Factor XII deficiency inhibits fibrinolysis and therefore can lead to thromboembolic complications.\textsuperscript{5} Patients with isolated factor XII deficiency have a high rate of venous thromboembolism and arterial thrombosis, leading to life threatening complications such as pulmonary embolism and myocardial infarction.\textsuperscript{6} In addition, surgical trauma and immobilization increase the risk of thromboembolic complications. This necessitates the use of thromboprophylaxis perioperatively (mechanical and pharmacological) and early aggressive mobilization postoperatively.\textsuperscript{7} We used IPCD intraoperatively and in the postoperative period, low molecular weight heparin, early mobilization, and daily screening for DVT was done in addition to continued use of IPCD.

In conclusion, patients with factor XII deficiency are at high risk of thromboembolism for which vigilant monitoring and measures for thromboprophylaxis should be undertaken perioperatively. The risk of bleeding, although present, is not so alarming. The prophylactic correction of prolonged aPTT should be given a second thought.

Conflict of Interest
None declared.

References

Dilated Cardiomyopathy and Prone Position: An Anesthetic Challenge

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Abstract

The anesthetic management of a patient with dilated cardiomyopathy (DCM) for non-cardiac surgery is challenging due to associated congestive heart failure, malignant dysrhythmias, sudden cardiac arrest, implanted rhythm devices, and thromboembolism. We report successful conduct of a case of DCM on cardiac resynchronization device with Cauda equina syndrome (CES) under general anesthesia in prone position. The anesthetic concerns specific to the pathophysiology of DCM are also discussed.

Keywords

- cardiac resynchronization therapy device
- dilated cardiomyopathy
- prone position

Case Report

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Introduction

Dilated cardiomyopathy (DCM) has an incidence of 5 to 8 per 100,000 and is characterized by decreased contractility, decreased cardiac output, and increased left ventricular (LV) filling pressures due to dilatation of one or both ventricles.1,2 There is progressive ventricular dilatation, eventually leading to congestive heart failure, thromboembolism, valvular regurgitant lesions, malignant arrhythmias, and sudden cardiac death. Patients of DCM are commonly on cardiac resynchronization therapy (CRT) device by means of biventricular pacing (CRT-P), automated internal cardioverter defibrillator (AICD), or a combo device (CRT-D with antiarrhythmia function). Surgery in the prone position in DCM patients with reprogrammed CRT-P (pacing in VOO mode) further increases the chances of perioperative cardiac failure, arrhythmias, and sudden cardiac arrest.3

We report the case of a 52-year-old (body mass index [BMI]–21.4 kg/m²) hypertensive and diabetic female with DCM on Medtronic’s CRT-P for severe systolic dysfunction, posted for urgent laminectomy L4 level with L4–5 discectomy for intradural lesions, malignant arrhythmias, and sudden cardiac death. The patient was under cardiology workup since the past 10 years for DCM. CRT-P was placed as her ejection fraction (EF) dropped to 25% 3 years ago and she became progressively limited in her physical capacity. After CRT-P implantation, she was in New York Heart Association (NYHA) class II and there was no evidence of congestive heart failure. Two-dimensional (2D) echocardiography showed LV EF of 40% with global hypokinesia, elevated left ventricle end-diastolic pressure (LVEDP) (E/A ratio 0.67), left ventricular end-diastolic dimension (LVEDD) of 65 mm, and relative wall thickness 0.4, all suggestive of combined LV systolic dysfunction (mild) and grade I diastolic dysfunction. The rest of the investigations were unremarkable. The patient was on losartan, aspirin, carvedilol, aldactone, and metformin. All medications were continued perioperatively; angiotensin-receptor blocker (ARB), losartan was however stopped 24 hours prior to avoid intraoperative hypotension. Preoperatively, pacemaker interrogation and resetting to VOO mode with a pacing rate of 85 bpm, availability of automated external defibrillator (AED), arterial monitoring, central venous access and preparation of vasopressor and inotrope infusions were ensured. Written informed consent was obtained and nil per oral (NPO) status was confirmed. Standard monitoring was ensured with additional right radial artery cannulation for heat-to-beat blood pressure (IBP) monitoring and central venous cannulation for the purpose of vasopressor medications. Baseline blood gas parameters were assessed for any acid base and electrolyte abnormalities. The volume assessment was done using stroke volume variation (SVV) as measured by Vigileo monitor (Vigileo; FloTrac; Edwards; Lifesciences, Irvine, CA, USA) using third-generation software (version 3.02), aiming for SVV values < 12. Automated external defibrillator (AED) pads were placed in an anterior–posterior configuration to ensure that cardioversion if attempted would not cause pacemaker lead dislodgement/malfunction while allowing synchronized cardioversion/defibrillation in the prone position. The right internal jugular vein (IJV) was cannulated under ultrasound guidance and infusion of noradrenaline and dobutamine were started at the rate of 0.05 μg/kg/min and 5 μg/kg/min, respectively before induction. GA was administered with IV fentanyl 100 μg, titrated dose of etomidate 15 mg and vecuronium 5 mg IV. Postintubation heart rate (HR) and blood pressure (BP) remained within 10% of the baseline. Anesthesia was maintained with desflurane in air–oxygen mixture to allow rapid titration in order to keep hemodynamics stable. On turning the patient prone, BP dropped drastically to 80/48 mm Hg from 150/96 mm Hg and cardiac rhythm deteriorated to atrial fibrillation (AF). With the AED pads in place and significant hypotension, a synchronized cardioversion with 25 J was attempted. The rhythm soon reverted to pacing rhythm of 85 beats per minute and infusion rates of noradrenaline and dobutamine were adjusted to keep BP and HR within 10% of the baseline. The patient’s BP fluctuated a lot in first 20 to 25 minutes after making prone, and frequent adjustments in infusion rates were needed. The rest of the perioperative period was uneventful. Anticipating similar drop in hemodynamics on turning patient supine at the conclusion of surgery, vasopressor and inotropic infusions were overtitrated; intuitively, the hemodynamics stayed stable and patient was extubated fully awake. Postoperatively, the patient was monitored in the intensive care unit (ICU), and infusion of noradrenaline and dobutamine tapered off by postoperative evening. Pacing was resumed to the preoperative DDD mode with CRT-D in the ICU, and the patient was discharged after 5 days with complete recovery of neurologic function.

