Evaluation of Red Degeneration of Uterine Leiomyoma with Susceptibility-weighted MR Imaging

Mayumi Takeuchi1*, Kenji Matsuzaki2, Yoshimi Bando3, and Masafumi Harada1

Purpose: Red degeneration of uterine leiomyoma (RDL) is a hemorrhagic infarction caused by peripheral venous thrombosis. The peripheral high-intensity rim on T1-weighted MRI is characteristic for RDL; however, it may not be observed at all the phases of RDL. Susceptibility-weighted MR sequences (SWS) have exquisite sensitivity to blood products, and we hypothesized that the low-intensity rim due to the T2* shortening effects of blood products may be more clearly demonstrated on SWS. The purpose of this study is to evaluate the capability of SWS for the diagnosis of RDL.

Methods: Surgically proven 15 RDL, which showed suggestive MRI findings (high-intensity rim or entirely high signal intensity on T1-weighted imaging) were retrospectively evaluated. MRI was qualitatively evaluated for the presence of high-intensity rim around a mass on fat-saturated T1-weighted images, and low-intensity rim on T2-weighted images and on SWS (susceptibility-weighted imaging [SWI] or T2-star-weighted angiography [SWAN]).

Results: The high-intensity rim on T1-weighted images, low-intensity rim on T2-weighted images and on SWS were observed in 47%, 47%, and 100% of RDL, respectively. The other 53% of lesions showed entirely high signal intensity on T1-weighted images. Pathological examination revealed coagulative necrosis in all 15 lesions.

Conclusion: SWS may be helpful for the diagnosis of RDL by revealing characteristic peripheral low-intensity rim.

Keywords: leiomyoma, red degeneration, susceptibility-weighted magnetic resonance sequence, uterus, uterine sarcoma

Introduction

Red degeneration of uterine leiomyoma (RDL) is a subtype of degeneration, which often occurs during pregnancy, or with the use of oral contraceptives.1,2 On gross pathological examination, it is characterized by a red, hemorrhagic appearance of the leiomyoma if the surgery was performed during the acute phase of red degeneration. Red degeneration is a massive hemorrhagic infarction of leiomyoma caused by venous thrombosis within the periphery of a leiomyoma. Characteristic signal intensity patterns have been described on MRI as a peripheral high-intensity rim on T1-weighted images due to the T1 shortening effects of methemoglobin of blood products confined to the thrombosed numerous dilated vessels surrounding the tumor.1–4 On T2-weighted images, it may show variable signal intensity with a low-intensity rim due to the T2* shortening effects of deoxyhemoglobin, intracellular methemoglobin, or hemosiderin of blood products also confined to the thrombosed vessels.1–4 However, the high-intensity rim on T2-weighted images may not be observed at the very acute phase of red degeneration because of insufficient conversion of deoxyhemoglobin into methemoglobin, the low-intensity rim on T2-weighted images due to deoxyhemoglobin may be observed even at the very acute phase. However, because usual leiomyomas show low signal intensity on T2-weighted images, a peripheral rim-like low-intensity due to the residual un-degenerated area could be observed in degenerated leiomyoma without hemorrhagic infarction.

Susceptibility-weighted MR sequences (SWS) such as susceptibility-weighted imaging (SWI) and T2 star-weighted angiography (SWAN) can maximize sensitivity to susceptibility effects, and have exquisite sensitivity to blood products.5–8 We hypothesized that the low-intensity rim due to the T2* shortening effects of blood products may be more...
clearly demonstrated on SWS and be helpful for the diagnosis of red degeneration. The purpose of this study was to evaluate the capability of SWS for the diagnosis of red degeneration of uterine leiomyomas.

**Materials and Methods**

**Patients**
The Institutional Review Board in our hospital approved this retrospective study, and waived the requirement for written informed consent of patients. We cross-referenced the database of the Department of Obstetrics and Gynecology to identify all patients with histologically proven leiomyomas with possible red degeneration who had undergone MRI examinations including SWS between May 2010 and April 2016. Lesions with suggestive MRI findings for red degeneration such as well demarcated uterine myometrial masses exhibiting high-intensity rim or entirely high signal intensity compared to the uterine myometrium on T1-weighted imaging were included in this study. The patients were with or without suggestive symptoms for red degeneration (lower abdominal pain and/or fever) and suspected risk factors for onset (pregnancy, use of oral contraceptives, or hormonal drug administration such as gonadotropin-releasing hormone antagonist). All lesions were re-reviewed by a pathologist with 19 years of experience in gynecological pathology. A total of 15 lesions in nine women with a mean age of 39 years (range, 27–52 years) were included in the current study (Table 1). The median time from MRI examination to operation was 91 days (range 12–188). The risk factors for onset were pregnancy for five patients and hormonal drug administration for one patient; however, the other three patients did not have any risk factors. The median lesion size, which was the longest diameter measured by MRI was 59 mm (range, 18–88 mm).

