Acute abdomen in a patient with rudimentary horn

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Abstract
Unicornuate uterus with a rudimentary horn is a rare condition. These patients present with dysmenorrhoea and chronic pelvic pain and infertility. We present an interesting case of hematometra with non-communicating rudimentary horn with unicornuate uterus. Strong suspicion to this condition helps in early diagnosis and treatment.

Key words: rudimentary horn, dysmenorrhea, hematometra

Introduction
Unicornuate uterus with non-communicating rudimentary horn is susceptible to many gynaecological and obstetric complications which can occur at any stage of reproductive life¹. High index of suspicion should be kept in teenagers and young patients presenting with dysmenorrhoea. Unicornuate uterus with a rudimentary horn is a rare type of mullerian duct malformation with a prevalence of 1:10000².

Case report
A 24 year old nulligravida was admitted to the Department of Obstetrics and Gynaecology, of Teerthanker Mahaveer Medical College, Moradabad, (U.P.), India, with the complaint of severe dysmenorrhoea for past 10-12 years along with primary infertility. She attained menarche at the age 13 and she started having dysmenorrhea since 14 years. She was married 4 years back and had been on and off treatment with GP but was not relieved.

Vital signs remained stable. On abdominal examination there was tenderness present in the right iliac fossa. Gynaecological examinations revealed a normal vagina and cervix with a bulky uterus. The uterus was deviated to right side and it was tender. She had an approximately 5×4 cm palpable tender hard mass with restricted mobility in the right adnexal region. Differential diagnosis of chronic endometriosis and chronic ectopic was made. Laboratory values were as follows: haemoglobin 10.5g/dl, haematocrit 37%, white blood cell count 12000 per mm³ platelet count 105000 per mm³. Her β HCG level was not detectable. CA 125 - was 100 mIU. Ultrasonography revealed a 40×38 mm smooth, contoured, homogeneous mass with collection in right adnexal region. MRI was done in which a diagnosis of rudimentary horn with hematometra was made. Sonography for kidneys and intravenous pyelography was done which was normal. Decision for laparotomy and excision of rudimentary horn with ipsilateral salpingectomy was taken.

Peroperatively the left round ligament arose from the left uterine cornua however the right round ligament arose from the rudimentary horn. The right sided non communicating functional rudimentary horn of 5×4 cm in size was attached to the uterus, ovaries were adhered posteriorly and endometrioma of 5×5 cm² was present on right side, which was adhered to the omentum. Pouch of Douglas was completely obliterated. Endometriotic implants were present on the left side and the left tube was blocked due to fimbrial adhesions. Adhesiolysis and right sided semi hysterectomy with right sided salpingectomy was done (Figure 1 and Figure 2), with removal of endometrioma. Adhesiolysis was done on
After Chromo-tubation the left tube was found to be patent. Postoperative period was uneventful and she was treated with GnRH analogues for prevention of endometriosis.

Discussion

Unicornuate uterus with a rudimentary horn is the rarest congenital anatomic anomaly of the female genital system. Urinary tract abnormalities are commonly associated with Mullerian anomalies. This pathology is classified into 4 groups by the American Society of Reproductive Medicine (ASRM) as unicornuate uterus with communicating rudimentary horn, unicornuate uterus with non-communicating rudimentary horn, isolated unicornuate uterus and non-cavitated unicornuate uterus with non-communicating rudimentary horn. Possible causes of abdominal pain in these patients are the distention of the uterus because of blood accumulation in the non-communicating cavity of the rudimentary horn, hematometra, pyometra, and torsion. Another cause of abdominal pain in these women is endometriosis. Retrograde menstruation from the ipsilateral tube results in endometriosis supporting retrograde menstruation theory. Sensitivity of USG is only 26%.

Diagnosis is missed by inexperienced hands. Three dimensional ultrasound may be used for diagnosis of uterine abnormalities. Laparoscopy is the most accurate diagnostic tool, yet MRI has emerged as a useful and non-invasive tool for diagnosis of uterine anomalies. Once diagnosis is strongly suspected, laparoscopy or laparotomy is a must, excision of the rudimentary horn is advised. Literatures show a low preclinical (8%) and preoperative detection rate (29%). Evaluation of renal system is advised because of high incidence of urological anomalies. Surgical removal of the non-communicating horn should be performed once the diagnosis is made especially if it contains functional endometrium, to prevent endometriosis and adverse outcome. In our case, removal of the horn resulted in relief of dysmenorrhoea. The patient is currently undergoing follow up.

REFERENCES