A Case of Spurious Hypoxemia in an ICU Patient With Leukemic Blast Crisis

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Abstract

Arterial blood gas analysis is one of the most common tests performed on patients in the intensive care unit (ICU) to diagnose hypoxemia, a condition with low oxygen tension in the blood (PaO\textsubscript{2}). Hypoxemia can be true or spurious/pseudo. One of the causes of spurious hypoxemia is the presence of metabolically active cells like white blood cells (WBCs) and platelets. We present a case of blast crisis due to acute myelogenous leukemia (AML) presenting with spurious hypoxemia which improved after leukapheresis in our ICU.

Keywords: Hypoxemia; Leukemia; Spurious

Introduction

Arterial blood gas analysis is one of the most common tests performed on patients in the intensive care unit (ICU) to diagnose hypoxemia, a condition with low oxygen tension in the blood (PaO\textsubscript{2}) [1, 2]. However, PaO\textsubscript{2} in the arterial blood sample can be affected by many reasons, one being the presence of metabolically active cells like white blood cells (WBCs) and platelets leading to spurious or pseudo hypoxemia [3]. We present a case of blast crisis due to acute myelogenous leukemia ((AML) presenting with spurious hypoxemia which improved after leukapheresis.

Case Report

An 85-year-old elderly female with past medical history of diabetes, hypertension and recent diagnosis of AML was admitted to our ICU with severe metabolic acidosis and acute respiratory failure. She received recent chemotherapy outside hospital. Her chest X-ray was normal, thereby, ruling out pneumonia or congestive heart failure. Suspicion of acute pulmonary embolism was high with acute hypoxemia in a cancer patient with unimpressive chest X-ray. Transthoracic echocardiography did not show any right heart strain pattern. On arrival, she was clinically in shock, disseminated intravascular coagulation (DIC) and had WBC count of 455,000/\mu L. Her initial arterial blood gas (ABG) on admission after endotracheal intubation showed pH of 6.64, pCO\textsubscript{2} of 67 mm Hg and pO\textsubscript{2} of 29 mm Hg on 100% FiO\textsubscript{2} but pulse oximetry read SpO\textsubscript{2} of 100%. Her ICU course was complicated by septic shock, multi-system organ failure and refractory metabolic acidosis. Oncology service was consulted, and patient underwent emergent leukapheresis for blast crisis with improvement in WBC count to 140,000/\mu L. Her ABG improved to pH of 6.89, pCO\textsubscript{2} of 30 mm Hg and pO\textsubscript{2} of 351 mm Hg with SpO\textsubscript{2} of 100% on pulse oximetry. Her PO\textsubscript{2} improved dramatically after leukapheresis and removal of WBC from the blood. Family soon opted for comfort care measures, and she developed asystolic cardiac arrest and died.

Discussion

Hypoxemia is a serious condition commonly seen in the ICU patients. Underlying physiologic derangements include ventilation/perfusion mismatch, hypoventilation, shunt and abnormality of diffusion capacity [1]. ABG is a common test employed to diagnose hypoxemia. Hypoxemia can be true or spurious. Common clinical conditions leading to hypoxemia in ICU include pneumonia, pulmonary embolism, acute respiratory distress syndrome and congestive heart failure. Pseudo hypoxemia or spurious hypoxemia is an entity where laboratory abnormalities of PaO\textsubscript{2} are not a true reflection of patient’s bloodstream oxygen tension. Causes of inaccurate measurements of PaO\textsubscript{2} include improper sampling, delay in transport with failure to cool the specimen, presence of bubbles in the sample, elevated cell count, blood gas analyzer malfunction or miscalibration or methemoglobinemia [1, 2]. Spurious hypoxemia has been reported in patients with hyperleukocytosis or thrombocytosis in the setting of hematologic conditions.
logical malignancies [2, 4, 5]. It was first described in 1979 by Fox et al and was coined leukocyte larceny [3]. This is due to the presence of high number of metabolically active WBCs with elevated consumption of dissolved oxygen in the arterial blood samples [6]. The rate of oxygen consumption by WBCs is typically not important clinically in majority of patients with normal blood counts. Spuriously low PaO₂ measurements are common when WBC count exceeds 50,000/µL or with severe thrombocytosis. This entity is not very well described in the critical care literature with not many cases reported in ICU setting [1, 2].

Patients with hyperleukocytosis in the setting of acute myelogenous or lymphocytic leukemia or blast crisis can develop true or spurious hypoxemia. Causes of true hypoxemia in these patients include sepsis, pulmonary embolism, pulmonary leukostasis, leukemic infiltration, hemorrhage and drug-related toxicities. Inconsistencies between SpO₂ with pulse oximetry and PaO₂ in the arterial blood can help to differentiate spurious hypoxemia from these causes. In leukocyte larceny, SpO₂ is normal and PaO₂ is low. Cooling of blood sample on ice and rapid analysis can help reduce the discrepancy.

Pulse oximetry is the most accurate method to assess oxygenation in patients with hyperleukocytosis [7, 8]. It measures hemoglobin saturation directly and is unaffected by the plasma oxygen tension. Understanding the concept of spurious hypoxemia in patients with hematologic malignancies will help to prevent unnecessary tests like CT scans and ventilation/perfusion scans to rule out pulmonary embolism, minimizing ICU admission of these patients and saving hospital resources and costs.

Our patient did not undergo these expensive tests for the workup of pulmonary embolism as PaO₂ improved after leukapheresis.

**Conclusion**

ICU physicians should be vigilant about the concept of pseudo/spurious hypoxemia and leukocyte larceny in patients with leukemias and blast crisis. Causes of true hypoxemia still need to be ruled out in the correct clinical setting. Pulse oximetry is the most reliable and easy method to clinically assess oxygenation in patients with leukocyte larceny.

**References**