Heterotopic triplet pregnancy after in vitro fertilization and intracytoplasmatic sperm injection complicated with ovarian hyperstimulation syndrome single twin fetal demise and cervical incompetence: a successful obstetrical outcome

I. Korkontzelos¹, A. Vlachioti¹, P. Mavridou², A. Rapi¹, G. Tsanadis³, T. Stefos³

¹Department of Obstetrics and Gynecology, ²Department of Anesthesiology, Ioannina State General Hospital “G. Chatzikosta”, Ioannina
³Department of Obstetrics and Gynecology, University Hospital of Ioannina, Ioannina (Greece)

Summary

Spontaneous heterotopic triplet pregnancy (HTP) was formerly considered to be extremely rare. Recently, the broad use of assisted reproductive techniques to assist infertile couples, has contributed to the constant rise of non-spontaneous heterotopic triplets. A case of 38-year-old woman who underwent in vitro fertilization (IVF) and intracytoplasmatic sperm injection (ICSI) is presented. A series of complications appeared and were treated successfully. Although this situation could be life threatening and should alert the medical community, misdiagnosis is still present. The clinicians need to be aware that after IVF procedures, one or more complications might occur. In those cases, early diagnosis and surgical or conservative intervention is of paramount importance.

Key words: Heterotopic triplet pregnancy; In vitro fertilization; Ovarian hyperstimulation syndrome; Vaginal ultrasound.
form, the pregnant woman was transferred to the operating theater where an exploratory laparotomy was performed under regional (spinal) anesthesia. Diagnostic laparoscopy is considered a contraindication in cases of OHSS and it was excluded. Hemoperitoneum with a ruptured left tubal pregnancy was noted and left salpingectomy was performed (Figure 4). Histopathological examination verified the initial diagnosis. The patient received blood and plasma transfusion, recovered gradually due to OHSS, and was discharged on the eighth day in good condition. At 13 weeks of gestation, she underwent the first trimester screening test for Down syndrome (nuchal translucency) and cervical cerclage. Although discussed but not suggested, the patient decided to use vaginal progesterone 200 mg once a day. At 22 weeks of gestation, she underwent the second trimester scan and the non-invasive blood test for chromosomal abnormalities. At 30 weeks of gestation she was re-admitted due to threatened preterm labor. At 36+1 weeks of gestation a cesarean section was performed due to cervical effacement and a normal female neonate of 1,980 grams was born. One year follow up of the neonate showed normal physical and mental development.

Discussion

The incidence of HP formerly considered to be rare, continues to be so, but is increasing steadily in recent years. The same applies also for HTP. It is remarkable that the first case was reported in the literature in 1991 by Goldman et al. [3]. Heterotopic triplets should be considered a part of the ectopic-heterotopic medical condition, however diagnosis and treatment of HTP appears difficult and peculiar.

In general, the most common sites of HP in descending order occur at the fallopian tube, the cornu, the cesarean scar, the cervix, and the abdomen [4]. It is reported that approximately 70% of HTP are diagnosed between five and eight weeks of gestation, 20% are diagnosed between nine and ten weeks, and the remaining 10% are diagnosed after 11 weeks [5]. Tubal cases tend to be diagnosed earlier because the confined space results in earlier pain or rupture. Factors predisposing to HP are similar to those predis-
posing to ectopic pregnancy, such as: a large number of transferred embryos, embryo transfer near the uterine horn, excessive pressure on the syringe and deep insertion of the catheter during transfer, the quality of embryos, the use of gonadotropins, the amount of fluid used as media for the embryos, and also adhesions related to endometriosis and PID [6, 7].

The diagnosis of HP is challenging. Serial β-hCG levels are not useful due to the concomitant intrauterine pregnancy. Transvaginal ultrasound is the imaging modality of choice for the diagnosis of tubal and non-tubal HPs, however it continuous to have low sensitivity because it is often missed or overlooked [8]. Li et al. in a retrospective study set certain sonographic criteria so as to diagnose HP using TVS, even if a visible intrauterine gestational sac was observed. These criteria were (i) an inhomogenous adnexal mass, (ii) an empty extraterine gestational sac seen as a hyperechoic ring, (iii) a yolk sac, and/or a fetal pole with or without cardiac activity in an extra-uterine sac. The sensitivity and specificity of TVS for the detection of HP in this study was reported to be 92.4% and 100%, respectively [9].

