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Contents

Thrice Monthly Volume 10 Number 26 September 16, 2022

REVIEW

Assisting individuals with diabetes in the COVID-19 pandemic period: Examining the role of religious 9180 factors and faith communities

Eseadi C, Ossai OV, Onyishi CN, Ilechukwu LC

9192 Role of octreotide in small bowel bleeding

Khedr A, Mahmoud EE, Attallah N, Mir M, Boike S, Rauf I, Jama AB, Mushtag H, Surani S, Khan SA

MINIREVIEWS

9207 Internet of things-based health monitoring system for early detection of cardiovascular events during COVID-19 pandemic

Dami S

9219 Convergence mechanism of mindfulness intervention in treating attention deficit hyperactivity disorder: Clues from current evidence

Xu XP, Wang W, Wan S, Xiao CF

9228 Clinical presentation, management, screening and surveillance for colorectal cancer during the COVID-19 pandemic

Akbulut S, Hargura AS, Garzali IU, Aloun A, Colak C

Early diagnostic value of liver stiffness measurement in hepatic sinusoidal obstruction syndrome induced 9241 by hematopoietic stem cell transplantation

Tan YW, Shi YC

ORIGINAL ARTICLE

Case Control Study

9254 Local inflammatory response to gastroesophageal reflux: Association of gene expression of inflammatory cytokines with esophageal multichannel intraluminal impedance-pH data

Morozov S, Sentsova T

Retrospective Study

Evaluation of high-risk factors and the diagnostic value of alpha-fetoprotein in the stratification of primary 9264 liver cancer

Jiao HB, Wang W, Guo MN, Su YL, Pang DQ, Wang BL, Shi J, Wu JH

One-half layer pancreaticojejunostomy with the rear wall of the pancreas reinforced: A valuable 9276 anastomosis technique

Wei JP, Tai S, Su ZL



World Journal of Clinical Cases		
Conte	Thrice Monthly Volume 10 Number 26 September 16, 2022	
9285	Development and validation of an epithelial-mesenchymal transition-related gene signature for predicting prognosis	
	Zhou DH, Du QC, Fu Z, Wang XY, Zhou L, Wang J, Hu CK, Liu S, Li JM, Ma ML, Yu H	
	Observational Study	
9303	Incidence and risk factor analysis for swelling after apical microsurgery	
	Bi C, Xia SQ, Zhu YC, Lian XZ, Hu LJ, Rao CX, Jin HB, Shang XD, Jin FF, Li JY, Zheng P, Wang SH	
	CASE REPORT	
9310	Acute carotid stent thrombosis: A case report and literature review	
	Zhang JB, Fan XQ, Chen J, Liu P, Ye ZD	
9318	Congenital ovarian anomaly manifesting as extra tissue connection between the two ovaries: A case report	
	Choi MG, Kim JW, Kim YH, Kim AM, Kim TY, Ryu HK	
9323	Cefoperazone-sulbactam and ornidazole for <i>Gardnerella vaginalis</i> bloodstream infection after cesarean section: A case report	
	Mu Y, Li JJ, Wu X, Zhou XF, Tang L, Zhou Q	
9332	Early-onset ophthalmoplegia, cervical dyskinesia, and lower extremity weakness due to partial deletion of chromosome 16: A case report	
	Xu M, Jiang J, He Y, Gu WY, Jin B	
9340	Posterior mediastinal extralobar pulmonary sequestration misdiagnosed as a neurogenic tumor: A case report	
	Jin HJ, Yu Y, He W, Han Y	
9348	Unexpected difficult airway due to severe upper tracheal distortion: A case report	
	Zhou JW, Wang CG, Chen G, Zhou YF, Ding JF, Zhang JW	
9354	Special epithelioid trophoblastic tumor: A case report	
	Wang YN, Dong Y, Wang L, Chen YH, Hu HY, Guo J, Sun L	
9361	Intrahepatic multicystic biliary hamartoma: A case report	
	Wang CY, Shi FY, Huang WF, Tang Y, Li T, He GL	
9368	ST-segment elevation myocardial infarction in Kawasaki disease: A case report and review of literature	
	Lee J, Seo J, Shin YH, Jang AY, Suh SY	
9378	Bilateral hypocalcaemic cataracts due to idiopathic parathyroid insufficiency: A case report	
	Li Y	
9384	Single organ hepatic artery vasculitis as an unusual cause of epigastric pain: A case report	
	Kaviani R, Farrell J, Dehghan N, Moosavi S	
9390	Congenital lipoid adrenal hyperplasia with Graves' disease: A case report	
	Wang YJ, Liu C, Xing C, Zhang L, Xu WF, Wang HY, Wang FT	



