Case Report

Website: www.ijhsr.org ISSN: 2249-9571

Unusual Presentation of Posterior Reversible Encephalopathy Syndrome After Spinal Anesthesia for Cesarean Section: A Case Report

Georges N. Mezher¹, M. Yasser. M. Abdel Azziz.²

¹M.D: Instructor, Department of Anesthesia and Reanimation, Faculty of Medical Sciences, Lebanese University, Anesthesiology Consultant, Alkindi Hospital, Manama, Kingdom of Bahrain.

²MD: Anesthesiology Specialist, Alkindi Hospital, Manama, Kingdom of Bahrain.

Corresponding Author: Georges Nicole Mezher

DOI: https://doi.org/10.52403/ijhsr.20250214

ABSTRACT

Background: Headache after spinal anesthesia (SA) is usually attributed to post dural puncture headache (PDPH). We report a case of posterior reversible encephalopathy syndrome (PRES) after spinal anesthesia to a normotensive pregnant patient presented for cesarean section (CS).

Case presentation: A 33-year-old pregnant woman underwent a CS under SA. The patient was obese, had gestational diabetes mellitus with a history of two spontaneous abortions and had a history of one uneventful CS. Immediately after the SA, the patient developed an acute severe headache relieved with medical treatment.

The patient complained after 8 hours postoperatively of headache, sudden blurred vision which was followed by a generalized tonic-clonic seizure treated by intravenous midazolam and magnesium sulfate. The headache and visual disturbance resolved completely one hour later. The MRI suggested the diagnosis of PRES.

Conclusion: We hereby contribute to the literature an uncommon neurological disorder, the PRES, occurred in a normotensive parturient.

Keywords: Posterior reversible encephalopathy syndrome, headache, pre-eclampsia, eclampsia, spinal anesthesia, post dural puncture headache.

INTRODUCTION

Spinal anesthesia is the most commonly used anesthesia technique for cesarean delivery. Post spinal puncture headache (PSPH) is a well-known complication of SA. It is a common and incapacitating complication following dura-arachnoid puncture [1, 2]. In addition, PSPH may occur immediately after puncture [3].

The differential diagnosis of headaches in pregnancy includes the primary headaches (Tension type headaches, Migraine, Cluster

headache) [4,5], and the secondary headaches (Hypertension/Pre-eclampsia, Idiopathic intracranial hypertension, Subarachnoid hemorrhage. Cerebral venous thrombosis. Reversible cerebral vasoconstriction syndrome, Posterior reversible encephalopathy syndrome) [6, 7, 8, 9]. The pathology of PRES is related most commonly pregnancy-related to hypertensive disorders. Other conditions could induce the onset of PRES normotensive patients like systemic lupus

chronic renal erythematosus, failure. rheumatoid arthritis, immune suppressive medications, anti-neoplastic agents, severe hypercalcemia, thrombocytopenic syndromes, Henoch-Schonlein purpura, hemolytic syndrome, amyloid uremic angiopathy, various causes of renal failure, regional anesthesia, sepsis, toxic agents especially chemotherapy, and drugs such as cocaine and methamphetamine [10, 11].

PRES is a rare clinic and neuroradiological entity introduced as late as 1996 by Hinchey et al ^[12]. Clinical features include headache, encephalopathy, seizures, and cortical visual disturbances or blindness. Neuroimaging shows parieto-occipital white and grey matter change ^[13, 14]. We report a rare case of PRES to a normotensive pregnant patient post cesarean section under SA.

CASE REPORT

A 33-year-old woman (gravida 4, para 1), ASA III is admitted for elective CS. According to the estimated due date of ultrasound in the first semester, the gestation age was 39 weeks and 3 days. The patient is obese (BMI= 31.59 Kg/m²) and gestational diabetes has controlled by metformin. The parturient has no hypertensive disorder, no comorbidities, and no history of smoking or substance abuse. Her obstetric history was uneventful CS under spinal anesthesia four years ago and two miscarriages under general anesthesia.

