MRI Appearance of Florid Cystic Endosalpingiosis of the Uterus: a Case Report

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Endosalpingiosis is a non-neoplastic proliferation of ectopic tubal epithelium. It may be found incidentally or the patients may present with chronic pelvic pain. It may resemble a gynecologic malignancy on imaging findings and clinicians and radiologists should be aware of this benign entity to render a correct diagnosis and to avoid over-treatment. We report here the MR imaging appearance of a case of florid cystic endosalpingiosis.

CASE REPORT

A 40-year-old woman presented with chronic pelvic pain and dysfunctional uterine bleeding. The patient had a history of undergoing uterine ablation twice and this had been done due to dysfunctional uterine bleeding about nine and six months prior to the current presentation. The CT scan that had been done elsewhere revealed an irregular soft tissue mass in the pelvis and the mass was indenting the posterolateral aspect of the uterus and cervix. The manual vaginal examination that was performed at presentation revealed a large soft tender mass that was felt in the left fornix. In addition, a pedunculated cervical polyp was also noted; this was removed and the histology showed it to be an endometroid polyp with simple hyperplasia and it was without atypia or focal squamous metaplasia.

A subsequent MRI of the pelvis was done on a 1.5T AVANTO (Siemens, Erlangan, Germany) MRI scanner with the T1 weighted (T1W) high resolution sequence, the T2 weighted (T2W) sequence and the short tau inversion recovery (STIR) sequence in the axial plane and the T1W and T2W fast spin echo sequence in the sagittal plane. The post-contrast T1W images with and without fat suppression were acquired in the axial and sagittal planes. The pre-contrast T1W images were digitally subtracted from the post-contrast T1W images for evaluation.
Florid Cystic Endosalpingiosis of Uterus

Fig. 1. Florid cystic endosalpingiosis of uterus in 40-year-old woman.
A-C. Sagittal T2-weighted (A), axial T2-weighted (B) and axial post-contrast fat suppressed T1-weighted (C) images show heterogeneously enhancing complex cystic mass involving posterior myometrium at uterocervical junction and extending into pouch of Douglas. Discrete cystic lesion is seen in posterior cervical stroma and this appears as being predominantly hyperintense on T2-weighted images (arrow).
D, E. Sagittal post-contrast fat suppressed T1-weighted image (D) shows no significant post-contrast enhancement in discrete cystic lesion in posterior cervical stroma (arrow). Sagittal pre-contrast T1-weighted image (E) shows small areas of hyperintensity interspersed within cystic lesion (arrow). There was no evidence of hemorrhage within lesion on pathologic examination and small areas of hyperintensity were presumed to be mucinous content.
F. Resected specimen of corpus uteri and cervix shows irregular grayish white tissue adherent to posterior serosal surface.
G, H. Microphotograph showing multiple foci of cystically dilated glands in posterior cervical stroma (Hematoxylin & Eosin stain × 100) (G). Microphotograph showing circular gland lined by tubal epithelium. Adjoining cystic structures also noted (H).
The MRI findings revealed a heterogeneously enhancing, complex cystic mass involving the posterior myometrium at the uterocervical junction with an extraserosal extension infiltrating into the pouch of Douglas and a thickening that extended posterolaterally to the upper left to the pararectal region (Fig. 1A–C). A discrete well circumscribed cystic lesion was also noted in the posterior cervical stroma with no appreciable post-contrast enhancement (Fig. 1D). Focal hyperintensity was noted to be interspersed in the discrete well circumscribed cystic lesion, and this focal hyperintensity was presumed to be subacute blood, but there was no evidence of any hemorrhage in the histopathology specimen and this possibly represented mucinous contents (Fig. 1E). There were no enlarged retroperitoneal/pelvic nodes. Our imaging differential diagnosis was endometriosis versus cystic neoplasm. Total abdominal hysterectomy was done together with bilateral salpingo-oophorectomy. The operative findings revealed nodule in the pouch of Douglas with no apparent cystic mass (possibly collapsed) and there was a fibrotic thickening along the left uterosacral ligament extending to the left pararectal area.

Gross examination of the resected specimen revealed irregular grayish white tissue adhered to the posterior cervical wall (Fig. 1F). Both ovaries and both fallopian tubes were unremarkable except for a few simple cysts. Microscopic examination revealed cystic structures lined by the tubal type of epithelium. On examination of the multiple sections from the lesion, these dilated tubal structures were devoid of any endometrial stromal tissue, and this excluded a diagnosis of endometriosis (Fig. 1G, H). Similar histological findings were noted at the corpus uterine serosa, the left cervical serosa and in the pouch of Douglas. The final diagnosis was florid cystic endosalpingiosis.