World Journal of Clinical Cases		
Contents Thrice Monthly Volume 10 Number 26 September 16, 2022		
9398	Cytokine release syndrome complicated with rhabdomyolysis after chimeric antigen receptor T-cell therapy: A case report	
	Zhang L, Chen W, Wang XM, Zhang SQ	
9404	Antiphospholipid syndrome with renal and splenic infarction after blunt trauma: A case report	
	Lee NA, Jeong ES, Jang HS, Park YC, Kang JH, Kim JC, Jo YG	
9411	Uncontrolled high blood pressure under total intravenous anesthesia with propofol and remifentanil: A case report	
	Jang MJ, Kim JH, Jeong HJ	
9417	Noncirrhotic portal hypertension due to peripheral T-cell lymphoma, not otherwise specified: A case report	
	Wu MM, Fu WJ, Wu J, Zhu LL, Niu T, Yang R, Yao J, Lu Q, Liao XY	
9428	Resumption of school after lockdown in COVID-19 pandemic: Three case reports	
	Wang KJ, Cao Y, Gao CY, Song ZQ, Zeng M, Gong HL, Wen J, Xiao S	
9434	Complete recovery from segmental zoster paresis confirmed by magnetic resonance imaging: A case report	
	Park J, Lee W, Lim Y	
9440	Imaging findings of immunoglobin G4-related hypophysitis: A case report	
	Lv K, Cao X, Geng DY, Zhang J	
9447	Systemic lupus erythematosus presenting with progressive massive ascites and CA-125 elevation indicating Tjalma syndrome? A case report	
	Wang JD, Yang YF, Zhang XF, Huang J	
9454	Locally advanced cervical rhabdomyosarcoma in adults: A case report	
	Xu LJ, Cai J, Huang BX, Dong WH	
9462	Rapid progressive vaccine-induced immune thrombotic thrombocytopenia with cerebral venous thrombosis after ChAdOx1 nCoV-19 (AZD1222) vaccination: A case report	
	Jiang SK, Chen WL, Chien C, Pan CS, Tsai ST	
9470	Burkitt-like lymphoma with 11q aberration confirmed by needle biopsy of the liver: A case report	
	Yang HJ, Wang ZM	
9478	Common carotid artery thrombosis and malignant middle cerebral artery infarction following ovarian hyperstimulation syndrome: A case report	
	Xu YT, Yin QQ, Guo ZR	
9484	Postoperative radiotherapy for thymus salivary gland carcinoma: A case report	
	Deng R, Li NJ, Bai LL, Nie SH, Sun XW, Wang YS	
9493	Follicular carcinoma of the thyroid with a single metastatic lesion in the lumbar spine: A case report	
	Chen YK, Chen YC, Lin WX, Zheng JH, Liu YY, Zou J, Cai JH, Ji ZQ, Chen LZ, Li ZY, Chen YX	



Carata	World Journal of Clinical Cases
Conter	Thrice Monthly Volume 10 Number 26 September 16, 2022
9502	Guillain-Barré syndrome and hemophagocytic syndrome heralding the diagnosis of diffuse large B cell lymphoma: A case report
	Zhou QL, Li ZK, Xu F, Liang XG, Wang XB, Su J, Tang YF
9510	Intravitreous injection of conbercept for bullous retinal detachment: A case report
	Xiang XL, Cao YH, Jiang TW, Huang ZR
9518	Supratentorial hemangioblastoma at the anterior skull base: A case report
	Xu ST, Cao X, Yin XY, Zhang JY, Nan J, Zhang J
	META-ANALYSIS
9524	Certain sulfonylurea drugs increase serum free fatty acid in diabetic patients: A systematic review and meta-analysis
	Yu M, Feng XY, Yao S, Wang C, Yang P
	LETTER TO THE EDITOR
9536	Glucose substrate in the hydrogen breath test for gut microbiota determination: A recommended noninvasive test
	Xie QQ, Wang JF, Zhang YF, Xu DH, Zhou B, Li TH, Li ZP
9539	A rare cause of acute abdomen after a Good Friday
	Pante L, Brito LG, Franciscatto M, Brambilla E, Soldera J
9542	Obesity is associated with colitis in women but not necessarily causal relationship
	Shen W, He LP, Zhou LL
9545	Risk stratification of primary liver cancer
2010	Tan YW



