Aggressive Anesthetic Management of a Patient with Severely Dilated Cardiomyopathy for Non Cardiac Surgery

Abstract

We report a case of a 37 year old man, with history of rhabdomyosarcoma as a child. Patient developed severe dilated cardiomyopathy secondary to the treatment of the cancer. He presented for resection of the small bowel. An intra-aortic balloon pump (IABP) was placed preoperatively for mechanical cardiac support. A combined general/epidural technique was used for the surgery. Transesophageal echocardiogram (TEE) was used to monitor patients cardiac status intraoperatively. Patient remained hemodynamically stable and tolerated the procedure well.

Keywords: Dilated cardiomyopathy; Intra aortic balloon pump; Epidural; Hemodynamics; Noncardiac surgery; Chemotherapy induced cardiomyopathy; Chemotherapy; TEE; Transesophageal echo; Intraoperative echo; Aggressive management

Abbreviations: IABP: Intra-Aortic Balloon Pump; TEE: Transesophageal Echocardiogram; LV: Dilated Left Ventricular; MR: Mitral Regurgitation

Introduction

A case of a patient with severe dilated cardiomyopathy, undergoing small bowel resection. Anesthesia was managed with a combination of general/epidural anesthesia, in addition to IABP and TEE.

Case Report

A 37 year old white male presented to our surgical department for small bowel resection. Patient’s history was significant for rhabdomyosarcoma, diagnosed at the age of five. The rhabdomyosarcoma was treated, but recurred four years later. Patient’s treatment regimen included Adriamycin. On this presentation, patient’s medical history was significant for malignancy in the ileostomy that was a result of radiation enteritis. In addition, as a result of the treatment regimen, patient has developed severe non-ischemic dilated cardiomyopathy. Preoperative transthoracic echocardiogram was significant for severely dilated left ventricular (LV) cavity, LV ejection fraction of <10% and severe mitral regurgitation (MR).

On the day of surgery, a left radial arterial line was placed for close and accurate monitoring of hemodynamics. Unsuccessful attempts were made to cannulate both the internal jugular and the left subclavian veins. All three sites had been cannulated multiple times over the years. Patient had an indwelling catheter in the right subclavian. An epidural catheter was placed, both for intraoperative use and post operative pain control. General anesthesia was induced, intravenously, through an 18 gauge angiocath in the right upper extremity. After a smooth induction of anesthesia, the trachea was intubated and the position of the endotracheal tube confirmed.

After emptying the stomach, the TEE probe was placed without difficulty. IABP was placed via the right femoral artery. Position of the IABP was confirmed with the TEE. Operation of the IABP was confirmed. A left femoral venous cordis was placed and central access was established. Preoperative TEE found an LV ejection fraction of approximately 5%, severely dilated LV (Figure 1a & Figure 1b), moderately dilated right ventricle, and moderate to severe right ventricular hypokinesis. Patient had a 4+ MR, with significant reversal of the systolic component of the right superior pulmonary venous profile (Figures 1c & Figure 1d).

Figure 1a & 1b: Transgastric Short Axis view of the left ventricle showing severely dilated left ventricle.

During the surgery, anesthesia was maintained with narcotics, inhalational agent and the epidural. Patient remained hemodynamically stable during the procedure. No significant changes were noted on the TEE during the surgery. IABP functioned adequately, providing mechanical support.
cardiotoxicity can range from arrhythmias to nonspecific ECG changes to a reduction in LV ejection fraction [1]. Endomyocardial biopsy remains the definitive test for diagnosing anthracycline-induced toxicity [4].

Risk factors for developing irreversible anthracycline-induced cardiotoxicity include high dose infusions, high cumulative doses, extremes of age (like our patient) and history of cardiac disease. History of previous mediastinal radiation therapy and the concurrent use of chemotherapeutic agents like paclitaxel or trastuzumab can contribute to anthracycline cardiotoxicity [5-8]. Incidence of cardiotoxicity ranges from 0.4 to 41% [1].

In the acute form, cardiomyopathy manifests as myocarditis-pericarditis, usually within 24 hours of the infusion, with good long term prognosis. In it’s subacute form, it may manifest as late as 30 months after Adriamycin therapy, with 60% mortality. Late form cardiomyopathy may not be seen until four to 20 years later. This form is associated with echocardiographic and pathologic changes, in addition to heart failure [9].

Severity of cardiac dysfunction in our patient warranted aggressive anesthetic management. Although there are reports of using TEE in patients with severely dilated cardiomyopathy, [10,11] it was of paramount importance for our case, since we were unable to place a pulmonary artery catheter preoperatively for adequate hemodynamic monitoring. As it was critical not to overload this patient with fluids, TEE helped with monitoring the volume status. In our literature search, we did not come across any other case of severely dilated cardiomyopathy in which IABP was utilized to provide mechanical cardiac support. In our case, the afterload reduction, provided by the IABP, played a big role in the hemodynamic stability of this patient during the surgery. Placement of the epidural catheter, made it possible for us to limit our use of general anesthetics and avoid unnecessary depression of myocardial function.

