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Anaesthetic Management of Pre Eclamptic Patient with Solitary Kidney and Uterine Didelphys Posted for Emergency Caesarean Section: A Case Report

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Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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Case Study

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ABSTRACT

Uterine didelphys results from impaired fusion of the paired Müllerian ducts. The incidence of uterine anomalies is believed to be 0.5–2.0% of reproductive-age women, with didelphic uterus renal agenesis accounting for approximately 10%. Uterine didelphys is associated with in approximately 25% of cases¹. Pre eclampsia is defined as SBP more than 160 mmhg, DBP more than 90 mmhg, associated with proteinuria and urine protein excretion >300mg in a 24 hour period or a protein creatinine ratio of atleast 0.3². Neuraxialblockade stands an effective mode of anaesthesia for these patients. Management of pre eclampticpatient with solitary kidney can be a challenge to anaesthesiologist due tovarious metabolic derangements including hyperkalemia, hypocalcemia, hyperphosphatemia and metabolic acidosis.Multidisciplinary approach is required to have good pregnancy outcome in these patients.

Keywords: Preeclampsia; uterine didelphys; solitary kidney.

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1. INTRODUCTION

Congenital solitary kidney is a disorder caused by an abnormal development of one of the two kidnevs. This abnormality can be either anatomical (unilateral renal agenesis) or functional (extreme forms of dysplasia)Uterine didelphys is a type of Mullerian duct anomaly where there is complete duplication of uterine horns as well as cervix with no communications in between [1-5]. Pre eclampsia is defined as SBP more than 160mmhg, DBP more than 90 mmhg associated with proteinuria and urine protein excretion >300mg in a 24 hour period or a protein creatinine ratio of atleast 0.3 [6-8]. We report a case of 21 years old pre eclamptic patient with uterine didelphys and solitary kidney to undergo Caeserean section.

2. CASE REPORT

A 20 years old G2A1@30 weeks +3 days of gestational age with uterine didelphys with

solitarv kidnev(diagnosed in previous abortion with ultrasound) and gestational hypertension on treatment with Tab. labetalol Tabdepine 100 mg, 10 mg, andIni. Dexamethasone 6 mg admitted for safe confinement of pregnancy 2 days back. and Basic investigations done found within normal limits. Her NST was reactive and showed fetal tachycardia, she had bilateral pitting pedal edema so she was administered with Pritchard regimen and she was posted for Emergency Caeserean section.

Under aseptic precautions, patient in sitting position, using 26 gauge quincke needle, spinal anaesthesia was administered. Patient made supine immediately and oxygen given via facemask at 6l/minute. During the procedure, the patient's heart rate maintained around 80 bpm and MAP maintained above 60 and lasted for one and half hour, postop uneventful.





Fig. 1-4. Ultrasound images

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3. DISCUSSION

Patient with solitary kidney can develop various metabolic derangements including hyperkalemia. hypocalcemia. hyperphosphatemia and metabolic acidosis. Judicious fluid management, maintenance of normovolemia and avoidance of hypotension are priorities for the successful prevention of intra op renal injury for patient with solitary kidney .In patient who had hypertension and proteinuria held a high risk for progression to renal insufficiency. General anaesthesia for Caeserean delivery in pre eclamptic women has increased risk of developing intra cranial hemorrhage and stroke. Thus neuraxial blockade does avoid the possibility of difficult intubation secondary to airway edema. Nonetheless there are situations in which general anaesthesia is the best anaesthetic option like severe ongoing hemorrhage, maternal sustained fetal bradycardia with a reassuring maternal airway examination and severe thrombocytopenia or other coagulopathy or a combination of these indications. But Spinal anaesthesia in contrary cardiovascular effects by sympathetic has blockade that extends beyond the level of sensory blockade by 2 to 6 segments. Bradycardia and hypotension may occur .In this condition Ephedrine hydrochloride can be used which increases the heart rate and blood pressure at the same time [9-11]. Increased production of circulating factors with potent press or effect and the increased sensitivity to vasopressor drugs in pre eclampsia along with the use of hyperbaric bupivacaine with opioids could decrease spinal induced hypotension in parturients. preeclamptic Cardiac output monitoring after spinal anaesthesia has shown that neither spinal anaesthesia nor the use of phenyl ephedrine to treat hypotension decrease cardiac output during Caesarean section delivery further supporting its safety in pre eclampticparturients [15-14].

4. CONCLUSION

Patients with solitary kidney due to congenital causes or nephrectomy need to be counsel regarding risks of developing pyelonephritis, preeclampsia and its associated complications during pregnancy. These patients require close monitoring throughout their antenatal and postpartum period to avoid any deterioration of renal function. Multidisciplinary approach is required to have good pregnancy outcome in these patients.