Contents

Thrice Monthly Volume 10 Number 26 September 16, 2022

ABOUT COVER

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The primary aim of World Journal of Clinical Cases (WJCC, World J Clin Cases) is to provide scholars and readers from various fields of clinical medicine with a platform to publish high-quality clinical research articles and communicate their research findings online.

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CASE REPORT

Congenital ovarian anomaly manifesting as extra tissue connection between the two ovaries: A case report

Myeong Gyun Choi, Jong Woon Kim, Yoon Ha Kim, A Mi Kim, Tae Young Kim, Hyun Kyung Ryu

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Abstract

BACKGROUND

Ovarian anomalies except for uni- or bilateral streak gonads are rare. We present a rare case of an ovarian anomaly in which both ovaries were connected by extra tissue.

CASE SUMMARY

A 32-year-old, primipara with a twin pregnancy at 36 weeks of gestation was admitted to the hospital with severe preeclampsia. She underwent emergency cesarean section owing to persistent headache, blurred vision, and general edema. Following a peritoneal incision, a thin rectangular-shaped tissue was seen in front of the uterus. After delivery, the extra tissue was removed; no other anomalies were reported in either the ovaries or uterus. Pathology results of the removed tissue disclosed a well-vascularized loose stromal tissue with few follicles and scattered luteinized cells. In this case, to prevent pelvic adhesion or intestinal obstruction resulting from volvulus, strangulation, and torsion, the extra tissue was removed.

CONCLUSION

We report a case of a rare ovarian anomaly where both ovaries were connected by extra tissue. If the extra tissue extends to the abdominal cavity, it should be removed to prevent pelvic adhesion or abdominal complications including intestinal volvulus, strangulation, and torsion.

Key Words: Connected ovaries; Extra tissue; Ovarian anomaly; Case report

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Core Tip: Herein we present the case of a rare ovarian anomaly where both ovaries were connected by extra tissue. If the extra tissue extends to the abdominal cavity, it should be removed to prevent pelvic adhesion or abdominal complications including intestinal volvulus, strangulation, and torsion.

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INTRODUCTION

Several ovarian anomalies have been reported. Among them is ovarian absence; in phenotypic females, the absence of both ovaries is generally linked to chromosome abnormalities and gonadal dysgenesis syndrome. In this case, individuals are likely to have streak gonads or underdeveloped gonads, which are a risk factor for malignancy[1]. Congenital unilateral ovarian agenesis in a normal female is extremely rare and asymptomatic in most cases. Ipsilateral renal or ureteric agenesis and/or ipsilateral malformation of the fallopian tube may be accompanied by congenital unilateral ovarian agenesis. The etiology of unilateral ovarian agenesis has yet to be explained. The two most likely causes of unilateral ovarian agenesis are an asymptomatic torsion of the ovary with consequent organ ischemia and reabsorption or a defect in the development of the Mullerian and gonadal structures underlying vascular anomalies[2]. Another example of ovarian anomalies is ectopic ovaries; whether accessory or supernumerary, ectopic ovaries are also extremely rare, 1 in 29000 to 1 in 70000 gynecologic admissions [3], and may be associated with other congenital genitourinary abnormalities[1]. An accessory ovary contains ovarian tissue and is usually connected to a normal ovary. In contrast, supernumerary ovaries are not attached to the ovary but may be found at various sites in or outside the pelvis. In this report, we present a rare case of an unreported anomaly in which both ovaries were connected by extra tissue.

