

## CASE REPORT

# Anesthetic management of a case of a large atrial septal defect with mild pulmonary hypertension for emergency cesarean section

V. R. R Chari, MD\*, Chhaya M. Suryawanshi, MD\*, Preety Sahu\*\*

*\*Professor, \*\*Postgraduate Resident*

*Dept of Anesthesiology, Padm. Dr. D. Y. Patil Medical College College & Research Centre, Sant Tukaram Nagar, Pimpri, Pune-18, Maharashtra-411018 (India)*

**Correspondence:** Dr. Preety Sahu, Dept of Anesthesiology, Padm. Dr. D. Y. Patil Medical College College & Research Centre, Sant Tukaram Nagar, Pimpri, Pune-18, Maharashtra-411018 (India); Phone: 08087582578; E-mail: preety.sahu@yahoo.com

## ABSTRACT

Atrial septal defect (ASD) is the most common congenital acyanotic heart disease in adults and accounts for 10% of congenital cardiac defects in adults. It is the most commonly seen congenital cardiac lesion in women of child-bearing age and the pregnancy is usually well tolerated. Pulmonary hypertension is defined as a mean pulmonary arterial pressure greater than 25mm Hg at rest or greater than 30mm Hg during exercise. We report a case of a large ASD with mild pulmonary hypertension in a patient who underwent emergency caesarean section under general anaesthesia for failure to progress. During the general anaesthesia for the procedure our objectives were to avoid hypotension, hypoxaemia, hypercarbia, hypothermia, reversal of shunt (Eisenmenger's syndrome) and fluid overload. The patient had an eventful perioperative course and discharged from the hospital on the 8<sup>th</sup> postoperative day in good physical condition.

**Keywords :** Atrial septal defect; Pulmonary arterial hypertension; Caesarean section; hypotension; hypoxemia; hypercarbia; Eisenmenger's syndrome

**Citation:** Chari VRR, Suryawanshi CM, Sahu P. Anesthetic management of a case of a large atrial septal defect with mild pulmonary hypertension for emergency cesarean section. *Anaesth Pain & Intensive Care* 2013; 17(2):182-184

## INTRODUCTION

Atrial septal defect (ASD) is the most common congenital acyanotic heart disease in adults and accounts for up to 10% of cases of congenital acyanotic cardiac defects in adults (you need a reference here). There are three distinct types of ASD namely, ostium primum, ostium secundum and sinus venosus defect. The ostium secundum accounts for 70% of ASD cases, with a male:female ratio 1:2<sup>1</sup>. ASD causes left to right intra- cardiac shunt with right ventricular volume overload, increased pulmonary blood flow (PBF), pulmonary hypertension, right ventricular hypertrophy and eventually congestive heart failure (CHF). We present a parturient mother known to have ostium secundum type of ASD since childhood and who underwent emergency caesarean section under general (intra venous) anaesthesia for failure to progress. Anaesthetic management and perioperative strategies are discussed.

## CASE REPORT

A 26-years- old primigravida, in labor at 37 weeks of gestation with failure to progress during labor was

transferred from a private hospital and emergency cesarean section was planned. Patient was a known case of ostium secundum type of ASD since childhood with the medical history of repeated upper respiratory infections (URTI), moderate breathlessness (NYHA II), but no history of chest pain, palpitation, or fatigue. On physical examination pulse was 88/min (Regular), BP was 130/90 mmHg, bilateral pitting pedal edema was present, but the jugular venous pressure (JVP) was not raised on clinical examination. CVS examination revealed pansystolic murmur in the pulmonary and mitral area (suggesting mild mitral regurgitation), wide split and fixed second heart tone (S2). The rest of the physical examination was unremarkable. All the biochemical, haematological and coagulation tests were within normal limits. ECG showed right axis deviation, incomplete RBBB. 2D ECHO revealed grossly dilated left and right atrium and right ventricle, large ostium secundum ASD (24 mm in Diameter), left to right shunt, mild pulmonary hypertension with an estimated right ventricle systolic pressure of 40 mmHg and ejection fraction of 60%.

### Anesthetic management:

The patient was transferred to the operating room; a peripheral intravenous (IV) catheter was inserted and lactated ringer's solution was started at maintenance. Heart rate, non-invasive BP, ECG, SpO<sub>2</sub> and EtCO<sub>2</sub> were monitored throughout the anaesthesia. As it was an emergency situation, invasive monitoring was not done and fluid assessment was done clinically.

