Use of Intravenous Hydroxocobalamin without Methylene Blue for Refractory Vasoplegic Syndrome After Cardiopulmonary Bypass

DEFG 1 Vincent Peyko
ABDE 2 Michael Finamore

Conflict of interest: None declared

Case series

Patients: Male, 71-year-old • Male, 71-year-old
Final Diagnosis: Vasoplegic syndrome
Symptoms: Refractory hypotension
Medication: —
Clinical Procedure: Cardiopulmonary bypass
Specialty: Anesthesiology

Objective: Unusual or unexpected effect of treatment
Background: Cardiac vasoplegic syndrome is a form of vasodilatory shock characterized by profound vasodilation and low systemic vascular resistance, which results in significant hypotension despite high cardiac output and appropriate fluid resuscitation. In up to 45% of patients, cardiopulmonary bypass (CPB) can precipitate vasoplegic syndrome. Vasoplegic syndrome after CPB that is refractory to other vasopressors, such as catecholamine and vasopressin, has been successfully treated with inhibitors of the nitric oxide (NO) system, such as methylene blue and hydroxocobalamin. Methylene blue has been the treatment of choice because of its effectiveness for both prevention and rescue therapy. Hydroxocobalamin has demonstrated efficacy in combination with methylene blue, and also on its own when vasoplegic syndrome is refractory to methylene blue.

Case Report: We present 2 cases that expand upon the existing evidence supporting the efficacy of hydroxocobalamin as a first-line option for inhibiting the NO system in vasoplegic syndrome that is refractory to other vasopressors. Specifically, we demonstrate the appropriate and successful use of hydroxocobalamin alone to treat refractory vasoplegic syndrome after CPB.

Conclusions: Refractory vasoplegic syndrome that occurs after CPB has been successfully treated with inhibitors of the NO system, such as methylene blue and hydroxocobalamin. The present cases expand upon the scant existing evidence of the efficacy of hydroxocobalamin as an appropriate option for refractory vasoplegic syndrome.

Keywords: Cardiopulmonary Bypass • Hydroxocobalamin • Methylene Blue • Vasoplegia

Full-text PDF: https://www.amjcaserep.com/abstract/index/idArt/930890
Background

Profound vasodilation and low systemic vascular resistance (SVR) are characteristics of cardiac vasoplegic syndrome, which is a form of vasodilatory shock and can result in significant hypotension despite high cardiac output and appropriate fluid resuscitation [1,2]. After cardiopulmonary bypass (CPB), up to 45% of patients may exhibit vasoplegic syndrome [1-4]. The result is inadequate tissue perfusion and metabolic acidosis [2]. Significant morbidity and mortality can occur when treatment becomes refractory to vasopressors and the incidence may be increased in up to 5% of patients who do not respond to conventional vasoconstrictive therapy [3]. After cardiac surgery, reduced plasma levels of arginine vasopressin and excess nitric oxide (NO) production lead to vasodilation [2]. High-dose vasopressors are often required to maintain adequate blood pressure [1,2]. Complex interactions among plasma proteins, leukocytes, platelets, endothelial cells, and cytokines factor into this syndrome [2]. However, the primary clinical manifestation involves systemic hypotension [2]. Preoperative risk factors associated with higher incidence of postoperative vasoplegic syndrome include preoperative use of beta-blockers and angiotensin-converting enzyme (ACE) inhibitors, which increase the relative risk of vasoplegic syndrome by 1.37 and 1.31, respectively [5,6]. Use of vasopressors before or during CPB and longer duration of the procedure also confer increased risk of developing vasoplegic syndrome [1]. For each additional 30-minute interval on CPB, the risk of vasoplegic syndrome increases by 38% [5,6].

Resistance to vasopressors is thought to occur through 3 mechanisms: activation of adenosine triphosphate-sensitive potassium channels in the plasma membrane of vascular smooth muscle, vasopressin deficiency, and activation of inducible NO synthase [7]. Refractory vasoplegic syndrome that occurs after CPB has been successfully treated with inhibitors of the NO system, such as methylene blue [3].

Hydroxocobalamin also mitigates the effects of NO [8]. It has efficacy in combination with methylene blue when vasoplegic syndrome is refractory to methylene blue, and in its own right, as shown in several case reports and series [3,4,9-12]. We present 2 cases that add to the existing evidence about the efficacy of hydroxocobalamin as an appropriate option for vasoplegic syndrome refractory to other vasopressors. Quickly initiating appropriate treatment of refractory vasoplegic syndrome after CPB is vital because the mortality rate may be as high as 25% when the condition lasts for more than 36 to 48 h [5].

