Clinical diagnostic value and analysis of MRI combined with ultrasound in prenatal pernicious placenta previa with placenta accreta

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Background: Placenta previa is one dangerous disease which threatens the health of pregnant women and their fetuses. The purpose of this study was to evaluate the clinical value of ultrasound combined with magnetic resonance imaging (MRI) in screening for placenta previa complicated by placenta accreta.

Methods: Seventy patients with abnormal fetal position admitted to our hospital from January 2019 to January 2020 were selected for the study. Patients were diagnosed by ultrasound alone, MRI alone, and ultrasound combined with MRI. Diagnostic accuracy, sensitivity, specificity and false positive and negative diagnosis rates were evaluated against the postoperative pathological examinations of the patients.

Results: The diagnostic accuracy, sensitivity and false negative rate for ultrasound combined with MRI were 86.27%, 97.78% and 72.00%, respectively. These results were significantly superior to those of MRI or ultrasound alone (P<0.05). The specificity and false positive rate for ultrasound combined with MRI were 13.73% and 5.26%, respectively, which were not significantly different from those for MRI or ultrasound alone (P>0.05).

Conclusions: Compared with ultrasound or MRI alone, ultrasound combined with MRI has higher accuracy and sensitivity in the diagnosis of placenta previa with placenta accreta, along with lower false positive diagnosis rates. These findings are clinically important for improving the diagnostic efficiency.

Keywords: Ultrasound; magnetic resonance imaging (MRI); ultrasound combined with MRI; placenta previa; placenta accreta

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Introduction

At present, the pathogenesis of placenta previa remains unclear, and the study has indicated that the incidence of placenta previa is more than 10% (1). Some studies suggest that this condition is closely related to the age, number of previous cesarean deliveries, and smoking history of the pregnant women, while other studies have shown that patients with uterine-related diseases are more prone to placenta previa during pregnancy. During normal gestation, the villi on the placenta can interact with the villi on the endometrium to promote placenta implantation. Subsequently, the thickness of the endometrium increases, thus forming the decidua where placental villus are usually unable to invade the myometrium of the uterus (2); however, in some cases, inflammation, surgery, and dysplasia may cause endometrial damage in pregnant women and
attachment of the embryo to these damaged sites will cause placental villi to cross the endometrium and invade the myometrium, resulting in placenta accreta (3). Clinical data has revealed that placenta accreta could lead to coagulation disorders in patients, resulting in massive and potentially fatal hemorrhage during or after delivery (4-6). Therefore, timely diagnosis of placenta previa combined with placenta accreta is of great importance during pregnancy. At present, the diagnosis of placenta previa combined with placenta accreta occurs mainly through magnetic resonance imaging (MRI) and ultrasound (7,8); however, patient physique, placenta position, external environment and other factors can seriously impact diagnostic results. These issues can interfere with timely and effective diagnosis of placenta previa with placenta accreta and increase the risk of adverse events during pregnancy (9,10). Therefore, it is imperative to investigate an efficient and rapid diagnostic method for placenta accreta. Some studies have confirmed that ultrasound combined with MRI can improve efficiency compared to a single diagnostic method (11). This study intends to compare the diagnostic value of ultrasound combined with MRI compared with ultrasound alone and MRI alone in the diagnosis of placenta previa with placenta accreta by evaluating the accuracy, sensitivity, specificity, false positive and negative diagnosis rates. Ultimately, these investigations aim to provide a standard reference for the clinical diagnosis of placenta previa combined with placenta accreta. We present the following article in accordance with the STARD reporting checklist (available at https://dx.doi.org/10.21037/apm-21-1285).

Methods

General information

Seventy patients admitted to our hospital from January 2019 to January 2020 with placenta previa or low-lying placenta (of varying degrees) were selected for the study. Participant age ranged from 25 to 43 years old, with an average age of 32.02±3.27 years old.

Inclusion criteria for participants: singleton pregnancy; second or third trimester of pregnancy; no metal implants in their body; physical condition met the criteria for MRI and ultrasound; conscious with normal communication ability.

Exclusion criteria for participants: state of tension, anxiety or tachypnea; obvious fetal movements, which affected the diagnosis effects; other diseases that seriously affected the results; refusal to cooperate with the study.

The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013) (12). This study was approved by the Ethics Committee of Linyi Central Hospital. Patients and their family were informed of the related contents of this study, and all subjects written informed consent.

Detection methods

Ultrasound examination: Patients were diagnosed using the GE-E8 diasonograph (GE Company, USA) and were informed to drink sufficient water to prior to examination to ensure a full bladder during imaging. Transabdominal ultrasonography and transvaginal probe were adopted with a frequency of 3.0-5.0 and 8–10 Hz, respectively. The color Doppler ultrasound was used to observe the lower segment of the uterus, including the thickness and location of the placenta, as well as the distribution and characteristics of blood flow between the posterior placental space and the patient's bladder.