**Magnetic resonance imaging**
Thirteen leiomyomas with red degeneration in seven patients were evaluated at 1.5T (Signa Excite HD or HDx, General Electric, Milwaukee, WI, USA), whereas two leiomyomas with red degeneration in two patients were evaluated at 3T (Signa 3T HD or Discovery MR750, General Electric) by using body-array torso coils. SWI consisting of both magnitude and phase images from 2D fast spoiled gradient recalled acquisition in the steady-state sequence (TR/TE, 650–700/30 ms; flip angle, 15–20 degrees; matrix size, 288 × 192; FOV, 28 × 28 cm; two signals acquired; section thickness, 8 mm; section gap, 1 mm) were obtained in two leiomyomas with red degeneration. Sequence parameters in 3T and 1.5T were almost identical except for flip angles (15 degrees in 3T, and 20 degrees in 1.5T). SWAN using multiple magnitude images with different echo times for the image generation: 3D multi-echo gradient echo (TR/TE, 42.9/27.5 ms for 3T and 78.8/49 ms for 1.5T; matrix size, 320 × 192–256; FOV, 28 × 28 cm; section thickness, 3–5 mm; spacing, 1.5–2.5 mm) were obtained in 13 leiomyomas with red degeneration. Fast spin-echo T1-weighted images (TR/TE, 4000–7000/99.3–100 ms) and spin-echo (TR/TE, 466.7–600/7.9–9.8 ms) or fast spoiled gradient-recalled echo (TR/TE, 2.9–4/1.3–1.7 ms) T1-weighted images with fat saturation were obtained in all patients.

**Table 1** Clinical courses, MRI findings, and pathological findings of patients

<table>
<thead>
<tr>
<th>Case</th>
<th>Age</th>
<th>Lesion</th>
<th>Risk factor</th>
<th>Symptom</th>
<th>Size</th>
<th>MRI findings</th>
<th>SWS-low</th>
<th>Pathological findings</th>
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<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<td>T1-high</td>
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<td>C.N.</td>
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<tr>
<td>1</td>
<td>27</td>
<td>1</td>
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<td>Pain</td>
<td>60</td>
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<td>rim</td>
<td>rim / s-m</td>
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<tr>
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<td>Pain</td>
<td>75</td>
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<td>rim</td>
<td>rim / s-m</td>
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<tr>
<td>3</td>
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<td>Pain</td>
<td>99</td>
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<td>rim / s-m</td>
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<tr>
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<td>rim</td>
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<td>25</td>
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<td>84</td>
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<td>rim / s-m</td>
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*Drug. Gonadotropin-releasing hormone antagonist; s-m, spotty to mottled signal voids; C.N., coagulative necrosis; H.D., hyalinized degeneration; Hem., hemosiderin deposits; V.T., venous thrombus. SWS, Susceptibility-weighted MR sequences.
**Analysis methods**

Two radiologists with 27 and 18 years of experiences in gynecological MRI qualitatively evaluated the images for the presence of a low-intensity rim around a mass on T2-weighted images and on SWS, and of a high-intensity rim around a mass on fat-saturated T1-weighted images. Both iso intensity mass compared to the myometrium with high-intensity rim and high-intensity mass compared to the myometrium with higher intensity rim were considered as “high-intensity rim” on fat-saturated T1-weighted images. The reviewers examined all MR images of the cases independently, and then resolved discrepancies by consensus.

**Results**

The high-intensity rim on T1-weighted images was observed in seven of 15 leiomyomas with red degeneration (47%) (Fig. 1), whereas the other eight leiomyomas with red degeneration (53%) showed entirely high signal intensity and the high-intensity rim was not clearly demonstrated (Fig. 2). The low-intensity rim on T2-weighted images was observed in seven of 15 leiomyomas with red degeneration (47%) (Fig. 1). The low-intensity rim on SWS was observed in all 15 leiomyomas with red degeneration (100%) (Figs. 1 and 2). Spotty or mottled signal voids within the tumors were observed in 13 of 15 leiomyomas with red degeneration (87%) (Fig. 1).

Pathological examination of the leiomyomas with red degeneration revealed coagulative necrosis in all 15 lesions, hyaline degeneration in 14 lesions, and obsolete hemorrhage (hemosiderin deposition) in 11 lesions. Probable peripheral venous thrombus was found in only one lesion.

**Discussion**

In the current study, the low-intensity rim on SWS was observed in all the leiomyomas with red degeneration, whereas the high-intensity rim on fat-saturated T1-weighted images was observed in 47% of the leiomyomas with red degeneration. According to the results, both SWS and fat-saturated T1-weighted images can provide specific information for the diagnosis of leiomyoma with red degeneration. SWS may be more sensitive for the presence of blood products than fat-saturated T1-weighted images.

