Cervical Ectopic Pregnancy of 45,XO Embryo After In Vitro Fertilization/Embryo Transfer
İn Vitro Fertilizasyon/Embriyo Transferi Sonrası 45,XO Embriyonun Servikal Ektopik Gebeliği

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We report a case of anembryonic pregnancy located in the cervico-isthmic region of the uterus following in vitro fertilization (IVF)/embryo transfer (ET) treatment cycle. Cervical pregnancy was diagnosed during 7 weeks of gestation and treated with successful aspiration and curettage. Cytogenetic analysis of abortus material was consistent with 45,XO, Turner syndrome. Early diagnosis of cervico-isthmic pregnancy is important and critical for effective conservative management.

Key Words: Ectopic pregnancy; 45 XO; Turner syndrome; in vitro fertilization

ÖZET


Anahtar Kelimeler: Ektopik gebelik; 45 XO; Turner sendromu; in vitro fertilizasyon

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Cervical pregnancy is usually considered as a life threatening event. It is a rare form of ectopic gestation and accounts for less than 1% of all pregnancies.1 Cervical pregnancy was defined as implantation of the gestation in the endocervical canal. The incidence is approximately 1 in 2500-12000 pregnancies.2 It is a well known complication of ART procedures. Early diagnosis is important and allows conservative treatment option to preserve fertility. Hysterectomy was the only treatment for cervical pregnancy in the past.2 Current treatment modalities include aspiration curettage with hypogastric/uterine artery embolization, curettage and local prostoglandin injection, insertion of foley catheter in the cervical canal after curettage, local administration of methotrexate (MTX) with/without local potassium chloride, systemic administration of MTX and hysteroscopic resection of the gestation in conjunction with uterine artery embolization.3,5 Early diagnosis is important to prevent serious complications such as
massive hemorrhage. We present a case with cervical ectopic pregnancy following IVF/ICSI and ET in the current report.

**CASE REPORT**

A 28 year old woman was referred to our center for secondary infertility. She was married for 8 years. The etiology of infertility was male factor with severe oligospermia. Her first spontaneous pregnancy was terminated with dilatation and curettage for missed abortion at 6 weeks of gestation a year ago. The patient underwent IVF/ICSI cycle. 10 oocytes were retrieved and 9 of them were mature. ICSI was performed to all oocytes. Embryo was transferred under abdominal ultrasound guidance. Initial serum hCG, progesterone and E2 levels, 14 days after embryo transfer were 49 mIU/ml, 60 ng/ml and 926pg/ml, respectively. Daily progesterone vaginal gel and weekly hCG 5000 IU ampoule intramuscularly were prescribed. Six days after the initial hCG test, the level raised to the 349 IU/ml. Serum hCG, progesterone and estradiol levels were measured every 3-7 days (Figure 1). The serum hCG levels increased moderately.

Transvaginal ultrasound (TVUSG) examination 4 weeks after ET, demonstrated an empty uterine cavity with a 9 mm gestational sac located in the cervicoisthmic area and serum hCG level was 1861 IU/ml. Follow up ultrasound examination after one week was consistent with slightly growing anembryonic gestational sac in the same localization. 10x18 mm gestation sac in the cervikoisthmic localization was observed by TVUSG examination in 7 weeks and 4 days of gestation. The yolk sac and embryonic pole were not visible. Doppler sonography also revealed trophoblastic vessels in the same area (Figure 2a and 2b). Confirmation of cervical ectopic pregnancy was informed and treatment options were discussed with the patient and her family. Dilatation and aspiration curettage was successfully performed under ultrasound guidance.

The procedure was uneventful, per and post-operative abnormal bleeding was not observed. The abortion material was sent for cytogenetic analysis. The patient was discharged from the hospital the following day, without complication. Follow up ultrasound examination one week after the procedure demonstrated a normal cavity and cervical
canal (Figure 2c). The cytogenetic analysis was consistent with 45 XO Turner Syndrome. Patient’s peripheral blood karyotype analysis from peripheral blood also demonstrated mosaicism involving normal and 45,XO cells (45 X[4%]/46 XX[96%]). The informed consent was obtained from the patient for the current case report.

DISCUSSION

Several risk factors for ectopic pregnancies are prior dilatation and curettage, cesarean section and pelvic inflammatory disease. There is an overall increase in the incidence of cervical and other sites of ectopic pregnancies with the advent of specific ART technologies. One of the explanations for this condition focus on the the embryo transfer techniques. Insertion of the transfer catheter into the uterine cavity under ultrasound guidance is suggested to make sure the tip of the catheter is in the right place. Alterations in hormonal milieu and endometrial receptivity secondary to controlled ovarian stimulation are proposed as risk factors for ectopic pregnancies after IVF. A history of tubal factor infertility and pelvic inflammatory disease are significant risk factors for ectopic pregnancies after IVF.

Although the most common site for ectopic pregnancies are uterine tubes, unusual locations such as cornual, isthmic or cervical sites can be observed. Serial serum hCG and TVUSG monitoring after IVF, leads to early diagnosis of the abnormal location and development of the pregnancy. This approach also provides a chance for conservative management especially when the diagnosis is made in early first trimester. Dilatation and curettage either with or without cervical balloon tamponade, hysteroscopic resection, embolization of uterine arteries, systemic or local metotrexate therapies are fertility conserving procedures. Serial monitoring of serum hCG and TVUSG allowed early diagnosis of anembryonic cervicoisthmic pregnancy and treatment by successful dilatation and curettage without bleeding, in the current case. Moreover the genetic analysis of the abortus material revealed 45, XO Turner syndrome. The proportions of chromosomal abnormalities in ectopic pregnancies are as high as those in early spontaneous abortions. Embryos with genetic abnormalities may result with abnormal implantation of gestations. Following the diagnosis of 45,XO of the gestational material, the patient’s karyotype analysis from peripheral blood also revealed mosaicism involving normal and 45,XO cells (45 X[4%]/46 XX[96%]) in the current case.

In conclusion, early diagnosis of cervicoisthmic pregnancies enables conservative management of patients especially following the ART procedures. Embryonic genome may also contribute to the etiology of ectopic pregnancies. We suggest genetic testing of the ectopic pregnancy material if possible, especially after ART treatments.
REFERENCES