Case Reports

Patient 1

A 71-year-old man with a history remarkable for coronary artery disease (CAD) with recent ST-elevation myocardial infarction, systolic heart failure, hyperlipidemia, and chronic cellulitis in both lower extremities presented for a coronary artery bypass graft (CABG) and apical aneurysm repair.

Before surgery, the patient’s vital signs were as follows: heart rate, 78 beats/min; blood pressure, 119/65 mmHg; respiratory rate, 20 breaths/min; and oxygen saturation of 94% on room air. Laboratory studies were remarkable for decreased renal function with a serum creatinine of 1.3 mg/dL (reference: 0.7-1.2 mg/dL). The patient was on a beta-blocker, which increased the relative risk of vasoplegic syndrome.

Transthoracic echocardiography (TTE) revealed a decreased left ventricular ejection fraction (LVEF) of 30% with an aneurysmal apex, normal right ventricular size and function, mild mitral and tricuspid valve regurgitation, and mild aortic stenosis. Left heart catheterization showed severe triple vessel disease with 100% mid-left anterior descending coronary artery (LAD), 60% proximal obtuse marginal-3, and 70% mid-right coronary artery (RCA) stenosis.

A transesophageal echocardiogram (TEE) was performed immediately after intubation for intraoperative monitoring and diagnosis. The patient required boluses of vasopressor before initiation of CPB for a total of 20 mg of epinephrine, 2 units of vasopressin, and 200 mcg of phenylephrine to maintain a mean arterial blood pressure (MAP) of 65 mmHg. The retrograde autologous prime was the main reason for blood pressure supplementation. During CPB, hemodynamic stability was maintained without vasopressor support. After cross-clamp removal, epinephrine (3 µg/min) was initiated to assist with separation. Vasopressin (0.04 units/min) was started soon after echocardiography. That exam showed that the LVEF had increased to 45% and the right ventricular function remained normal. Cardiac output (CO) as calculated intraoperatively with echocardiography was 6.0 L/min.

Multiple boluses of vasopressin were administered and the infusion was increased to 0.06 units/min in an attempt to maintain a MAP above 65 mmHg. Calcium chloride was also given 30 min after cross-clamp removal and fluid resuscitation was guided by echocardiography. A Swan-Ganz catheter was placed before leaving the operating room. The entire procedure took 71 min and the patient still required multiple boluses of vasopressin at the completion of surgery. On arrival in the Intensive Care Unit (ICU), norepinephrine (6 µg/min) was started, vasopressin was increased to 0.08 units/min, and the epinephrine...
infusion rate remained unchanged. Stress-dose steroids were given for refractory hypotension related to potential corticosteroid insufficiency. The CO via the Swan-Ganz catheter was 5.6 L/min with a cardiac index (CI) of 2.9 L/min/m²; SVR was 859 dynes/sec/cm⁵. Because the patient’s vasopressor requirements were increasing and his CI was adequate, hydroxocobalamin was administered for refractory vasoplegic shock. A total dose of 5 g was administered i.v. over 10 min. The SVR immediately increased to 1163 dynes/sec/cm⁵ and peaked at 1638 dynes/sec/cm⁵ several hours later with weaning of vasopressor support. Within 24 h, the norepinephrine infusion was discontinued and the epinephrine and vasopressor infusions were decreased; they were ultimately shut off later in the day, after hydroxocobalamin administration. The Swan-Ganz catheter was removed at this point.

The patient was extubated the day after CPB, and over the next 3 days, low-dose dobutamine was used to augment CO to the kidneys and vasopressin was used at low doses as needed to maintain the MAP above 65 mmHg. On the fifth day after surgery, the patient was discharged from the ICU. On the eighth day after surgery, the patient was on room air, able to ambulate 400 feet, and discharged home.

**Patient 2**

A 71-year-old man with a history remarkable for CAD with recent non-ST-elevation myocardial infarction, mitral stenosis, peripheral vascular disease, and type 2 diabetes mellitus presented for a CABG and mitral valve replacement.