CASE PRESENTATION

Chief complaints

A 32-year-old female presented with high blood pressure.

History of present illness

The primipara woman with a twin pregnancy at 36 weeks of gestation was admitted to the hospital with high blood pressure and proteinuria. Her blood pressure was 160/100 mmHg, while laboratory test results showed 3+ proteinuria. The pregnancy followed a successful *in vitro* fertilization-embryo transfer.

History of past illness

The patient had no history of past illness.

Personal and family history

The patient had no specific history of genetic diseases.

Physical examination

She underwent emergency caesarean section owing to persistent headache, blurred vision, and general edema. Following a peritoneal incision, a thin rectangular-shaped tissue was seen in front of the uterus (Figure 1); it formed a connection between the two ovaries (Figure 2). We displaced this tissue, incised the uterus, and delivered the fetuses.

Laboratory examinations

There were no specific findings related to laboratory examinations.

Imaging examinations

There was no specific findings observed on imaging examinations.

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Figure 1 Thin rectangular-shaped tissue in front of the uterus (arrow).



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Figure 2 The tissue connected to both ovaries at the posterior view of the uterus after a cesarean section.

FINAL DIAGNOSIS

Congenital ovarian anomaly manifesting as an extra tissue connection between the two ovaries.

TREATMENT

After delivery, we set the margins of both ovaries to avoid injuring the normal ovaries. First, both ends of the extra tissue were ligated and excised. Then the extra tissue was removed; no other abnormal findings were observed in both the ovaries and uterus. We explored the lower abdomen as much as possible to check for extra ovary tissues and any other malformations such as renal anomalies, but there were no specific findings. Pathology results of the removed tissue disclosed a well-vascularized loose stromal tissue with few follicles and scattered luteinized cells (Figure 3).

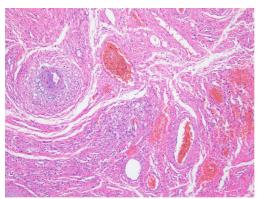
OUTCOME AND FOLLOW-UP

After surgery, the patient recovered and was discharged on the third postoperative day. During an outpatient follow-up after one month, the patient had no abdominal symptoms and ultrasonography revealed no abnormal findings on both adnexa.

DISCUSSION

Several studies have reported numerous cases of ovarian anomalies including bilateral and unilateral ovary absence, accessory, and supernumerary ovary. However, there are no reports on cases of connected ovaries, and to the best of our knowledge, this is the first case report. Our patient was





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Figure 3 Well-vascularized loose stromal tissue with few follicles and scattered luteinized cells (x100).

admitted to the hospital with severe preeclampsia. She did not present other symptoms such as abdominal pain or pelvic pain and the pregnancy followed a successful in vitro fertilization-embryo transfer. She underwent an emergency caesarean section owing to severe headache, blurred vision, and general edema. A tissue connection between both ovaries was discovered by chance. In vitro fertilization is associated with several complications including ovarian hyperstimulation syndrome (characterized by swollen and painful ovaries), which results from the use of injectable fertility drugs, such as human chorionic gonadotropin (HCG)[4]. However, there seems to be no association between connected ovaries and ovulation induction complications. The extra tissue attached to an ovary may be asymptomatic and is not associated with infertility. In most cases, it may be left untreated with observation. However, as in the present case, if both ovaries are connected and a rectangular-shaped tissue lies in the abdominal cavity, this extra tissue can cause pelvic adhesion or intestinal volvulus, strangulation, and torsion resulting in intestinal obstruction[5]. Therefore, the extra tissue should be removed.

CONCLUSION

Herein we report the case of a rare ovarian anomaly where both ovaries were connected by extra tissue. In this case, to prevent pelvic adhesion or intestinal obstruction resulting from volvulus, strangulation, and torsion, the extra tissue was removed.

FOOTNOTES

Author contributions: Choi MG wrote the manuscript, and edited all its revisions; Kim AM, Ryu HK and Kim TY retrieved the data, assisted in writing and editing the manuscript; Kim JW and Kim YH participated in designing the study, retrieved the data, assisted in writing the manuscript, and edited all its revisions; all authors have read and approved the final manuscript.

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