Leiomyomas are the most common benign uterine neoplasm, which is composed of smooth muscle and varying amounts of fibrous tissue with/without various types of degeneration, such as hyaline, hydropic, myxoid, cystic, fatty, or red degeneration. Red degeneration (carneous degeneration) is a hemorrhagic infarction of uterine leiomyoma, which may cause acute abdominal pain. Because leiomyomas with red degeneration can be treated conservatively, preoperative diagnosis is important.1 Leiomyomas with red degeneration may show diffuse high signal intensity or a peripheral high-intensity rim on T1-weighted images depending on the degree of intra-tumoral hemorrhage, coagulative necrosis, or hyaline degeneration.1,4 The peripheral high-intensity rim on T1-weighted images reflecting methemoglobin of blood products confined to the thrombosed vessels is characteristic; however, this finding may not be observed at the very acute phase of red degeneration. Because oxyhemoglobin and deoxyhemoglobin do not cause the signal increase on T1-weighted images, high signal intensity due to the T1 shortening effect of methemoglobin may be observed at least 3 days after the onset.9,10 The peripheral low-intensity rim on T2-weighted images due to the T2 shortening effect of deoxyhemoglobin, intracellular methemoglobin or hemosiderin reflecting venous thrombosis is also characteristic for red degeneration, and may be observed even at the very acute phase due to deoxyhemoglobin. However, usual leiomyoma itself may show low signal intensity on T2-weighted images and evaluation of the low-intensity rim may occasionally be difficult. In our study, the prevalence of the high-intensity rim

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**Fig. 1** A 45-year-old woman with uterine leiomyoma with red degeneration. (A) On the fat-saturated T1-weighted image, a peripheral high-intensity rim (arrow) is clearly demonstrated surrounding a uterine myometrial mass exhibiting iso to slight high signal intensity compared with the myometrium. (B) The mass shows high signal intensity with a low-intensity rim (arrow) on T2-weighted image. (C) A low-intensity rim (arrow) on susceptibility-weighted imaging (SWI) is more prominent than that on T2-weighted image. Spotty low intensities are also observed within the mass on SWI.
on T₁-weighted images was only 47% (7 of 15 lesions), and the other eight lesions showed entirely high signal intensity compared to the myometrium. Nakai et al. evaluated the MRI and pathological findings of 10 leiomyomas with red degeneration, and the high-intensity rim on T₁-weighted images was found in only four lesions (40%). The other six lesions showed entirely high-intensity on T₁-weighted images and pathological examination revealed that coagulative necrosis without hemorrhage may cause high signal intensity of the lesions, and our results support their observation.

Intra-tumoral necrosis and hemorrhage are suggestive findings for high-grade uterine sarcomas. Because uterine sarcomas such as leiomyosarcoma could also show diffuse high signal intensity due to massive hemorrhagic necrosis, diagnosis of tumors exhibiting diffuse high signal intensity on T₁-weighted images may be problematic. The low-intensity rim on T₂-weighted images was found in 7 of 15 leiomyomas with red degeneration (47%) exhibiting diffuse high signal intensity in our study, and the other eight lesions showed entirely or inhomogeneously low signal intensity so the presence of low-intensity rim was obscure on T₂-weighted images.

SWS such as SWI and SWAN are sensitive MRI techniques for blood products, and could demonstrate hemorrhage of various time phases. SWS may reveal venous thrombus as signal voids due to the T₂* shortening effects of both deoxyhemoglobin and intracellular methemoglobin at the acute phase and hemosiderin at the chronic to obsolete phases of thrombus. In the current study, SWS demonstrated the peripheral low-intensity rim in all cases of leiomyomas with red degeneration. However, probable peripheral venous thrombus was pathologically revealed only in one lesion of leiomyomas with red degeneration, SWS could reveal hemosiderin deposits as remaining traces of obsolete venous thrombus in the other lesions. From the result, SWS may be helpful in distinguishing leiomyomas with red degeneration from usual leiomyomas with degeneration or uterine sarcomas with massive hemorrhagic necrosis by revealing the peripheral low-intensity rim.

The retrospective nature and small population are limitations in this study. Use of multiple field strength and multiple SWS may be another limitations. Further studies in larger populations to verify the results are needed.

**Conclusion**

We conclude that SWS may be helpful for the diagnosis of leiomyomas with red degeneration by revealing characteristic peripheral low-intensity rim.

**Conflicts of Interest**

The authors declare that they have no conflicts of interest.

**References**

arteries compared to TOF-MRA. Eur J Radiol 2012; 81: 1238–1245.