The patient never showed any trace of proteinuria. The blood laboratory investigations made before the CS were normal. Pre-spinal anesthesia non-invasive blood pressure was 118 mmHg systolic and 67 mmHg diastolic. The heart rate was 108 beats per minute, the respiratory rate was 16 per minute, and her oxygen saturation was 100% on nasal oxygen 4 l/minute. The SA was uneventful except of the occurrence of

a severe headache 5 minutes after. The patient's vitals remained within the normal range. A normal weight female baby was extracted (APGAR score at 1 minute and 5 minutes were respectively equal to 8 and 10). Paracetamol and midazolam were administered intravenously and a total of 1000 ml of Ringer's lactate was infused during the entire procedure which lasted for 1 hour. Postoperatively, she was shifted to a post-anesthesia care unit where her headache resolved partially for a pain visual scale equal to 3.

8 hours after delivery, the patient reported experiencing a sudden blurred vision and recurrence of her headache rated on the visual scale equal to 8. The patient's vitals remained unchanged and the neurological exam was normal. The pupils were equal and reactive to light, the fundoscopy was normal. A complete blood count (CBC), coagulation profile, LDH, blood sugar, serum electrolytes, liver function test (LFT), renal function test (RFT), urine analysis, and MRI of head were ordered.

The patient developed few minutes after a generalized tonic-clonic seizure treated by intravenous midazolam and magnesium sulfate. The headache and visual disturbance resolved completely within one hour. Blood tests and urine analysis suggested no abnormalities. These findings were followed by a MRI head which showed small foci of signal abnormality affecting the grey and white matter of both cerebellar hemispheres, without restriction of diffusion or associated brain edema. The findings were suggestive of PRES. Fig 1.

The patient was explained about the diagnosis and potential reversibility. She was transferred to the intensive care unit for 24 hours where she was treated with analgesics and magnesium sulfate and kept on strict clinical monitoring.

MIC BRAIN PLA
AX TZW. TI
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17 1
11/17/2024, 11:24:17

Fig 1. MRI image showing T2 high signal intensity in the cerebellar hemispheres (yellow arrows)

DISCUSSION

Complaint of headache after SA arouses the suspicion of post-dural puncture headache. It is due to dural and arachnoid puncture. The onset of post spinal headache puncture may be immediately. Therefore, the patient was managed for post spinal puncture headache PSPH.

The parturient maintained a normal blood pressure throughout her pregnancy and peripartum periods. Hence, hypertensive etiologies for headache were less likely to be suggested.

However, the occurrence of the visual disturbance preceded by a severe headache and followed, few minutes later by tonic-clonic seizure raised the possibility of PRES. In fact, PRES is a rare and serious entity of the central nervous system, characterized by headaches, seizures, altered mental status, and visual impairment [15]. Proteinuria may be absent [16].

PRES frequently associated is hypertensive gestation (gestational hypertension, pre-eclampsia, eclampsia). However, PRES could rarely occur to a normotensive pregnant woman [17]. PRES may be caused by several conditions which include acute or chronic renal diseases, uremic syndrome, hemolytic use cytotoxic and immunosuppressant drugs, blood transfusion, electrolyte disturbances, and regional anesthesia [18].

Findings of PRES on neuroimaging are common in women with neurological symptoms (headaches, vision disturbance, seizures). It reveals edema involving the white matter in the posterior portions of the cerebral hemispheres especially in the parieto-occipital regions The pathological condition in PRES relies mainly on two theories; The disruption of the cerebrovascular autoregulation caused by acute fluctuations of systemic blood pressure. In fact, the susceptible posterior areas of the cerebral hemispheres have a reduced density of sympathetic innervation in the posterior, compared to the anterior, circulation, the latter being more densely innervated by the superior cervical ganglion [21]. The second theory is that the syndrome is triggered by endothelial dysfunction caused by circulating endogenous exogenous toxins [13].

Cerebral venous thrombosis may also share the same critical presentation as that of PRES. Hence, it must be excluded via imaging tools since it is the most frequent cerebrovascular disorder in the puerperium [19]

Administration of midazolam and magnesium sulfate intravenously treated the neurologic symptoms and prevented the recurrence of seizures. The visual deficits from PRES usually fully recover [20].

The prognosis of PRES is good. In general, the patient improves with symptomatic treatment, with no adverse effects ^[21]. With adequate treatment 70-90% of people with PRES make a full recovery within hours to days ^[22]. Of those who have residual symptoms after PRES, this attributable

largely to hemorrhage ^[23]. In our case the patient experienced complete recovery of her symptoms.

CONCLUSION

The relevance of this case report is due to the rare occurrence of posterior reversible encephalopathy syndrome to a normotensive parturient after spinal anesthesia for cesarean section. Although the pathophysiology is not well established, the prognosis is usually good.

