Fetal Entrapment in the Myometrium Caused by Ectopic Implantation Post Myomectomy

Fetal entrapment in myometrial dehiscence after myomectomy has been rarely reported. A 38-year-old woman, gravida 4, parity 2, with suspected amniotic or adhesion band was referred to our hospital at 234/7 weeks. She had undergone open myomectomy and cesarean delivery twice. Ultrasonography revealed two separate compartments divided by a septum-like structure with a small orifice. Magnetic resonance imaging (MRI) finding suggested an ectopic implantation on the endometric and myometrial dehiscence. At 25 weeks, the thoracic cage and umbilical cord escaped through the orifice. Emergent cesarean delivery and subtotal hysterectomy were performed for the placental increta at the myometrial dehiscent site. In fetal entrapment in myometrial dehiscence, MRI is valuable for the assessment of the fetus and spatial arrangement in the uterine cavity.

Key Words: Dehiscence, Entrapment, Fetal, Myomectomy, Uterus

Introduction

The risk of uterine rupture or dehiscence in pregnant women with prior myomectomy is relatively well recognized. According to the literature and meta-analysis, the risk of uterine rupture after laparoscopic myomectomy is 0.75%, similar to that of laparotomic myomectomy. A recent single-center study in Korea also showed that in 523 pregnancies, uterine rupture after laparoscopic myomectomy occurred in 0.6% (3 cases); miscarriage, in 13%; and preterm delivery, in 10.4%. Uterine rupture after a previous uterine surgery can manifest as a frank rupture or dehiscence. Unlike frank rupture, uterine dehiscence is incidentally observed at the time of cesarean delivery. However, fetal entrapment in myometrial dehiscence after myomectomy is rarely reported. Here, we present the diagnosis and management of a stuck fetus in the myometrium, which was initially suspected as an intrauterine adhesion band, in a woman with a history of myomectomy.

Case

A 38-year-old woman, gravida 4, parity 2, was referred to our clinic at 234/7 weeks for the evaluation of an abnormal pregnancy with suspected amniotic or adhesion band. She had undergone an open myomectomy due to a fist-sized myoma. Her obstetric history indicated that she had undergone dilatation and curettage twice, once due to artificial abortion at 5 weeks and the other for spontaneous abortion at 7 weeks, and cesarean delivery twice, which were both uneventful. Ultrasonographic examination at 20 weeks at a local clinic revealed a complete septum-like structure suspected as uterine synechia, while that performed in our center also revealed a horizontal synechia or band-like structure in the upper third of the uterus (Fig. 1). Two separate compartments were found: the upper pocket where the fetus was stuck showed nearly absent amniotic fluid, and the lower pocket was mainly...
filled with amniotic fluid. The deepest pocket in the upper and lower compartments was 1.1 and 14.6 cm deep, respectively. The diameter and band thickness of the orifice between the two compartments were 3.3 and 0.7 cm, respectively, and the left shoulder of the fetus protruded through the band orifice but could move up and down while showing moderate edema. The patient was hospitalized for lower abdominal discomfort and continuous fetal surveillance.

Fetal magnetic resonance imaging (MRI) performed the following day after admission indicated that the pregnancy was based on an ectopic implantation probably through the endometrial and myometrial dehiscence, and the main amniotic sac was herniated into the original uterine cavity as the fetus grew (Fig. 2). We assumed an extrauterine adhesion due to abnormal ectopic implantation. The amniotic sac was herniated through the myometrial dehiscence into the original uterine cavity.

Finally, at 25 weeks, not only the fetal arm but also part of the thoracic cage and umbilical cord escaped through the orifice. The fetal heartbeat showed recurrent variable deceleration followed by prolonged deceleration of approximately 60 beats per minute (bpm) for 15 minutes. Therefore, an emergent cesarean delivery was performed via a vertical incision in the upper compartment of the uterus. A male neonate weighing 790 g was delivered, with an Apgar score of 1 at 1 minute and 5 at 5 minutes. We could not take a cord blood sample because of umbilical cord collapse. As the placental increta was complicated and the patient did not want further pregnancy, a subtotal hysterectomy was performed. She remained hemodynamically stable with a
postoperative hemoglobin level of 10.4 g/dL and was discharged on postoperative day 4. Unfortunately, the baby died 15 days after birth because of severe pulmonary hypoplasia and proven late-onset neonatal sepsis caused by *Staphylococcus aureus*.

According to the pathological examination of the uterus, the right upper mid-portion of the uterine cavity demonstrated a myometrial dehiscence rather than an intrauterine adhesion band, forming the second uterine cavity connected to the original one by a narrow opening (Fig. 3). The placental disc adhered and invaded the myometrium surrounding the dehiscent site. The placental tissue showed increased syncytiotrophoblast and fetal vessels with thrombi, suggesting maternal underperfusion and fetal thrombotic vasculopathy; however, neither acute chorioamnionitis nor funisitis was observed.