MRI examination: SIEMENS Aerate 1.5T MRI scanner was used. The placenta was taken as the center for plain scanning. With the long axis of the patient’s uterus as the reference, plain scanning was conducted in the transverse, sagittal and coronal planes. The scanning slice thickness and gap were 5 and 1 mm, respectively, while the number of scanning slices was between 25 and 30. The transverse scanning, which was perpendicular to the uterine wall, started from the uterine fundus to the pelvic floor to ensure full view of the cervix and placenta, while the sagittal and coronal scans (parallel to the long axis of the uterus) ensured full view of the uterus and cervix. Scanning sequences were as follows:

(I) In T2-weighted turbo spin-echo imaging (TSE/T2WI) sequence, transverse scans were performed with repetition time (TR) of 3,010 ms, echo time (TE) of 71 ms, flip angle of 90° and 2 times of radio frequency (RF) excitation;

(II) In Haste/T2WI sequence, sagittal, transverse and coronal scans were performed with scanning parameters TR of 1,350 ms, TE of 91 ms, flip angle of 90° and 2 times of RF excitation;

(III) In True-Fisp sequence, sagittal, coronal and transverse scans were performed with scanning parameters TR of 3.8 ms and TE of 1.9 ms;

(IV) In volumetric interpolated breath-hold T1-weighted (VIBE-T1WI) sequence, transverse scans
Table 1 Calculation formula of diagnostic efficiency

<table>
<thead>
<tr>
<th>Accuracy rates</th>
<th>True positive rates</th>
<th>True negative rates</th>
<th>False positive diagnosis rate</th>
<th>False negative diagnosis rate</th>
</tr>
</thead>
<tbody>
<tr>
<td>= (TP + TN)/(TP + TN + FP + FN) × 100%</td>
<td>= TP/(TP + FN) × 100%</td>
<td>= NP/(FP + TN) × 100%</td>
<td>= FP/(TP + FP) × 100%</td>
<td>= FN/(TN + FN) × 100%</td>
</tr>
</tbody>
</table>

TP, true positive; TN, true negative; FP, false positive; FN, false negative.

Table 2 Gold standard evaluation of different detection methods

<table>
<thead>
<tr>
<th>Ultrasound</th>
<th>Pathological examination</th>
<th>MRI</th>
<th>Pathological examination</th>
<th>Ultrasound + MRI</th>
<th>Pathological examination</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>P</td>
<td>N</td>
<td>S</td>
<td>P</td>
<td>N</td>
</tr>
<tr>
<td>P</td>
<td>35</td>
<td>8</td>
<td>43</td>
<td>P</td>
<td>33</td>
</tr>
<tr>
<td>N</td>
<td>10</td>
<td>17</td>
<td>27</td>
<td>N</td>
<td>12</td>
</tr>
<tr>
<td>S</td>
<td>45</td>
<td>25</td>
<td>70</td>
<td>S</td>
<td>45</td>
</tr>
</tbody>
</table>

P, positive; N, negative; S, summation; MRI, magnetic resonance imaging.

Calculation methods

The results of pathological examination were treated as the gold standard (Table 1).

Statistical analysis

SPSS 22.0 software was used to analyze data. Measurement data were expressed as mean value ± standard deviation and tested using a t-test. Enumeration data were tested using the chi-square test. Results with P<0.05 were considered to be statistically significant.

Results

Different diagnostic methods

Pathological examination confirmed that 45 of the 70 participants were diagnosed as implantable types (true positive) and 25 as non-implantable types (true negative). From ultrasound examination, 43 patients were classified as placenta previa complicated with placenta accreta, while 27 as placenta previa. From MRI examination, 42 patients were classified as placenta previa complicated with placenta accreta, while 28 as placenta previa. From combined MRI and ultrasound diagnosis, 51 patients were classified as placenta previa complicated with placenta accreta, while 19 as placenta previa. There were some deviations in the pathological examination results among the three diagnostic methods (Table 2).

Comparison of single ultrasound and combined diagnosis

Pathological examination results were used as the gold standard to compare the efficacy of ultrasound alone and combined ultrasound and MRI for diagnosis of placenta previa complicated with placenta accreta (Figure 1, Table 3). The results showed that the diagnostic accuracies for single ultrasound versus combined ultrasound and MRI were 81.40% and 86.27%, respectively (P<0.05). Sensitivity for each diagnostic method was 77.78% and 97.78%, respectively (P<0.05) and specificity was 68.00% and 72.00%, respectively (P>0.05). False positive diagnosis rates were 18.60% and 13.73%, for each method respectively (P>0.05) while the false negative diagnosis rates were 37.04% and 5.26%, respectively (P<0.05).
Comparison of single MRI and combined diagnosis

Pathological examination results were used as the gold standard to compare the efficacy of MRI alone and combined MRI and ultrasound for diagnosis of placenta previa complicated with placenta accreta (Figure 2, Table 4). The results showed that the diagnostic accuracies for MRI alone versus combined MRI and ultrasound were 77.78% and 86.27%, respectively (P<0.05). Sensitivity for each diagnostic method was 73.33% and 97.78%, respectively (P<0.05) and specificity was 64.00% and 72.00%, respectively (P>0.05). False positive diagnosis rates were 21.43% and 13.73%, respectively (P>0.05) while false negative diagnosis rates were 42.86% and 5.26%, respectively (P<0.05).

