Preoperative anemia in major elective surgery

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An estimated 23%–45% of patients undergoing major surgery have anemia, with the most common causes being iron deficiency anemia and anemia of inflammation or chronic disease.^{1,2}

Preoperative anemia leads to adverse outcomes

Regardless of its severity, preoperative anemia is an independent risk factor for postoperative death, major morbidity, increased length of hospital stay and transfusion.^{1,3} In patients undergoing cardiac surgery, a 10 g/L decrease in preoperative hemoglobin levels increased mortality odds by 16% (95% confidence interval 10%–22%).²

3 A preoperative hemoglobin of 130 g/L or higher should be targeted for both sexes

Females have lower circulating blood volumes and greater proportional operative blood loss than males.⁴ Females with a hemoglobin of 120 g/L were shown to be twice as likely as males with a hemoglobin of 130 g/L to receive postoperative blood transfusions.⁴ When treating preoperative anemia, targeting the same hemoglobin level in both sexes minimizes the risk of unfavourable outcomes and transfusions.⁴

Patients undergoing major elective surgery, with expected blood loss of more than 500 mL, should be screened for anemia 6-8 weeks before their operation

Clinicians should order a complete blood count and ferritin levels, as iron deficiency anemia (ferritin < 30 ng/mL) is the most common cause. ^{1,4} When underlying inflammation is present, ferritin is less sensitive, and iron deficiency anemia can be diagnosed with a ferritin of 30–100 ng/mL and a transferrin saturation of less than 20%. ^{1,4} Patients with iron deficiency anemia should be investigated for an underlying cause (e.g., gastrointestinal blood loss, menorrhagia, malabsorption).

5 Preoperative iron deficiency anemia should be treated with iron supplementation

Patients with iron deficiency anemia at least 8 weeks from surgery should be treated with oral supplementation at equivalent doses of 40–60 mg elemental iron daily or 80–100 mg every other day.^{1,4} If patients are within 8 weeks of surgery, or if they are unable to tolerate oral supplementation, they should receive intravenous iron.¹ For patients with refractory or other forms of anemia, erythropoiesis-stimulating agents can be considered along with a specialist referral.^{1,5}

References

- Greenberg JA, Zwiep TM, Sadek J, et al. Clinical practice guideline: evidence, recommendations and algorithm for the preoperative optimization of anemia, hyperglycemia and smoking. Can J Surg 2021;64:E491-509.
- Klein AA, Collier TJ, Brar MS, et al.; Association of Cardiothoracic Anaesthetists (ACTA). The incidence and importance of anaemia in patients undergoing cardiac surgery in the UK: the first Association of Cardiothoracic Anaesthetists national audit. *Anaesthesia* 2016;71:627-35.
- Mueller MM, Van Remoortel H, Meybohm P, et al.; ICC PBM Frankfurt 2018 Group. Patient blood management: recommendations from the 2018 Frankfurt Consensus Conference. JAMA 2019;321:983-97.
- Muñoz M, Acheson AG, Auerbach M, et al. International consensus statement on the peri-operative management of anaemia and iron deficiency. Anaesthesia 2017;72:233-47.
- Kei T, Mistry N, Curley G, et al. Efficacy and safety of erythropoietin and iron therapy to reduce red blood cell transfusion in surgical patients: a systematic review and meta-analysis. Can J Anaesth 2019:66:716-31.

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