Declaration by Authors Acknowledgement: None

Source of Funding: None

Conflict of Interest: The authors declare no conflict of interest.

REFERENCES

- Cook TM, Counsell D, Wildsmith JA; Royal College of Anaesthetists Third National Audit Project. Major complications of central neuraxial block: report on the Third National Audit Project of the Royal College of Anaesthetists. Br J Anaesth. 2009 Feb;102(2):179-90. doi: 10.1093/bja/aen360. Epub 2009 Jan 12. PMID: 19139027.
- Rodgers A, Walker N, Schug S, McKee A, Kehlet H, van Zundert A, Sage D, Futter M, Saville G, Clark T, MacMahon S. Reduction of postoperative mortality and morbidity with epidural or spinal anaesthesia: results from overview of randomised trials. BMJ. 2000 Dec 16;321(7275):1493. doi: 10.1136/bmj.321.7275.1493. PMID: 11118174; PMCID: PMC27550.
- 3. Mathew G, Agha R, Albrecht J, Goel P, Mukherjee I, Pai P, D'Cruz AK, Nixon IJ, Roberto K, Enam SA, Basu S, Muensterer OJ, Giordano S, Pagano D, Machado-Aranda D, Bradley PJ, Bashashati M, Thoma A, Afifi RY, Johnston Challacombe B, Ngu JC, Chalkoo M, Raveendran K, Hoffman JR, Kirshtein B, Lau WY, Thorat MA, Miguel D, Beamish AJ, Roy G, Healy D, Ather HM, Raja SG, Mei Z, Manning TG, Kasivisvanathan V, Rivas JG, Coppola R, Ekser B, Karanth VL, Kadioglu H, Valmasoni M, Noureldin A: **STROCSS** Group. **STROCSS** 2021: Strengthening the reporting of cohort, cross-

- sectional and case-control studies in surgery. Int J Surg. 2021 Dec; 96:106165. doi: 10.1016/j.ijsu.2021.106165. Epub 2021 Nov 11. PMID: 34774726.
- Silberstein SD, Young NB. Headache and Facial Pain. In Textbook of Clinical Neurology 3rd edn. Goetz CG 2007. Saunders Elsevier Philadelphia 2007.
- 5. Revell K, Morrish P. Headaches in Pregnancy. The Obstetrician & Gynecologist 2014; 16: 179-184. doi: 10.1111/tog.12101.
- 6. Ferro JM, Canhão P, Stam J, Bousser MG, Barinagarrementeria F; ISCVT Investigators. Prognosis of cerebral vein and dural sinus thrombosis: results of the International Study on Cerebral Vein and Dural Sinus Thrombosis (ISCVT). Stroke. 2004 Mar;35(3):664-70. doi: 10.1161/01.STR.0000117571.76197.26. Epub 2004 Feb 19. PMID: 14976332.
- 7. Mehraein S, Ortwein H, Busch M, Weih M, Einhäupl K, Masuhr F. Risk of recurrence of cerebral venous and sinus thrombosis during subsequent pregnancy and puerperium. J Neurol Neurosurg Psychiatry. 2003 Jun;74(6):814-6. doi: 10.1136/jnnp.74.6.814. PMID: 12754362; PMCID: PMC1738512.
- 8. Sattar A, Manousakis G, Jensen MB. Systematic review of reversible cerebral vasoconstriction syndrome. Expert Rev Cardiovasc Ther. 2010 Oct;8(10):1417-21. doi: 10.1586/erc.10.124. PMID: 20936928; PMCID: PMC3020907.
- Digre KB. Headaches during pregnancy. Clin Obstet Gynecol. 2013 Jun;56(2):317-29. doi: 10.1097/GRF.0b013e31828f25e6. PMID: 23563877.
- 10. Ansari B, Saadat Nia, M. Prevalence and Risk Factors of Posterior Reversible Encephalopathy Syndrome in Isfahan, Iran. Adv Biomedical Res. 2021; 10:53. doi: 10. 53. 10.4103/abr.abr_243_19.
- 11. Anderson RC, Patel V, Sheikh-Bahaei N, Liu CSJ, Rajamohan AG, Shiroishi MS, Kim PE, Go JL, Lerner A, Acharya J. Posterior Reversible Encephalopathy Syndrome (PRES): Pathophysiology and Neuro-Imaging. Front Neurol. 2020 Jun 16; 11:463. doi: 10.3389/fneur.2020.00463. PMID: 32612567; PMCID: PMC7308488.
- 12. Hinchey J, Chaves C, Appignani B, Breen J, Pao L, Wang A, Pessin MS, Lamy C, Mas JL, Caplan LR. A reversible posterior