Patient was premedicated with IV glycopyrrolate 0.2 mg, metoclopramide 20 mg, ondansetron 4 mg, ranitidine 150 mg. The patient was then preoxygenated with 100% oxygen for 3 mins and then induced with propofol 50 mg, ketamine 50 mg, and suxamethonium chloride 100 mg. The patient was intubated orally with a size 7.0 mm cuffed endotracheal tube. Rocuronium 40 mg was given as a neuromuscular blockade agent. The right hip was elevated with a soft pillow in order to prevent supine hypotension syndrome. Anaesthesia was maintained with oxygen 40%, N<sub>2</sub>O 60%, isoflurane (0.6-0.8%), and controlled ventilation. EtCO<sub>2</sub> was maintained around 32-35 mmHg.

After the delivery of a 2.8 kg female baby with an Apgar scores of 8 (1min) and 10 (5min), oxytocin infusion (15 Units in 500ml saline) was administered over 20 minutes followed by fentanyl (50 mcg) by slow IV push. The procedure lasted 45 minutes. Intraoperatively pulse was 90-110 beats/min, BP was 110/60-140/90 mm Hg, SpO<sub>2</sub>-100%. Lactated ringer solution 500 ml and 0.9% sodium chloride were given during the anaesthesia and the urine output was 150 ml during the 45 min procedure. The estimated blood loss during the procedure was 400ml.

For reversal of neuromuscular block neostigmine (2.5 mg)+glycopyrrolate (0.5 mg) given. To have adequate spontaneous respiratory efforts neostigmine (0.5 mg) was repeated. This was done to avoid residual neuromuscular block and to improve recovery. Post extubation patient was conscious, responding to verbal commands, normal muscle power, stable BP, HR and SpO<sub>2</sub>. Patient was monitored in recovery room for 1 hour and then transferred to ICU and managed with ICU protocols and discharged on 8th postoperative day. Her hospital stay was uneventful.

### DISCUSSION

We present a case of secundum ASD in a parturient patient who successfully underwent emergency caesarean section for failure to progress. We employed the usual anaesthetic approach with particular attention to control of factors that may lead sudden pulmonary hypertension with resultant hypoxemia and the potential for the development of acute heart failure, which would have been detrimental to the mother and the fetus.

There are three types of ASD. The sinus venosus type occurs high in the atrial septum near the entry of superior

vena cava (SVC) into the right atrium(RA) and is frequently associated with anomalous pulmonary venous connection from the right lung to SVC/RA. Ostium primum anomalies lie adjacent to AV valves either of which may be deformed / regurgitant. Ostium secundum ASD, which accounts for 70% of cases of ASD, involves the fossa ovalis and is midseptal in location.<sup>11</sup>

It enables blood flow between the left and right atria via the inter-atrial septum, which may not be clinically significant. Patients are usually asymptomatic during childhood. However, symptoms appear later in life and by age 40, 90% of untreated patients have symptoms of exertional dyspnea, fatigue, palpitation or sustained arrhythmia.<sup>3</sup> Complications of uncorrected secundum type of ASD include pulmonary arterial hypertension, right sided heart failure, atrial fibrillation/flutter, stroke and Eisenmenger's syndrome.<sup>4</sup>

Large ASD (>9mm), may result in a clinically remarkable left-to-right shunt. This extra blood from the left atrium may cause a volume overload of both the right atrium and ventricle. If untreated, it can result in enlargement of the right side of the heart and ultimately heart failure. Changes in SVR during the perioperative period have important implications for patients with ASD.<sup>5</sup> The magnitude of left to right shunt depends on size of ASD, ventricular diastolic properties and the relative impedance in pulmonary and systemic circulation.<sup>6</sup>

The echocardiography can establish the size and location of the ASD, magnitude and hemodynamic impact of the left to right shunt, and the presence and the degree of pulmonary hypertension.<sup>7,8</sup> Pulmonary hypertension is classified as mild (36-49 mm of Hg systolic), moderate (50-59 mm of Hg), severe (>60 mm of Hg) based on the estimated right ventricular systolic pressure gradient calculated from Doppler Echocardiography.<sup>4</sup>

Anticipated problems during general anaesthesia include, Air embolism during vascular access, dysrhythmias (5-10%), heart failure, heart block and infective endocarditis.

Anaesthetic techniques commonly used for lower segment caesarean section (LSCS) include regional anaesthesia i.e spinal / epidural anaesthesia. However, these techniques carry the risk of sudden and uncontrolled hypotension and unstable hemodynamics with the possibility of reversal of intra cardiac shunt.