**Discussion**

We report an unusual case of fetal entrapment secondary to an ectopic implantation in a patient with a previous myomectomy. Most ectopic implantation (95%) occur in the ampulla, infundibular, and isthmic segments of the fallopian tube. Fewer than 5% of ectopic implantation occur in the interstitial segment of the fallopian tube, cervix, and anterior lower uterine segment of the cesarean delivery scar, ovary, or peritoneal cavity. Herein, we report a case of implantation in the myometrial dehiscence site that was confirmed by gross pathological examination.

Through a search on PubMed (English language; through April 2020; search terms: “uterus,” “dehiscence,” “synechia,” “fetal,” and “entrapment,” both individually and with several combinations), we identified four references on intrauterine fetal entrapment. In 2003, Deering et al. reported a case with intrauterine head entrapment of the second twin by uterine synechia that progressed to fetal distress with a cesarean delivery. In 2016, Bouet et al. published a case with fetal legs in a large herniated amniotic sac through a 2.5-cm uterine dehiscence in a patient who had five previous cesarean deliveries. In 2017, fetal leg entrapment was reported in a patient with uterine diverticulum secondary to a previous uterine instrumentation complicated by compartment syndrome and rupture of the diverticular portion. In the three previous cases, the patients were previously indicated for uterine surgery such as uterine curettage or cesarean delivery. One case without surgical intervention was fetal entrapment secondary to the compartmentalization due to a postoperative hemoglobin level of 10.4 g/dL and was discharged on postoperative day 4. Unfortunately, the baby died 15 days after birth because of severe pulmonary hypoplasia and proven late-onset neonatal sepsis caused by *Staphylococcus aureus*.

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submucosal myoma. To our best knowledge, this is a rare case of fetal entrapment with ectopic implantation in myometrial dehiscence.

A herniated amniotic sac, which is a fetal membrane bulging into the abdominal cavity, is an unusual manifestation of uterine rupture or dehiscence during pregnancy. According to a study by lemura et al., who summarized 24 cases of herniated amniotic sac, 21 patients had a history of cesarean delivery, uterine curettage, and myomectomy, whereas 2 patients had no prior intrauterine intervention and the remaining patient had only a history of amniocentesis. Mishina et al. also reported a case without a history of intrauterine intervention except amniocentesis after that amniotic sac was prolapsed through the small uterine wall defect. In these cases, the fetus was in the intrauterine cavity and herniated amniotic sac was located in the abdominal cavity. Our case is the reverse of the above-mentioned cases in terms of herniation, that is, ectopic implantation in the myometrial dehiscence caused fetal entrapment and herniation of the amniotic sac into the original intrauterine cavity. As the amniotic fluid volume in the compartment where the fetus was scant, severe pulmonary hypoplasia of the neonate seemed inevitable.

The fetal entrapment associated with oligohydramnios can lead to similar fetal and neonatal consequences of early onset preterm premature rupture of membranes. Usually, severe oligohydramnios in the second trimester cause fetal lung hypoplasia, frequently resulting in neonatal death. In general, the prevalence of lung hypoplasia in cases of early onset oligohydramnios (occurring at <24 weeks) is known to be 11.1–25%. Meanwhile, the optimal timing of delivery with mid-trimester oligohydramnios has not been clearly determined. In ongoing pregnancies of fetal entrapment complicated with oligohydramnios, the risks and potential sequelae of cord compression, preterm birth, and neonatal abnormalities need to be discussed. Serial ultrasonography for the assessment of amniotic fluid volume and continuous fetal heart rate monitoring for abrupt events related to entrapment may be necessary. Also, according to the American College of Obstetrics and Gynecologists guideline for periviable births, antenatal corticosteroids can be considered for infants born between 230/7 and 256/7 weeks of gestation. In our case, we decided on continuous fetal heart rate monitoring at 234/7 weeks of gestation and acute cord compression happened through the orifice of the cavity of entrapment during the hospital stay.

In conclusion, we report a rare case of fetal entrapment in myometrial dehiscence in a patient who had a prior myomectomy. In a previously reported case of uterine synechia or amniotic band syndrome, fetoscopic surgery released the fetal entrapment. However, we did not consider fetoscopic intervention because of the myometrial dehiscence. Once this condition is diagnosed, close surveillance of the fetal heart rate is vital. The use of MRI is valuable for understanding the spatial arrangement in the uterine cavity and precise assessment of the fetus and placental implantation.

Financial Disclosure

The authors did not report any potential conflicts of interest.

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