- leukoencephalopathy syndrome. N Engl J Med. 1996 Feb 22;334(8):494-500. doi: 10.1056/NEJM199602223340803. PMID: 8559202.
- Bartynski WS, Boardman JF. Distinct imaging patterns and lesion distribution in posterior reversible encephalopathy syndrome. AJNR Am J Neuroradiol. 2007 Aug;28(7):1320-7. doi: 10.3174/ajnr. A0549. PMID: 17698535; PMCID: PMC7977645.
- 14. Casey SO, Sampaio RC, Michel E, Truwit CL. Posterior reversible encephalopathy syndrome: utility of fluid-attenuated inversion recovery MR imaging in the detection of cortical and subcortical lesions. AJNR Am J Neuroradiol. Aug;21(7):1199-206. PMID: 10954269: PMCID: PMC8174901.
- 15. Kastrup O, Gerwig M, Frings M, Diener HC. Posterior reversible encephalopathy syndrome (PRES): electroencephalographic findings and seizure patterns. J Neurol. 2012 Jul;259(7):1383-9. doi: 10.1007/s00415-011-6362-9. Epub 2011 Dec 22. PMID: 22189837.
- Zhang Y, Liang B, Zhao C, Zhou Y, Yan C. Posterior reversible encephalopathy in a pregnant woman without preeclampsia or eclampsia: A case report. Medicine (Baltimore). 2022 Sep 9;101(36): e30519. doi: 10.1097/MD.0000000000030519. PMID: 36086692; PMCID: PMC10980412.
- Gewirtz AN, Gao V, Parauda SC, Robbins MS. Posterior Reversible Encephalopathy Syndrome. Curr Pain Headache Rep. 2021 Feb 25;25(3):19. doi: 10.1007/s11916-020-00932-1. PMID: 33630183; PMCID: PMC7905767.
- Stott VL, Hurrell MA, Anderson TJ.
 Reversible posterior leukoencephalopathy syndrome: a misnomer reviewed. Intern Med J. 2005 Feb;35(2):83-90. doi:

- 10.1111/j.1445-5994.2004.00750. x. PMID: 15705136.
- 19. Zhang L, Wang Y, Shi L, Cao J, Li Z, Wáng YX. Late postpartum eclampsia complicated with posterior reversible encephalopathy syndrome: a case report and a literature review. Quant Imaging Med Surg. 2015 Dec;5(6):909-16. doi: 10.3978/j.issn.2223-4292.2015.12.04. PMID: 26807372; PMCID: PMC4700241.
- Rao N, Raychev R, Kim D, Liebeskind D. Elucidating the Mechanism of Posterior Reversible Encephalopathy Syndrome. Neurologist. 2012; 18 (6): 391-4. doi:10.1097/NRL.0b013e31826a9954.
- 21. Fugate JE, Rabinstein AA. Posterior reversible encephalopathy syndrome: clinical and radiological manifestations, pathophysiology, and outstanding questions. Lancet Neurol. 2015 Sep;14(9):874. doi: 10.1016/S1474-4422(15)00195-7. PMID: 26184985.
- 22. Liman TG, Siebert E, Endres M. Posterior reversible encephalopathy syndrome. Curr Opin Neurol. 2019 Feb;32(1):25-35. doi: 10.1097/WCO.0000000000000040. PMID: 30531559.
- 23. Tetsuka S, Ogawa T. Posterior reversible encephalopathy syndrome: A review with emphasis on neuroimaging characteristics. J Neurol Sci. 2019 Sep 15; 404:72-79. doi: 10.1016/j.jns.2019.07.018. Epub 2019 Jul 17. PMID: 31349066.

How to cite this article: Georges N. Mezher, M. Yasser. M. Abdel Azziz. Unusual presentation of posterior reversible encephalopathy syndrome after spinal anesthesia for cesarean section: a case report. *Int J Health Sci Res.* 2025; 15(2):116-120. DOI:

https://doi.org/10.52403/ijhsr.20250214