We preferred general anaesthesia (GA) with controlled ventilation because our patient had arrived in emergency and patient was in labour and in agony. GA provides better haemodynamic stability,

100% oxygen can be given, as the patient had mild pulmonary hypertension. Hypercarbia can be avoided by mechanical ventilation and adjusting tidal volume and respiratory rate. Inflation of the lungs with intermittent

## Anesthetic management of a case of a large atrial septal defect

positive pressure ventilation (IPPV) leads to release of endogenous nitric oxide and prostaglandins which cause pulmonary vasodilatation.<sup>10</sup> IPPV of lungs is well tolerated when there is increased pulmonary blood flow.<sup>5</sup> Intraoperatively we remained vigilant to avoid hypothermia, hypercarbia and hypoxaemia were avoided as these factors can cause reversal of shunt.

Our aim during GA was to maintain adequate preload, avoid increase in the shunt fraction, maintain optimum cardiac contractility, maintain normal SVR and PVR (achieved in this case by administration of IV ketamine), near normal heart rate and close monitoring to detect atrial arrhythmias.

Balanced GA in this clinical scenario is unlikely to cause drastic changes in SVR / PVR with minimum effect on

shunt. GA will maintain systemic blood flow within normal limits so that pharmacokinetics of inhaled anesthetic is not altered. Intravenously administered medications may be diluted due to increased pulmonary blood flow, however this is unlikely to alter the clinical response to these medications because the pulmonary circulation time is very short. Perioperative antibiotic cover was provided to prevent sub acute bacterial endocarditis.

## CONCLUSION

In patients with ASD we should tailor anesthesia technique to avoid increase in pulmonary vascular resistance, decrease in systemic vascular resistance, hypotension, tachycardia, hypoxia, hypercarbia and hypothermia.

## REFERENCES

1. Feldt RH, Avasthey P, Yoshimasu F, Kurland LT, Titus JL. Incidence of congenital heart disease in children born to residents of Olmsted County, Minnesota, 1950-1969. *Mayo Clin Proc.* 1971 Dec; 46(12):794-799. [PubMed]
2. Kuczkowski KM. Labour analgesia for parturient with cardiac disease - what does an obstetrician need to know? *ActaObstetGynecolScand* 2004 ;83: 223-33. [PubMed]
3. Saxena A, Divekar A, Soni NR. Natural history of secundum atrial septal defect revisited in the era of transcatheter closure . *Indian Heart J.* 2005;57:35-38. [PubMed]
4. Balint OH, Samman A, Haberer K, Tobe L, McLaughlin P, Siu SC, Horlick E, Granton J, Silversides CK. Outcomes in patients with pulmonary hypertension undergoing percutaneous ASD closure. *BMJHeart* 2008;94:1189-1193. [PubMed]
5. Hines RL, Marschall KE. In: Livingstone C, editor. *Stoelting's anesthesia and co-existing disease.* 6th edition. India: Elsevier ,2008 .p.60.
6. Siu SC, Sermer M, Colman JM, Alvarez AN, Mercier LA, Morton BC et al. Prospective multicentre study of pregnancy outcomes in womens with heart disease. *Circulation*104.2001:515-521. [PubMed] [Free Full Text]
7. Valdes-Cruz LM, Cayre RO. Anomalies of right ventricular outflow tract and pulmonary arteries. In: Valdes-Cruz LM, Cayre RO, editors. *Echocardiographic diagnosis of congenital heart disease.* Philadelphia: Lippincott-Raven ; 1999.p.325-348.
8. Lee MG, Ko JS. An unusual presentation of an atrial septal defect . *J Cardiovasc Ultrasound* 2009;17(4):151-152. [PubMed] [Free Full Text]
9. Weiss BM, Hess OM. Pulmonary vascular disease and pregnancy: current controversies, management strategies, and perspectives. *Eur Heart J* 2000; 21:104-115. [PubMed] [Free Full Text]
10. Morray JP, Lynn AM, Mansfield PB. Effect of pH and PaCO<sub>2</sub> on pulmonary and systemic haemodynamics after surgery in children with congenital heart disease and pulmonary hypertension. *J Paediatr* 1998;113:474-479. [PubMed]
11. Brickner ME, Hillis LD, Lange RA: Congenital heart disease in adults. *N Engl J Med* 2000; 342:256-263. [PubMed] [Free Full Text]