The diagnosis of HP is becoming more complicated in cases of ruptured ones associated with OHSS. Both medical conditions could be associated with hypotension, tachycardia, abdominal pain, and free fluid in the abdominal cavity with the simultaneous presence of an intrauterine pregnancy. Ruptured corpus luteum could also be misdiagnosed as HP since both conditions may exhibit the “ring fire” sign on ultrasound. On the contrary, women with OHSS usually have normal hemoglobin and high hematocrit and enlarged ovaries [8, 10].

The management of HTP aims to remove the ectopic pregnancy without disturbing the viable intrauterine embryo or embryos. Currently, there is no consensus on the treatment of HTP and the majority of data derive from case reports. Treatment depends on the gestational age at diagnosis, the clinical status of the patient, the site of the ectopic implantation, and the experience of the treating physician. HP can be treated by a conservative or surgical approach. Laparotomy and removal of the ectopic gestation is the preferred technique in ruptured cases and in hemodynamically unstable patients. Laparoscopic approach could be considered when there are no contraindications. Salpingectomy, is preferable for tubal pregnancy in patients with uncontrolled bleeding or extensive tubal damage. Salpingostomy, is indicated when the patient is stable, when there is an unruptured ectopic sac < 5 cm, or when the contralateral tube is absent or damaged or the patient strongly wishes to conserve fertility [11, 12].

Conservative treatment requires early diagnosis. Agents like methotrexate (MTX), potassium chloride (KCL), and hyperosmolar glucose could be used. The use of MTX with a viable pregnancy is contraindicated as the drug could cause miscarriage or congenital malformations. Sonographically guided local injection of KCL into the heart of the ectopic fetus can induce cardiac asystole. Hyperosmolar glucose could also be injected into the gestational sac causing local dehydration, necrosis, and resolution of the trophoblastic tissue. Cornual and interstitial pregnancies can be treated by surgical excision or ultrasound guided aspiration and injection of KCL. Cervical HP could be treated by ultrasound-guided aspiration with cervical sutures, intracardiac injection of KCL, vacuum aspiration after ligation of the descending cervical branches of the uterine arteries, electrodessication, and hysteroscopic resection. In cases of ovarian ectopic pregnancy laparoscopic wedge resection or ipsilateral oophorectomy is the treatment choice [2, 13].

In cases of single twin fetal demise, the chorionicity of the multiple pregnancy and the gestational age at which fetal demise occurs are the two most important factors for the survival of the second twin [14]. In monochorionic twins the perinatal survival rate is reported 83%, while in dichorionic twins the perinatal survival rate is reported 100% [15].

There has been a lot of discussion concerning the role of cervical cerclage and vaginal progesterone for the treatment of cervical incompetence. In a systematic review, Conde-Aguedelo et al. stated that in singleton pregnancies with sonographic short cervix in the second trimester and previous spontaneous preterm birth, either vaginal progesterone or cerclage, are equally efficacious [16]. On the contrary, Wang et al. noticed that cerclage is more beneficial than progesterone [17]. The same author mentioned that the effectiveness is similar when vaginal progesterone is added to the cervical cerclage. In another article, cerclage, vaginal progesterone, and cervical pessary appear to have similar effectiveness, while the use of single or combined therapy is still under discussion [18].

**Conclusion**

The present case report confirms the importance of the early diagnosis of HP for a successful obstetrical outcome. Additionally, it emphasizes the implication of large number of embryos transferred in utero contrary to the current guidelines. Searching the literature, the authors found only one similar report with heterotopic triplets complicated by OHSS following IVF, where it is noted that enlarged ovaries and free fluid in the abdominal cavity could mask early accurate diagnosis [8]. In the present case, two parameters led us to early diagnosis although the patient was referred as OHSS syndrome in a stable situation, the relatively low hematocrit and the mass noticed in the left adnexa after thorough vaginal ultrasound. Another case referred as OHSS in a stable condition with low hematocrit and a sac in the right adnexa is also reported and treated [19].

Based on the present authors’ experience, they suggest
serial early vaginal scans in all pregnancies induced by ART every week, starting from the 5th until the 9-10th week of gestation. Furthermore, they strongly agree with Mustafa et al. who noticed that in cases of pregnancy and ART the first thought coming in mind should be “think ectopic” [19] and they would further add “think heterotopic”. Obstetrical complications treatment strategy if present, should take into account clinicians experience and woman’s preferences.

References


Corresponding Author:
I. KORKONTZELOS, M.D.
Department of Obstetrics and Gynecology
Ioannina State General Hospital “G. Chatzikosta”
Ioannina (Greece)
e-mail: i.korkontzelos@yahoo.com