Before surgery, the patient’s vital signs were as follows: heart rate, 71 beats/min; blood pressure, 141/81 mmHg; respiratory rate, 16 breaths/min; and oxygen saturation of 96% on room air. Laboratory studies were remarkable for decreased renal function with a creatinine of 1.3 mg/dL (reference: 0.7-1.2 mg/dL) and glucose of 233 mg/dL (reference: 74-99 mg/dL). The patient was on a beta-blocker and an ACE inhibitor before surgery, which increased the relative risk of vasoplegic syndrome. TTE revealed a normal LVEF of 60%, normal right ventricular size and function, mild-to-moderate mitral regurgitation with multiple jets, moderate mitral stenosis, significant calcification, mild tricuspid valve regurgitation, and mild aortic stenosis. Left heart catheterization demonstrated severe multi-vessel disease with 80% proximal to mid-LAD stenosis, mid-95% stenosis in the circumflex, and 90% diffuse RCA stenosis of the more anterior branch.

A TEE was placed immediately after intubation for intraoperative monitoring and diagnosis. The patient required boluses of vasopressor prior to initiation of CPB, for a total of 450 µg of phenylephrine, to maintain a MAP of 65 mmHg. The retrograde autologous prime was the main reason for blood pressure supplementation. Soon after administration of CPB, cross-clamping, and cardioplegia administration, maintenance of blood pressure became difficult. A norepinephrine infusion was started at 4.4 µg/min and multiple boluses of vasopressin were given (8 units total), but minimal MAP elevation occurred. It was determined that there was a problem with the CPB pump, because the revolutions per minute were elevated and there was no pressure in the aorta. At this point, stress-dose steroids were given for refractory hypotension related to potential corticosteroid insufficiency. Propofol (20 mg) was initiated to achieve a bispectral index monitor of 0, epinephrine boluses were given (1 mg followed by 100 µg), and the patient’s head was packed with ice while he was in the Trendelenburg position. One surgeon unclamped the aorta and initiated manual compressions of the patient’s heart to facilitate blood flow to the brain. Another surgeon cannulated his femoral artery. Once femoral arterial cannulation was achieved, arterial flow was obtained and the patient’s hemodynamics stabilized. The norepinephrine infusion was continued throughout the bypass period to maintain a MAP around 70 mmHg. A cross-clamp was reapplied and the surgery proceeded. The patient was kept cool for the majority of the bypass time and he was rewarmed prior to weaning. After cross-clamp removal, epinephrine (7.6 µg/min) was started to assist with separation and the norepinephrine was continued at 6.5 µg/min. An echocardiogram showed that the LVEF had decreased to 45%, right ventricular function was mildly impaired, and the prosthetic mitral valve (27-mm Medtronic Mosaic) was well-seated and functioning normally with no signs of regurgitation. The patient was given calcium chloride 30 min after cross-clamp removal and fluid resuscitation was guided by echocardiography.

The patient received 5 units of packed red blood cells, 2 units of fresh frozen plasma, 6 units of random donor platelets, and 15 units of cryoprecipitate secondary to a bloody surgical field and anemia. No bolus of vasopressors was required to maintain MAP prior to leaving the operating room. The length of CPB was 209 min. Norepinephrine was started at 6 µg/min and epinephrine at 7 µg/min.

Overnight in the ICU, the patient’s norepinephrine requirements increased from 6 µg/min to 14 µg/min with vasopressin started at 0.02 units/min. The epinephrine infusion rate remained the same. A noninvasive CO monitor placed on the patient showed an SVR of 620 dynes/sec/cm⁵ and a CI of 2.7 L/min/m². Because his vasopressor requirements were increasing and his CI was adequate, hydroxocobalamin was administered for refractory vasoplegic shock (total dose 5 g i.v. over 10 min). SVR increased to 854 dynes/sec/cm⁵ and the norepinephrine infusion was discontinued within 1 h. The patient was weaned off epinephrine over the next 48 h and off vasopressin over the next 72 h.
After surgery, the patient developed an acute kidney injury that required dialysis. He also developed new-onset atrial fibrillation that required amiodarone infusion on the day after surgery. He eventually developed junctional bradycardia that required dual-chamber pacemaker placement. The patient ultimately required a tracheostomy and placement of a percutaneous endoscopic gastrostomy (PEG) tube to facilitate complete recovery because he was still demonstrating signs of encephalopathy.

On the ninth day after surgery, the patient was transferred to a long-term acute care facility. Over the course of the following week, his tracheostomy was capped and he was eating on his own without use of the PEG tube. He continued to require dialysis.