ETHICAL APPROVAL

As per international standard or university standard written ethical approval has been collected and preserved by the author(s).

CONSENT

As per international standard or university standard, patients' written consent has been collected and preserved by the author(s).

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

- Madureira AJ, Mariz CM, Bernardes JC, Ramos IM. Case 94: Uterus didelphys with obstructing hemivaginal septum and ipsilateral renal agenesis. Radiology 2006 May;239(2):602-606. DOI:10.1148/radiol.2392031187 [PubMed] [CrossRef] [Google Scholar]
- Kim TE, Lee GH, Choi YM, Jee BC, Ku SY, Suh CS, et al. Hysteroscopic resection of the vaginal septum in uterus didelphys with obstructed hemivagina: a case report. J Korean Med Sci 2007 Aug;22(4):766-769.

DOI:10.3346/jkms.2007.22.4.766 [PMC free article] [PubMed] [CrossRef] [Google Scholar]

- Nahum GG. Uterine anomalies. How common are they, and what is their distribution among subtypes? J Reprod Med. 1998 Oct;43(10):877-887 [PubMed] [Google Scholar]
- Golan A, Langer R, Bukovsky I, Caspi E. Congenital anomalies of the müllerian system. Fertil Steril. 1989 May;51(5):747-755 [PubMed] [Google Scholar]
- 5. Smith NA, Laufer MR. Obstructed hemivagina and ipsilateral renal anomaly (OHVIRA) syndrome: management and follow-up. FertilSteril. 2007 Apr;87(4):918-922.

DOI:10.1016/j.fertnstert.2006.11.015 [PubMed] [CrossRef] [Google Scholar]

 Shavell VI, Montgomery SE, Johnson SC, Diamond MP, Berman JM. Complete septate uterus, obstructed hemivagina, and ipsilateral renal anomaly: pregnancy course complicated by a rare urogenital anomaly. Arch GynecolObstet 2009. Sep;280(3):449-452. DOI: 10.1007/s00404-008-0919-6 [PubMed] [CrossRef] [Google Scholar]

 Burgis J. Obstructive Müllerian anomalies: case report, diagnosis, and management. Am J ObstetGynecol 2001. Aug;185(2):338-344. DOI:10.1067/mob.2001.116738 [PubMed]

[CrossRef] [Google Scholar]

- García González P, MeanaMorís AR, GracíaChapullé A, Matesanz Pérez JL. The role of MRI in congenital cystic lesions in the pelvis: a case of uterus didelphys with double vagina, hematocolpos, and ipsilateral renal agenesis. Radiologia. 2009 Mar-Apr;51(2):194-197 DOI:10.1016/j.rx.2009.01.001 [PubMed] [CrossRef] [Google Scholar]
- Rock JA, Breech LL. Surgery for anomalies of the Mullerian ducts. In: Rock JA. And Thompson JD, editor, Te Linde Operative Gynecology, 9thed, Philadelphia:Lippincott Wilkins. 2003;746-8. [Google Scholar]
- Rehman A, Hasan Z, Amanat S, Shaukat T, Saeed A, Jamil K, et al. Combined persistent mullerian duct syndrome, Transverse Testicular Ectopia and Mosaic

Klinefelter's Syndrome. J Coll Physicians Surg Pak. 2008. Jun;18(6):375-377 [PubMed] [Google Scholar]

- Kaufman Y, Lam A. The pelvic uterus-like mass-a primary or secondary Müllerian system anomaly? J Minim Invasive Gynecol. 2008 Jul- Aug;15(4):494-497. DOI:10.1016/j.jmig.2008.03.002 [PubMed] [CrossRef] [Google Scholar]
- 12. Shulman LP. Müllerian anomalies. Clin Obstet Gynecol. 2008 Jun;51(2):214-222 DOI:10.1097/GRF.0b013e31816feba0 [PubMed] [CrossRef] [Google Scholar]
- 13. Jindal G, Kachhawa S, Meena GL, Dhakar Uterus didelphys with unilateral G. hemivagina obstructed with hematometrocolpos and hematosalpinx with ipsilateral renal agenesis. J Hum Jul;2(2):87-89. 2009. Reprod Sci. DOI:10.4103/0974-1208.57230 [PMC free article] [PubMed] [CrossRef] [Google Scholar]
- Olpin JD, Heilbrun M. Imaging of Müllerian duct anomalies. Clin Obstet Gynecol. 2009. Mar;52(1):40-56. DOI:10.1097/GRF.0b013e3181958439 [PubMed] [CrossRef] [Google Scholar]

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