Discussion

Clinical data has shown that placenta previa complicated with placenta accreta may lead to coagulation disorders, resulting in massive and potentially fatal bleeding during or after delivery (13,14). Other studies have also shown that patients with uterine-related diseases are more likely to develop placenta accreta during pregnancy (15,16). During pregnancy, the villi on the placenta can interact with the villi on the endometrium and promote placenta accreta. Recently, both MRI and ultrasound are major methods for diagnosing placenta previa combined with placenta accreta; however, these methods have certain deficiencies pertaining to diagnostic sensitivity and specificity (17,18). This study adopted the diagnostic methods of MRI, ultrasound, and
ultrasound combined with MRI, to diagnose the presence of placenta previa complicated with placenta accreta in 70 patients with placenta previa. The clinical diagnostic value of MRI combined with ultrasound was evaluated using measures of accuracy, sensitivity, specificity, false positive and false negative diagnosis rates. From ultrasound examination, 43 patients were classified as placenta previa complicated with placenta accreta, while 27 as placenta previa. From MRI examination, 42 patients were classified as placenta previa complicated with placenta accreta, while 28 as placenta previa. From combined MRI and ultrasound diagnosis, 51 patients were classified as placenta previa complicated with placenta accreta, while 19 as placenta previa. There were some deviations in the pathological examination results among the three diagnostic methods.

Accurate diagnosis of patients with placenta accreta can reduce the risk of hemorrhage during childbirth. Some studies have pointed out that misdiagnosis and missed diagnosis can occur when ultrasound and MRI diagnosis are used in isolation (19,20). Einerson et al. (21) found that the misdiagnosis rate of placenta accreta using MRI was 23% and the missed diagnosis rate was approximately 14%; additionally, only two out of 14 patients with placenta accreta detected by ultrasound were confirmed using MRI. Some studies have also shown that missed diagnosis that occurs when MRI or ultrasound are used in isolation may be related to changes in bladder filling due to inflammation, the thickness of abdominal fat layer, and placenta position (22-24). Xia et al. (25) found that combined ultrasound and MRI had higher sensitivity for placenta accreta diagnosis in the second trimester of pregnancy than ultrasound or MRI alone. This study also revealed that the combined ultrasound and MRI had significantly higher specificity and accuracy for the placenta accreta diagnosis either modality alone, reducing the risk of missed diagnosis and its implications.

The results of our study showed that the diagnostic accuracies for single ultrasound versus combined ultrasound and MRI were 81.40% and 86.27%, respectively (P<0.05). Sensitivity for each diagnostic method was 77.78% and 97.78%, respectively (P<0.05) and specificity was 68.00% and 72.00%, respectively (P>0.05). False positive diagnosis rates were 16.80% and 13.73%, for each method respectively (P<0.05) while the false negative diagnosis rates were 37.04% and 5.26%, respectively (P<0.05). In addition, our results showed that the diagnostic accuracies for MRI alone versus combined MRI and ultrasound were 77.78% and 86.27%, respectively (P<0.05). Sensitivity for each diagnostic method was 73.33% and 97.78%, respectively (P<0.05) and specificity was 64.00% and 72.00%, respectively (P>0.05). False positive diagnosis rates were 21.43% and 13.73%, respectively (P>0.05) while false negative diagnosis rates were 42.86% and 5.26%, respectively (P<0.05).

The false negative diagnosis rate for combined ultrasound and MRI diagnosis was significantly lower than that of MRI or ultrasound diagnosis alone, which suggests combining these methods could reduce the probability of missed diagnosis and allow for improved patient management and reduce potentially avoidable adverse events.

In conclusion, compared with single ultrasound or MRI, ultrasound combined with MRI has higher accuracy and sensitivity in the diagnosis of placenta accreta in patients with placenta previa, and has a lower rate of false negative diagnosis. These results have important clinical implications for the diagnosis and timely management of this condition.

**Acknowledgments**

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**Footnote**

*Reporting Checklist:* The authors have completed the STARD reporting checklist. Available at [https://dx.doi.org/](https://dx.doi.org)
Guo et al. Value of MRI combined with ultrasound on placenta accreta

Data Sharing Statement: Available at https://dx.doi.org/10.21037/apm-21-1285

Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at https://dx.doi.org/10.21037/apm-21-1285). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). This study was approved by the Ethics Committee of Linyi Central Hospital. Patients and their family were informed of the related contents of this study, and all subjects written informed consent.

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