Discussion

Patients with DCM have a complicated pathophysiology with elevated LV filling pressures, myocardial contractile dysfunction, and tendency for malignant arrhythmias and sudden cardiac arrest. Additionally, these patients are precarious balanced on an inverse relationship between stroke volume and afterload. Therefore, the anesthetic goals are to maintain normovolemia, avoid increases in ventricular afterload, prevent myocardial depression, and avoid overdosage of drugs during induction as circulation time is slow in these patients.3 Further, preoperative symptomatic diastolic dysfunction is associated with a higher risk of postoperative major adverse cardiac events (MACEs), the risk being more with a grade III diastolic dysfunction.4 Commonly, practicing anesthesiologists pay more attention to the preoperative EF as a measure of LV systolic function for cardiac risk stratification, while overlooking the diastolic dysfunction, particularly heart failure with preserved ejection fraction (HFpEF). There is a strong relationship between diastolic dysfunction, HFpEF-induced LA structural remodeling and AF, emphasizing avoidance of acute changes in HR, and hemodynamic overload. Therefore, titration of anesthesia should be approached with altered pharmacodynamics at the core, employing one-third to one-half of the anesthetic dosages at induction and using cardio stable drugs, to account for the slow transit time and reduced cardiac reserve.5,6
Literature research shows that infusion of noradrenalin and dobutamine before induction is effective in countering the depressive effect of anesthetic drugs. Further, cardio-stable and short-acting drugs such as etomidate, fentanyl, and vecuronium are employed as part of anesthetic regime. Noninvasive continuous cardiac output monitoring is used to evaluate the ventricular performance in response to fluid therapy, inotropes, and prone positioning. Prone position leads to increased intrathoracic pressure reducing venous return, and decrease in cardiac output (CO) and cardiac index (CI), which is exaggerated in DCM patients.

A noteworthy point with regard to pacing in these patients: preoperative conversion to asynchronous mode of CRT-P is necessary, in order to prevent pacemaker inhibition by electromechanical interference and use of bipolar cautery and also prevent pacemaker malfunction. For AICD, since these patients are not pacemaker-dependent, simple deactivation of the ICD intraoperatively may suffice. Additionally, AED pads should be placed before induction in the anterior–posterior configuration, so that the cardioversion/defibrillation current would not cross the pacemaker path.

**Conclusion**

Anesthetic management of patients with DCM on CRT device poses unique challenge for the anesthesiologist, with strong likelihood of catastrophic hemodynamic perturbations, dysrhythmias, and even sudden cardiac arrest. Particularly in surgery in prone position with limited access to the patient in the event of a catastrophe, meticulous planning, vigilant monitoring, adequate preparation with anterior–posterior configuration of transthoracic pads, judicious use of pharmacological agents, and tailor-made anesthetic regime, can lead to a favorable outcome.

**Conflict of Interest**

None declared.

**References**


**Case Report**

Management of a Difficult Airway Scenario in a Case of Hurler’s Syndrome with a D-Blade Video Laryngoscope

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