Case Report

A Case of Ruptured Decidualized Ovarian Endometrioma: Usefulness of Serial MRI for Determining Adequate Management

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Received 1 March 2022; Revised 10 July 2022; Accepted 15 July 2022; Published 31 July 2022

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Decidualization can originate in ovarian endometrioma by elevated serum progesterone levels during pregnancy, which mimics malignancy on ultrasonography. Moreover, decidualized ovarian endometrioma may rupture and cause acute abdominal pain during pregnancy. Magnetic resonance imaging (MRI) is reportedly useful in differentiating decidualized ovarian endometriomas from malignancies. However, to our knowledge, serial MRI of decidualized ovarian endometrioma before and after rupture has not been reported. Herein, we report the case of a 39-year-old woman with a ruptured decidualized ovarian endometrioma in which serial MRI findings were useful for adequate management.

1. Introduction

Ovarian endometrioma occurs in 17–44% of patients with endometriosis and accounts for 35% of all benign ovarian cysts [1]. During pregnancy, increased progesterone levels may cause the decidualization of ovarian endometriomas, which mimics malignancies on imaging [2]. Recently, magnetic resonance imaging (MRI) has been reported to be useful in differentiating decidualized tissue from malignancies [3]. Moreover, decidualization can increase the risk of rupture of ovarian endometrioma [4]. Therefore, accurate diagnosis of a ruptured decidualized ovarian endometrioma during pregnancy is important for its adequate management.

Herein, we report a case of ruptured decidualized ovarian endometrioma in which serial MRI findings before and after the rupture were useful for determining the appropriate management.

2. Case Presentation

A 39-year-old woman (gravida 0, para 0) presented to our hospital for infertility treatment. She had a history of a 30 mm diameter right ovarian endometrioma. She became pregnant by oral clomiphene-intruterine artificial insemination. At 20 weeks of gestation, transvaginal ultrasonography revealed that the ovarian endometrioma had enlarged...
The patient’s serum cancer antigen-125 levels were slightly elevated (37 U/mL; normal, <35 U/mL), and carbohydrate antigen 19-9 levels were within normal limits (16 U/mL; normal, <37 U/mL). MRI for further evaluation showed a well-circumscribed teardrop-shaped cystic lesion with mural nodules measuring 68 × 45 × 35 mm in the right ovary (Figure 1). The cystic lesion showed uniformly marked high signal intensity on the T1-weighted image (a, asterisk) and intermediate signal intensity (b, asterisk) on T2-weighted images consistent with ovarian endometrioma. The mural nodules show intermediate to high signal intensity on T2-weighted images (b, c; short arrows) similar to the placenta (c, long arrows). Diffusion-weighted image (d, short arrows) and ADC map (e, short arrows) show the mural nodules with a high ADC value (1.6 × 10^{-3} mm²/s). Based on these imaging findings, we diagnosed a decidualized ovarian endometrioma.

At 27 weeks of gestation, the patient complained of sudden abdominal pain. Emergent MRI was performed. The decidualized ovarian endometrioma had disappeared; however, bloody ascites were found (Figure 2). Based on serial MRI findings, rupture of the decidualized ovarian endometrioma was suspected. The abdominal pain subsided immediately after injection of pentazocine 15 mg, and conservative observational management was chosen.

At 37 weeks of gestation, the patient underwent an elective cesarean section and right ovarian cystectomy simultaneously. Ruptured ovarian cystic lesions were confirmed during the surgery. The cystic lesion contained chocolate-like bloody fluid. Histopathological findings revealed decidual reaction of the ovarian endometrioma and no evidence of malignancy (Figure 3). The final diagnosis was a ruptured decidualized ovarian endometrioma. The postoperative course was uneventful, and the patient was discharged 7 days later.

3. Discussion
We have reported the case of a ruptured decidualized endometrioma during pregnancy, which was useful to diagnose accurately by serial MRI and determine the patient’s treatment options. In the present case, we diagnosed a decidualized ovarian endometrioma upon the first MRI examination during pregnancy. Further, we suspected rupture of the endometrioma due to its disappearance and bloody ascites found on serial MRI at the onset of sudden abdominal pain. We further employed a conservative observational management approach, and elective cesarean section and right ovarian cystectomy were simultaneously performed at full term. Intraoperative and histologic examination demonstrated a ruptured decidualized endometrioma.

Endometriosis is a common disease occurring in 5–10% of women of reproductive age [2]. It is one of the main causes of infertility due to mechanical factors, including adhesions, tubal blockage, and ovulation disorders [5]. The number of pregnancies with endometriosis has been increasing due to the progression of assisted reproductive technologies [2]. Decidualization can also originate in ovarian...
endometrioma by elevating serum progesterone levels during pregnancy [2]. Previous reports revealed that approximately 12% of ovarian endometriomas have decidual reactions during pregnancy [2]. Decidualization primarily occurs in the second trimester and regresses during the third trimester or postpartum period [2].

Ovarian endometriomas can rupture for various reasons, including adhesions, tissue fragility, increased external pressure, or increased internal pressure due to mass enlargement. During pregnancy, increased levels of progesterone tend to shrink ovarian endometrioma [2]. However, decidualization is expected to increase ovarian endometrioma rupture due to enlargement and wall softening caused by a severe inflammatory response [2, 6]. To date, there have been six case reports of ruptured decidualized ovarian endometrioma.