**Discussion**

Patient 2 was on an ACE inhibitor and both he and Patient 1 were on beta-blockers before surgery, with vasopressors initiated prior to CPB, which increased the relative risk of vasoplegic syndrome [5]. Prolonged CPB time is considered to be >180 min and confers increased risk of post-CPB vasoplegic syndrome [5,13]. Patient 2 had a total CPB time of 209 min. A retrospective study of 2823 cardiac surgery cases by Levin et al demonstrated a mean CPB time of 164.4 min [6]. Because each additional 30-min interval on CPB increases the risk of vasoplegic syndrome by 38%, the risk was nearly 76% higher in Patient 2 than in the average patient in the study by Levin et al [5,6]. Thus, both our patients had increased risks of developing post-CPB vasoplegic syndrome, particularly Patient 2.

There is no universal definition for vasoplegic syndrome. Accepted definitions include SVR <800 dynes/s/cm², MAP <60 to 65 mmHg, CI >2.5 to 3 L/min/m², and MAP <65 mmHg within 24 h after CPB. Because of product availability and evidence from the literature supporting its safety and efficacy, hydroxocobalamin was chosen for refractory vasoplegic syndrome for our patients because of product availability and evidence from the literature supporting its safety and efficacy. A recently published retrospective study comparing the drugs for treatment of vasoplegic syndrome that occurs after CPB [3].

Methylene blue is a water-soluble dye that inhibits NO synthase and guanylate cyclase to reverse vasodilation caused by excessive NO signaling [15]. Methylene blue has been the treatment of choice, given its effectiveness as both a preventive and rescue therapy [3,16-18]. However, drug shortages and drug-drug interactions through inhibition of monoamine oxidase to perpetuate serotonin syndrome may lead to the need for alternative therapy for vasoplegic syndrome that occurs after CPB [3].

Many case reports detailing hydroxocobalamin efficacy include concomitant use of methylene blue [3,7,8]. Methylene blue was given in addition to hydroxocobalamin in 54% of patients in the Armour et al case series and 45% in the Shah et al case series and there may be an advantage to the combination versus methylene blue alone [11,12,19].

Like the report by Roderique et al, our study demonstrates efficacy without concomitant methylene blue [4]. Both patients in the present case study received a single, 5-g dose of hydroxocobalamin given i.v., with very rapid hemodynamic responses, as MAP and SVR increased while the vasopressor requirement was reduced within hours. Hydroxocobalamin was chosen for refractory vasoplegic syndrome for our patients because of product availability and evidence from the literature supporting its safety and efficacy. A recently published retrospective study comparing hydroxocobalamin and methylene blue did not demonstrate differences between the ability to increase MAP, increase SVR, or lead to changes between groups in norepinephrine equivalents [20]. This equivalency suggests that either agent is appropriate for refractory vasoplegic syndrome that occurs after CPB. Unfortunately, a clinical trial comparing the drugs for treatment of vasoplegic syndrome was discontinued in 2020 due to lack of funding [21].

Both of our patients exhibited chromaturia and erythema, but otherwise, we report no significant adverse events (AEs) due to the administration of hydroxocobalamin in either case. A study of 136 healthy volunteers showed that chromaturia and erythema were the most common AEs in those that received 5- or 10-g doses of hydroxocobalamin [22]. Oxalate crystals were found in the urine of 61% of those that received 5-g doses and 56% of those that received 10-g doses [22]. There were no reports of photosensitivity, angioedema, or anaphylaxis [22].

It is not entirely known how i.v. hydroxocobalamin mitigates catecholamine-resistant vasodilatory shock, but it scavenges, binds, and prevents the formation of NO and hydrogen sulfide, which may also potentiate vasodilation and hypotension [9]. NO oxidizes the cobalt from hydroxocobalamin [9]. This complex of cobalt and NO may transfer NO to hemoglobin to reduce the NO level. Less NO and hydrogen sulfide may stabilize capillary membranes and restore vascular tone [7]. Hydroxocobalamin also likely exerts its vasopressor effects through inhibition of inducible NO synthase and guanylate synthase [4,14].
Conclusions

The 2 cases in our report represent further evidence that hydroxocobalamin can be substituted for methylene blue to treat vasoplegic syndrome occurring after CPB that is refractory to other vasopressors and in situations in which methylene blue is contraindicated or unavailable, to reduce vasopressor demand and restore appropriate hemodynamic stability. Development of vasoplegic syndrome after CPB leads to death or a hospital stay of >10 days in 57.4% of patients [5]. Both of our patients avoided such outcomes, with vasoplegic syndrome resolving shortly after hydroxocobalamin utilization.

References:


Conflict of Interest

None.