**DISCUSSION**

Endosalpingiosis is non-neoplastic proliferation of ectopic tubal epithelium (2). According to Zinsser and Wheeler (3), the frequency of endosalpingiosis is up to 12.5%, on the basis of the surgically removed omentum that was histologically examined. These patients are usually asymptomatic and their condition is usually incidentally diagnosed at the time of operation or by microscopic examination of the resected/biopsied specimen. The symptomatic cases may present with chronic pelvic pain (4), as in our case.

Cystic endosalpingiosis has been described on MRI as a well circumscribed, intramural serous fluid-filled unilocular cystic mass in the uterine fundus and as a simple cyst in the right ovary with no hemorrhagic component (5). Multiple disseminated pelvic calcifications in endosalpingiosis have been described on CT (6). Yet we found no evidence of any calcifications on the retrospective review of the CT study that was done prior to MRI. Tumor-like foci of endosalpingiosis have only rarely been described in the urinary bladder (7) and in the vermiform appendix (8).

Florid cystic endosalpingiosis is rare. The pathogenesis of florid cystic endosalpingiosis is largely unknown; however, mullerianization wherein the coelomic epithelium lining of the peritoneal cavity might undergo a change towards primary mullerian epithelium, including tubal, endometrial and endocervical epithelium, has been described (9). Clement and Young (1), in their series of four cases of endosalpingiosis, described the clinical and histopathology findings in a case of florid cystic endosalpingiosis that had tumor-like masses, which were characterized by a polypoidal mass composed of multiple cysts lined by tubular type epithelium, hyperplastic smooth muscle tissue and a myofibromatous stroma. Atypical endosalpingiosis has been reported by the same authors with marked cellular stratification and a varying degree of cellular atypia. No cellular atypia was noted in our case.

The morphology of the lesion in our case resembled a multicystic mass with intrusion into the cervical stroma. Various differential diagnoses have been proposed for cystic cervical masses (10), and these include deep nabothian cysts of the cervix, florid deep glands of the uterine cervix, endometriosis, cystic adenocarcinoma and adenoma malignum. Of the tumor-like lesions, florid deep glands of the uterine cervix can infiltrate the cervical stroma with no evidence of any atypia, as in our case.

Moreover, in our case, a multicystic mass was seen in the pouch of Douglas on imaging. Tubal and tuboendometroid metaplasia may penetrate deeply into the cervical stroma, but it should originate from the endocervical canal, rather than arising from or beneath the serosa (10), as in our case.

We describe this case to demonstrate the MRI appearance of florid cystic endosalpingiosis, which can clinically and radiologically mimic neoplasia. Awareness of the imaging appearance of this malady may broaden differential diagnosis of cystic pelvic masses and help prevent over diagnosis and over treatment.

**References**

1. Clement PB, Young RH. Florid cystic endosalpingiosis with tumor like manifestations: a report of four cases including the first reported case of transmural endosalpingiosis of the uterus. *Am J Surg Pathol* 1999;23:166-175
2. Bazot M, Vacher Lavenu MC, Bigot JM. Imaging of endosalpingiosis. *Clin Radiol* 1999;54:482-485
3. Zinsser KR, Wheeler JE. Endosalpingiosis in the omentum: a study of autopsy and surgical material. *Am J Surg Pathol* 1982;6:109-117
4. deHoop TA, Mira J, Thomas MA. Endosalpingiosis and chronic...
pelvic pain. *J Reprod Med* 1997;42:613-616
5. Cil AP, Atasoy P, Kara SA. Myometrial involvement of tumor-like cystic endosalpingiosis: a rare entity. *Ultrasound Obstet Gynecol* 2008;32:106-110
6. Tutschka BG, Lauchlan SC. Endosalpingiosis. *Obstet Gynecol* 1980;55:S57-S60
7. Young RH, Clement PB. Müllerianosis of the urinary bladder. *Mod Pathol* 1996;9:731-737
8. Cajigas A, Axiotis CA. Endosalpingiosis of the vermiform appendix. *Int J Gynaecol Pathol* 1990;9:291-295
9. Shim SH, Kim HS, Joo M, Chang SH, Kwak JE. Florid cystic endosalpingiosis of the uterus: a case report. *Korean J Pathol* 2008;42:189-191
10. Heatley MK, Russell P. Florid cystic endosalpingiosis of the uterus. *J Clin Pathol* 2001;54:399-400