All that Clots Is Not Blood: Bone Cement Implantation Syndrome Presenting as an Intracardiac Mass and Pulmonary Embolism

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Abstract

Bone cement implantation syndrome (BCIS) is a rare and potentially fatal perioperative complication of cemented orthopaedic surgery. A computed tomography pulmonary angiogram and echocardiogram images were acquired from an 88-year-old patient who had a perioperative collapse while undergoing a revision operation for a peri-implant fracture of the right femoral neck. Findings were suggestive of an intracardiac clot connected to a saddle pulmonary artery embolus. The patient also developed disseminated intravascular coagulation. Overall findings were suggestive of BCIS.

Keywords

► thrombosis
► surgery
► disseminated intravascular coagulation

Bone cement implantation syndrome (BCIS) is a rare and potentially fatal perioperative complication of cemented orthopaedic surgery. A computed tomography pulmonary angiogram and echocardiogram images were acquired from an 88-year-old patient with a peri-implant fracture of the right femoral neck. A new implant was inserted with bone cement. During cementation, she developed a cardiac arrest with pulseless electrical activity.

CT-pulmonary angiogram showed the intracardiac mass connected to a saddle pulmonary artery embolus. This was seen on a noncontrasted view, consistent with a radio-opaque bone cement (►Fig. 1). The transthoracic echocardiogram on the apical four-chamber view showed a 2.5 cm pedunculated mass transiting between the right atrial and right ventricle (►Fig. 2, arrow). This mass was seen on the right ventricular outflow tract view and was connected to an echo-dense structure in the pulmonary artery (►Fig. 3, arrow). McConnell’s sign was demonstrated, which is a regional pattern of right ventricular dysfunction and is an indicator of the presence of a pulmonary embolus.1 The patient also developed severe coagulopathy with a prolonged prothrombin time of 22.9 seconds, an activated partial thromboplastin time of 72.9 seconds, a decreased fibrinogen of 0.69 g/L, and an elevated D-dimer of 5.5 mg/L suggestive of disseminated intravascular coagulation.

BCIS is a rare, perioperative thrombotic phenomenon which hematologists should consider as a clinical factor and management principles defer slightly from traditional venous thromboembolism. The syndrome is characterized by hypoxia, hypotension, increased pulmonary vascular resistance, and cardiac arrest. Associated thrombotic masses have commonly been reported.2 The mechanism for thrombosis is postulated to be multifactorial, including cement embolization, complement activation, and histamine release.3

Management of BCIS is mainly supportive based on acute critical care principles. As the contributor of cardiovascular collapse is predominately right ventricular cardiac failure, fluid resuscitation and vasopressor support should be considered. Alpha-adrenergic agents are preferred. Extracorporeal membrane oxygenation may also be considered for patients with very unstable hemodynamics. The role and efficacy of anticoagulation is not clear.3 It may also be difficult to be administered as patients often develop disseminated intravascular coagulation. Surgical or percutaneous embolectomy can be considered but usually reserved for...
patients with central thrombus. The risk factors for BCIS include patient factors, e.g., old age, osteoporosis, and surgical factors, e.g., revision surgery. In light of the potentially devastating clinical consequences, for patients at high risk, preventative interventions such as increased monitoring at periods of high risk and use of cementless or low-viscosity cement prosthesis should be considered.

The patient demised 12 hours postoperation.

Author Contributions
All authors have accepted responsibility for the entire content of this manuscript and approved its submission. M. A. Cheong and K. M. Chew wrote the manuscript.

Conflict of Interest
None declared.

References