The incidence of ectopic pregnancy reaches nearly 1.5–2% of all pregnancies. Although the incidence has risen about six-fold over the past 30 years, the risk of death related to ectopic pregnancy has been reduced by almost 90%.\(^1\) The highest mortality rate occurs at 15–19 years of age.\(^2\) However, the likelihood of death due to an ectopic pregnancy is still higher than that from a live birth. An ectopic pregnancy mostly has a classical triad of symptoms: delayed menses, vaginal bleeding, and lower abdominal pain. The other symptoms associated with a ruptured ectopic pregnancy are shoulder pain, lightheadedness, and shock. Interference with normal tubal transport mechanisms may predispose to ectopic pregnancy. It is possible, but unproven, that endocrine factors predisposing to premature implantation take part in the pathogenesis of ectopic pregnancy.\(^3\)
most common site of ectopic implantation is a fallopian tube, accounting for 98% of all ectopic pregnancies. Overall, 70% of ectopic pregnancies are implanted in the ampulla, 12% are implanted in the isthmus, 11% are implanted in the fimbria, and 2% are implanted in interstitial (cornual) segments.**4**5** Ovarian, abdominal, and cervical locations are relatively rare.

Herein, we present an unusual case of a live 16-week isthmic ectopic pregnancy endangering the life of the mother.

CASE REPORT

A 23-year-old woman, gravida 2, para 1, was admitted to the university polyclinic for a routine pregnancy examination. She had a gestation period of 16 weeks and 5 days according to the last menstrual period and was suffering from right-sided lumbar pain. Her medical history did not report any risk factors for ectopic gestation. Abdominal examination revealed a tenderness to palpation in the right upper and lower quadrants of the abdomen without guarding and rebound tenderness. Her blood pressure and heart rate were in the normal ranges, 100/75 mmHg and 92 bpm, respectively. Her complete blood count showed a normal hemoglobin level of 12.1 g/dL and normal levels of leukocytes and platelets. The ultrasonographic examination confirmed a live intrauterine pregnancy at 16 weeks and 3 days of gestation. On the ultrasonogram, a slight dilatation and a crystalloid appearance of the right maternal renal pelvis were detected. It was decided to refer the case for consultation to an urologist. Thereafter, her skin became pale, and the patient suffered from a shortness of breath in a short time. She was urgently transferred to the emergency department. The blood pressure reading was in the normal range, but the heart rate increased to 125 bpm. A second ultrasonography was performed, and an accumulation of fluid in the maternal abdominal cavity and a live intrauterine fetus were reported. A repeat test showed that hemoglobin dropped to 7.1 g/dL. The patient gave the written informed consent before surgical intervention. An urgent laparotomy was decided upon to determine the cause of the clinical condition.

Under general anesthesia, a median laparotomy was performed, which revealed 2.5 L of intraabdominal blood. In addition, there was an unruptured right-sided ectopic pregnancy, which was located in the isthmic portion of the tube and bled from the fimbrial end (Figures 1, 2). A total right salpingectomy was performed. When the excised fallopian tube was opened, a male fetus weighing 96 grams, with normal external appearance, was delivered (Figure 3). Four units of erythrocyte suspension were administered to the patient. The postoperative period was uneventful, and the patient was discharged on the 10th day of hospitalization.

The histological results revealed that the development of the fetus was compatible with 16–17 weeks of gestation. The results of examination of fetal organs were reported to be normal. A microscopic study revealed chorionic villi and trophoblastic cells in the tubal lumen and attenuation of the muscular layer of the tube uterine (Figure 4).

DISCUSSION

The present case is a rare example of isthmic ectopic pregnancy, which was viable at 17 weeks of gestation after a spontaneous conception. We were urged to perform a laparotomy to determine the reason why the life of the mother was at risk.
Few risk factors have been described that predispose women to ectopic pregnancy, but 50% of the women do not show them. Some of the reported risk factors are previous ectopic pregnancy, tubal surgery, intrauterine device, tubal pathology, infertility, the use of assisted reproductive technology, previous genital infections, smoking, and prior cesarean delivery. However, our case had not used any contraception method, and had no history of the other risk factors.

The histopathology of ectopic pregnancies shows a variety of implantation sites. Ectopic implantation in the tubal isthmus typically results in penetration of the tubal wall at a relatively early stage. In contrast, ampullary and especially interstitial ectopic implantations mostly lead to delayed penetration of the involved tubal segment, because the more muscular segments are less distensible. As a rule, a tubal rupture in the first weeks of pregnancy gives an impression that the ectopic pregnancy is situated in the isthmic portion of the tube. When a fertilized ovum is implanted within the interstitial portion, rupture occurs after 14–16 weeks and is usually spontaneous or occurs after coitus or a bimanual examination. There are few cases of advanced isthmic pregnancies reported in the literature. Noteworthy, the ectopic gestation in the present case progressed to 17 weeks without any clinical warning signs.

Ultrasonography is the usual diagnostic procedure of choice in most cases. The confirmation of an intrauterine pregnancy is performed by observing the uterine wall surrounding the fetus and placenta. Occasionally, sonographic findings may be vague, depending on examiner’s experience and the quality of ultrasound equipment. Diagnosis of advanced ectopic gestations is only possible in 50-90% of unsuspected cases. However, even trans vaginal sonography cannot reveal ectopic implantation in up to 10% of the cases. In advanced ectopic cases, surgical intervention is essential for diagnosis and treatment to prevent likely dangerous hemorrhage. In our case, an appropriate diagnosis of the implantation site was missing even at the late stage of gestation.

The incidence of uterine congenital anomalies due to Müllerian defects in normal fertile women
is 3.2%. Unicornuate uterus is thought to be seen in 2.4-13% of all Müllerian anomalies.\textsuperscript{11} The pregnancy in a rudimentary horn (RH) is estimated between 1/76,000 and 1/140,000.\textsuperscript{12} In most cases, the gestational duration is longer than a tubal pregnancy. As the uterine wall in RH is thicker and has the variable musculature, the risk of uterine horn rupture is near to 50% especially during second and third trimester. Tubal pregnancy, cornual pregnancy, intrauterine pregnancy and abdominal pregnancy are common sonographic misdiagnoses.\textsuperscript{13} Tsafrir et al. proposed the following criteria for ultrasonographic diagnosis: (1) a pseudo pattern of an asymmetrical bicornuate uterus, (2) absent visual continuity tissue surrounding the gestational sac and the uterine cervix, and (3) the presence of myometrial tissue surrounding the gestational sac.\textsuperscript{14} In a review, sensitivity of sonography for diagnosis of RH pregnancy was estimated at 26%. RH pregnancy should always be considered as a differential diagnosis of tubal pregnancy. A tubal pregnancy will not have a ring of myometrium surrounding the gestational sac. Additionally, hyper-vascularity typical to placenta accreta may support the diagnosis of RH pregnancy.\textsuperscript{15}

In conclusion, if an isthmic pregnancy is not diagnosed at an earlier gestational age, it poses a risk of potentially catastrophic intra-abdominal bleeding. Therefore, an urgent laparotomy becomes a necessity for hemodynamically unstable patients.\textsuperscript{9,10} Clinicians must be aware of the implantation site at an early stage of gestation to avoid a misdiagnosis as an intrauterine pregnancy at later stages.

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\section*{REFERENCES}