Figure 2: Magnetic resonance imaging (MRI) of the pelvis performed at 27 weeks of gestation. (a) Axial T1-weighted image. (b) Axial T2-weighted image. (c) Sagittal T2-weighted image. (d) Axial diffusion-weighted image (b factor: 1000 s/mm²). (e) ADC map. (f) Coronal T1-weighted image with fat saturation. The cystic lesion disappears (a–e). Ascites around the uterus shows high signal intensity on fat-suppressed T1-weighted images, which is consistent with bloody ascites (f, arrows). Based on serial MRI findings, rupture of the decidualized ovarian endometrioma is suspected.

Figure 3: Microscopic view of a mural nodule (hematoxylin and eosin staining, magnification: 100x). Edematous tissue with abundant stromal cells’ cytoplasm is visible (asterisk). These findings are consistent with decidualization.
<table>
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<tr>
<th>Authors (year)</th>
<th>Patient age (years)</th>
<th>Size (mm)</th>
<th>Time of rupture</th>
<th>Imaging</th>
<th>Preoperative diagnosis</th>
<th>Treatment during pregnancy</th>
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<tbody>
<tr>
<td>Garcia-Velasco et al. (1998) [7]</td>
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<td>83 × 54</td>
<td>First trimester (9 weeks pregnant)</td>
<td>US</td>
<td>Hemorrhagic corpus luteum or endometrioma</td>
<td>Laparotomy (left salpingo-oophorectomy)</td>
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<tr>
<td>Gregora et al. (1998) [8]</td>
<td>44</td>
<td>60</td>
<td>Second trimester (17 weeks pregnant)</td>
<td>US</td>
<td>Endometriomas*</td>
<td>Laparotomy (opening the cyst and stripping the lining)</td>
</tr>
<tr>
<td>Reif et al. (2011) [10]</td>
<td>25</td>
<td>NA (after rupture)</td>
<td>Second trimester (27 weeks pregnant)</td>
<td>US</td>
<td>NA</td>
<td>Laparotomy (operative hemostasis and preterm cesarean section)</td>
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<tr>
<td>Present study</td>
<td>39</td>
<td>68 × 45 × 35</td>
<td>Second trimester (27 weeks pregnant)</td>
<td>US MRI</td>
<td>Rupture of decidualized endometriotic cyst</td>
<td>Conservative treatment, an elective cesarean section, and right ovarian cystectomy at 37-week pregnancy</td>
</tr>
</tbody>
</table>

NA: not applicable; US: ultrasonography; MRI: magnetic resonance imaging. *The preoperative diagnosis of the ruptured tumor was not described.
endometrioma (Table 1) [2, 7–11]. Among the six cases, rupture occurred during the first trimester in two cases, second trimester in three cases, and third trimester in one case.

Ultrasonography is the first choice for the screening of decidualized ovarian endometrioma. However, decidualized tissues appear as mural nodules in the cyst mimicking malignant tumors, such as endometrioid carcinoma and clear cell carcinoma arising from ovarian endometrioma [12]. Moreover, no previously reported cases (Table 1) were accurately diagnosed preoperatively with ruptured decidualized ovarian endometrioma. It is considered that ultrasonography may be limited of the visual field during pregnancy.

MRI is reported to be useful in the diagnosis of decidualized tissue, which has a thin broad structure and shows intermediate to high signal intensities on T2-weighted images, similar to the placenta. Morisawa et al. reported that the ADC values of the decidualized regions were significantly higher than those of the mural nodules of endometrioid carcinoma or clear cell carcinoma (1.77 × 10⁻³ mm²/s vs. 1.13 × 10⁻³ mm²/s) on diffusion-weighted images [3]. In the present case, the patient’s MRI findings were similar, and we diagnosed a decidualized ovarian endometrioma. Additionally, MRI of a ruptured endometrioma can identify shrinkage or disappearance of the cyst and bloody ascites, as in the present case [13].

To our knowledge, serial MRI before and after the rupture of decidualized ovarian endometrioma during pregnancy has not been reported. In the present case, serial MRI findings before and after rupture showed disappearance of the cyst with MRI findings characteristic of decidualized ovarian endometrioma, which contributed to the accurate diagnosis.

There is no definite management for ruptured benign ovarian cystic lesions during pregnancy. All of the six patients mentioned in Table 1 underwent surgery. However, the risk of miscarriage associated with abdominal surgery during pregnancy increases in the first trimester. In addition, the risk of adverse outcomes such as miscarriage, preterm delivery, and intraperitoneal death is increased in surgeries performed after 23 weeks of gestation [14]. Therefore, we believe that conservative treatment can be a management option in cases without torsion or malignancy and with stable hemodynamics or pain.

In conclusion, here we accurately diagnosed a ruptured decidualized ovarian endometrioma by serial MRI and chose conservative observation. In the future, the indications for conservative treatment of benign ovarian cystic lesions during pregnancy should be considered with the addition of MRI findings.

Conflicts of Interest

The authors declare no conflicts of interest regarding the publication of this article.

References


