Spontaneous heterotopic pregnancy with live birth – a case report, clinical and ultrasonographical aspects

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Introduction

Heterotopic pregnancy is a rare pathology, in which an intrauterine and an extraterine pregnancy coexist. It can be a life-threatening condition (Hassani 2010). The incidence is approximately 1 in 30000 spontaneous pregnancies (Lee 2016). The incidence in cases of medical assisted reproduction is 1-3 in 100 pregnancies (Levine 2007). The incidence has increased in recent years as a consequence of the more frequent use of medical assisted reproduction techniques (Li 2016). The risk factors for heterotopic pregnancy are the same as those for ectopic pregnancy (Rasuli 2020). A history of pelvic inflammatory disease, prior tubal and pelvic surgery, as well as tubal damage can increase the incidence of heterotopic pregnancy (Karkoe 2019). Heterotopic pregnancy presents with no specific symptoms, thus early diagnosis is difficult. The symptoms are similar to those of ectopic pregnancy, adnexal torsion or cyst and tubo-ovarian abscess; therefore, every fertile patient presenting with amenorrhea, acute lower abdominal pain, peritoneal irritation or signs of hypovolemic shock should be suspected of ruptured ectopic or heterotopic pregnancy (Li-Ping 2018). The differential diagnosis can be made with a bicornuate uterus with gestation in both cavities, or intra-uterine gestation with hemorrhagic corpus luteum (Stanic 2020).

Case report

Patient ETM, age 31, presented in the emergency department of our hospital with acute abdominal pain, mild vaginal bleeding and 8 weeks of amenorrhea. She had a medical history of repeated pneumothorax.

The clinical exam revealed tenderness in the lower abdomen, of higher intensity in the left iliac fossa. The vaginal exam revealed mild bleeding with fresh blood, tenderness in the left vaginal sack and enlarged uterus. Rectal exam revealed tenderness. The ultrasonographic exam showed a normal intrauterine pregnancy, with a normal embryo, a crown-lump length (LCC) of 4 mm (Fig. 1) and a fetal heart rate (FHR) of 157 b/min (Fig. 2), without any signs of hematoma.

On the left side of the uterus the ultrasonography revealed the left ovary with the corpus luteum and a lateral-uterine mass of 43.5/38.4 mm, with inhomogeneous content resembling an embryo (Fig 3). A Pulse Wave Doppler exam revealed a positive embryonic cardiac activity. The Douglas sack had a fine hypoechogenic line interpreted as normal liquid.

Regarding the patient history, the anaesthesiology team recommended an open surgical approach and spinal anesthesia. We performed a Pfannensteil laparotomy that revealed the normal pregnant uterus, a dilated left fallopian tube, a yellow body located on the left ovary and a small amount of blood intraabdominally. We performed a left salpingectomy. The fallopian tube was sent to the pathology department for histopathological examination.

The patient had a good postoperative recovery. She was administered 200 mg of progesterone intravaginally as supportive therapy for the intrauterine pregnancy. Dismissal was in the third day. The macroscopic evaluation was performed after fixation in 10% Formaldehyde. The investigated tissue consisted of fallopian wall with intraluminal haematic material.
an intrauterine pregnancy as well as the characteristic lateral-uterine mass with a gestational sac (Petrides 2015).

Human chorionic gonadotropin levels are difficult to interpret because of the intrauterine pregnancy that can increase them accordingly (Stanic 2020).

Management of the ectopic pregnancy can be conservative or non-conservative. The in-situ injection of potassium chloride or methotrexate can be used, but the literature describes cases of infection, internal haemorrhage or fetal toxicity for the intra-uterine pregnancy when following this approach (Li 2016).

Potassium chloride or a hyperosmolar solution can be considered in the event of unusual extrauterine locations that would be challenging for a surgical approach (e.g. at the cervix), or in the presence of caesarean scars (Talbot 2011).

There are no teratogenic effects known to occur secondary to the anesthesia medications (Ninke 2015).

Multiple paraffin segments were examined after Hematoxylin-Eosin coloration. The pathology department used a Leica DMC 2900 microscope.

The pathology report confirmed the existence of an ectopic pregnancy through the presence of chorionic structures and decidualisation (Fig 4,5).

After the surgery, the patient had a normal pregnancy follow-up. She delivered by caesarean section at 39 weeks and 4 days a healthy boy with a birth weight of 3450g and an APGAR score of 10.

**Discussions**

Heterotopic pregnancy is difficult to diagnose before 9 weeks of gestation (Wang 2014). The gold standard in diagnosis is represented by transvaginal ultrasonography which identifies an intrauterine pregnancy as well as the characteristic lateral-uterine mass with a gestational sac (Petrides 2015).

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Although many studies claim that the laparoscopic approach is the gold standard of surgical management, mainly because of minimal manipulation of the uterus, better operative field exposure and less postoperative pain (Guan 2017), there is no consensus regarding this aspect (Chadee 2016).

**Conclusion**

Heterotopic pregnancy is a condition that should be suspected in all fertile patients with amenorrhea and abdominal pain. Early management has a vital impact on the patient’s life and the outcome of the intrauterine pregnancy.

Patient consent was obtained for publication of this manuscript.

**References**


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