of two in the right ovary and an AMH 0.16 ng/mL were found. At this point, the patient was in amenorrhea and surgery on the right ovarian cyst was pending. Therefore, fertility preservation was postponed.

At present, the patient is well after one year of post-surgical follow-up. No residual tumors have been detected through regular examinations with ultrasounds and scans. The patient regained normal menstrual cycles and another ovarian reserve study found an antral follicle count of nine in the right ovary and an AMH of 1.46 ng/mL.

Given the possibility of a recurrence of ovarian teratomas in the remaining ovarian tissue, the patient was offered the possibility of hormonal stimulation to vitrify oocytes in order to preserve fertility. The patient was informed about the prognosis secondary to the chemotherapy received and the remaining tissue in the right ovary. She accepted the treatment and at present, she is waiting for her period in order to start ovarian stimulation.

**Discussion**

Benign teratomas are believed to arise from primordial germ cells in the fetus. They are distributed along the lines of migration from the embryonic yolk sac endoderm to other tissues such as the primitive gonads during early embryogenesis. Although these tumors are mainly located in the ovaries, extragonadal tumors can appear anywhere along germ cell migration routes and account for 2% to 5% of all germ cell tumors [4,5]. As they follow migration routes, these tumors typically grow in middle structures like the retroperitoneum, brain, thymus, lungs, liver, and uterus [6].

Growing Teratoma Syndrome (GTS) was first described by Logothetis et al. in 1982. This syndrome leads to the appearance of large masses of metastatic mature teratoma in diverse locations during or after chemotherapy in patients with immature germ cell tumors [7]. The diagnosis of GTS is based on three main criteria: personal history of germ cell tumor, radiological
evidence of growing lesions during or after chemotherapy, and normal tumor markers [8]. Surgery is the primary treatment and the outcome depends on the size and location of GTS [9]. Our patient met all the three criteria. A differential diagnosis was made between recurrence of the immature teratoma and GTS. In this case, the main suspicion was GTS, as recurrences of immature teratomas often appear as larger masses with prominent solid components.

Although an ovarian location is infrequent, these patients have a high risk of infertility due to multiple abdominal interventions that reduce the ovarian reserve and the secondary formation of intraabdominal adhesions [10,11]. In our case, the patient had two important risk factors for developing infertility in the future: treatment with gonadotoxic chemotherapy and multiple ovarian surgeries.

The increase in survival rates of young patients with cancer, new reproductive techniques, and a growing interest in quality of life after gonadotoxicity in cancer survivors have highlighted the importance of fertility preservation in young patients [12,13]. Different techniques can be used in order to preserve fertility in oncological patients. The cryopreservation of oocytes and embryos is the standard procedure. However, these procedures require ovarian stimulation for eight to 12 days, which delays the start of oncological treatment. In addition, embryo cryopreservation raises serious ethical dilemmas given that in some oncological patients, the prognosis of the disease is guarded in terms of survival and the technique requires a male partner or the use of sperm from a donor. For this reason, the main technique used is oocyte vitrification. This procedure also requires the patient to have menstrual cycles and most authors consider the upper age limit to be between 35 and 40 years [12]. In prepubertal patients or patients who need to start chemotherapy urgently, cryopreservation of ovarian tissue is the only alternative, but it continues to be an experimental technique [13,14].

In this case, oocyte cryopreservation was the first choice for fertility preservation, but was conditional upon the histological nature of the ovarian tumor. Cryopreservation of ovarian tissue was also a possibility in the event of major difficulty in the surgical approach that would have led to requiring a total adnexitomy instead of a cystectomy.

Conclusion

The appearance of clinical or radiological masses in patients treated with chemotherapy for malignant germ cell tumors should alert clinicians to the possibility of GTS. The early identification and surgical treatment of the syndrome improves the prognosis. Fertility preservation in young patients must be carefully evaluated, as the risk of secondary infertility is high given the exposure to gonadotoxic drugs and multiple gynecological intervention that may lead to a decrease in ovarian reserve.

Declarations

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References