Conclusion

At the conclusion of the surgery, patient was adequately reversed and extubated on the operating table. Patient was awake, alert, breathing spontaneously and pain free. Since the patient still had an IABP in place, he was transported to the cardiac intensive care unit for further postoperative care. Patient’s postoperative course remained uneventful.

References

1. Swerdlow S, Campo E, Harris N, Jaffe E, Pileri S, et al. World Health Organization (WHO) (2008) classification of tumours of haematopoietic and lymphoid tissues. Fourth Edition.

2. Jares P, Colomer D, Campo E (2007) Genetic and molecular pathogenesis of mantle cell lymphoma: perspectives for new targeted therapeutics. Nat Rev Cancer 7(10): 750-762.

3. Bosch F, Jares P, Campo E, Lopez-Guillermo A, Piris MA, et al. (1994) PRAD-1/cyclin D1 gene over expression in chronic lymphoproliferative disorders: a highly specific marker of mantle cell. Blood 84(8): 2726-2732.

4. Bosch F, Lopez-Guillermo A, Campo E, Ribera JM, Conde E, et al. (1998) Mantle cell lymphoma: presenting features, response to therapy, and prognostic factors. Cancer 82(3): 567-575.

5. Schniederjan SD and Osunkoya AO (2009) Lymphoid neoplasms of the urinary tract and male genital organs: a clinicopathological...
study of 40 cases. Mod Pathol. 22(8): 1057–1065.

6. Monfared A, Orangpoor RO, Fakheri TF, Babahakkar S (2009) Acute Renal Failure and Bilateral Kidney Infiltration as the First Presentation of non-Hodgkin Lymphoma. Iran J Kidney Dis 3(1): 50-53.

7. Obrador GT, Price B, O’Meara Y, Salant DJ (1997) Acute renal failure due to lymphomatous infiltration of the kidneys. J Am Soc Nephrol 8(8): 1348-1354.

8. Pras M, Franklin EC, Shibolek S, Frangione B (1992) Amyloidosis associated with renal cell carcinoma of the AA type. Am J Med 73(3): 426-428.

9. Chin KC, Perry GJ, Dowling JP, Thompson NM (1999) Primary T-cell-rich-cell lymphoma in the kidney presenting with acute renal failure and a second malignancy. Pathology 31(4): 325-327.

10. Davies J, Healey DA, Wood KM, Jones K, Kanagasundaram NS, et al. (2007) Acute renal failure due to mantle cell lymphoma-a case report and discussion of the literature. Clin Nephrol 67(6): 394-396.

11. Çolak N, Dedik C, Canbakam B, Odabas A, Akyürek N, et al. (2007) Acute tubulo-interstitial nephritis associated with mantle cell lymphoma presented as acute renal failure. Nephrology 12(1): 107-110.

12. Rault RL, Holley JL, Banner BF, el-Shahawy M (1992) Glomerulonephritis and Non-Hodgkin’s Lymphoma: A report of Two Cases and Review of the Literature. Am J Kidney Dis 20(1): 84-89.

13. Karim M, Hill P, Pillai G, Gatter K, Davies DR, et al. (2004) Proliferative Glomerulonephritis associated with mantle cell lymphoma-natural history and effect of treatment in 2 cases. Clin Nephrol 61(6): 422-428.

14. Da’as N, Pollack A, Cohen Y, Amir G, Darman D, et al. (2001) Kidney involvement and renal manifestations in non-Hodgkin’s lymphoma and lymphocytic leukemia: a retrospective study in 700 patients. Eur J Hematol 67(3): 158-164.

15. Rerolle JP, Thervet E, Beaufils H, Vincent F, Rousset P, et al. (1999) Crescentic Glomerulonephritis and centrocytic lymphoma. Nephrol Dial Transplant 14(7): 1744-1745.

16. Rosas SE, Tomaszewski JE, Feldman HI, Foster MH (1999) Membranoproliferative glomerulonephritis type I, mixed cryoglobulinemia and lymphoma in the absence of hepatitis C infection. Am J Nephrol 19(5): 599-604.

17. Smith KD, Alpers CE (2005) Pathogenic mechanisms in Membranoproliferative Glomerulonephritis. Curr Opin Nephrol Hypertens 14(4): 396-403.

18. Korbert SM, Schwartz MM, Lewis EJ (1991) Immunotactoid glomerulopathy. Am J Kidney Dis 17(3): 247-257.

19. Alpers CE, Rennke HG, Hopper J Jr, Biava CG (1987) Fibrillary Glomerulonephritis: An entity with unusual immunofluorescence features. Kidney Int 31(3): 781-789.

20. Alshayeb H, Wall BM (2009) Non Hodgkin’s Lymphoma associated Membranoproliferative glomerulonephritis: rare case of long term remission with chemotherapy: a case report. Cases J 2: 7201.