

Ovarian Torsion after Ovarian Hyperstimulation Syndrome

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Abstract

We here report two case of ovarian torsion We checked the presence of all possible factors that could explain the onset of the syndrome. Ovarian hyperstimulation syndrome (OHSS) is almost exclusively associated with ovulation induction with gonadotropins[3] OHSS continues to be a serious complication of assisted reproductive therapy (ART), with no universally agreed upon best method of prevention If OHSS is diagnosed, the etiology should be determined in order to focus the treatment and avoid future complications.



INTRODUCTION

Ovarian hyperstimulation syndrome (OHSS) is rather frequent (1–5%) in women submitted to superovulation with gonadotropins for in vitro fertilization (IVF). Multiple pregnancy, gestational trophoblastic disease, primary hypothyroidism, thyroid-stimulating hormone/ gonadotropin-secreting adenomas, and mutations of the FSHR gene may trigger spontaneous OHSS.[1,2]

PURPOSE

The aim of the study was to investigate the incidence, progress, management, and outcome of adnexal torsion after controlled ovarian hyperstimulation in embryo transfer cycles.

PRESENTATION OF CASE

A 25-year-old woman (Gravida 0 Para 0) with first infertility of 2-year duration with bilateral polycystic ovaries and timed intercourse came for further treatment to our clinic. Her body mass index was 20.

She was planned for controlled ovarian stimulation for IVF and was given short protocol. Stimulation with urinary FSH was started on day 2 of menstrual cycle, and starting Recagon dose was 225 IU s/c for 8 days. Moreover, antagonist was started on day 7 of menstrual cycle. Decaptople 0/1 subcutaneous was given for oocyte maturation. On day 12, there were 15 dominant follicles in the left and right ovary. Oocyte retrieval was done on day 14 of cycle. She came with the sudden onset of severe right- and left-sided

abdominal pain with increasing severity in the next 2 h.

On the day of pick up, rotation occurred.

She had no medical or surgical history. On examination,

she had tachycardia +108/min, blood pressure – 110/80,

and respiratory rate –18/min. She had moderate abdominal

distension, and severe right- and left-sided guarding. Abdominal ultrasound one suggested enlarged bilateral

ovary - right 14 × 10 cm and left 8 × 9 cm with multiple

cysts, fluid in paracolic gutter, and Morison’s pouch. She

was diagnosed with severe OHSS. Doppler study was not

conclusive about torsion. Pain decreased with injection

tramadol. With the further episode of severe abdominal pain

after 2 h of admission, decision was made for emergency

laparoscopy.

Ultrasonography showed that bipolar ovaries of normal

diammonotic and large ovaries with thick blood flow in the

left and right ovary, in the abdominal cavity indicative of

hemoperitoneum, were shown. Emergency laparoscopy was done and bilateral ovarian torsion with retained vascularity was noted. Biochemistry and thyroid function were normal.

Laparoscopy for suspicion of OHSS complicated by ovarian torsion confirmed bilateral ovarian enlargement and hemoperitoneum.[5] The left adnexa was twisted around the stem and an ischemic adenoectomy was performed and two liters of blood was sprayed. Later, anatomopathological examination confirmed ovarian ischemia.[5,7]

The pathogenesis of OHSS involves hyperstimulation of FSHR by FSH or other glycoproteins with similar structures. Recently, mutations of the FSHR gene leading to an increased sensitivity of the receptor to normal levels of hormones have been also described, although a standard genetic test has yet to be developed. Several circumstances may increase the levels or biological activity of these glycoproteins.[6,3]

CASE REPORT
Woman 28years old (gravida 0 para 0). with first infertility of 1years duration with top normal ovaries and timed intercourse came for further treatment to our clinic. Her body mass index was 19/8. She was planned for controlled ovarian stimulation for IVF and was given short protocol. Stimulation with urinary FSH was started on day 3 of the menstrual cycle, and starting recagon dose was 225 IU s/c for 8 days.

And antagonist was started on day 7 of menstrual cycle.

Decapeptide 0/1 subcutaneous was given for oocyte maturation.

On day 10 there were 20-30 dominant follicles in left&right ovary.

Oocyte retrieval was done on day 12 of cycle.

She came with sudden onset of severe right-left sided abdominal pain with increasing severity in next 2 hours.

On the day of pick up rotation occurred.

The left ovary torso, which had been rotating, turned open.

CONCLUSION

OHSS can be associated with life-threatening complications that require early diagnosis for successful management. The etiology should be determined in order to focus the treatment and avoid future complications.

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