Myocardial injury after noncardiac surgery

Anjali Kulkarni MD MSc, Maura Marcucci MD MSc, Julie M.V. Nguyen MD MSc

Cite as: CMAJ 2024 May 21;196:E661. doi: 10.1503/cmaj.231355

1 Myocardial injury after noncardiac surgery (MINS) is frequent and often asymptomatic

Defined as troponin elevation exceeding the 99th percentile of reference values within 30 days of noncardiac surgery due to ischemia, the estimated incidence is 12%–24% and is higher after nonelective procedures.¹ More than 80% of patients present without either symptoms related to ischemia or electrocardiographic changes, and the condition is missed without measurement of troponin.¹

2 Postoperative morbidity and mortality are increased regardless of symptoms

Myocardial injury after noncardiac surgery is associated with increased 30-day and 1-year mortality, compared with those without MINS.² Within 16 months, 1 in 6 patients with MINS will experience a major vascular complication, including vascular death, myocardial infarction, stroke, peripheral arterial thrombosis, or symptomatic venous thromboembolism.³

3 High-risk patients undergoing noncardiac surgery should have routine troponin measurement

The Canadian Cardiovascular Society guideline recommends immediate postoperative electrocardiography in the recovery room, daily troponin measurement for 48 to 72 hours, and multidisciplinary management for high-risk patients.⁴ For patients undergoing emergency or urgent or semiurgent surgery, those aged 65 years or older, or aged 18–64 years with cardiovascular disease, are deemed high risk. For patients undergoing elective surgery, those aged 65 years or older, or 45–64 years with cardiovascular disease, or with a Revised Cardiac Risk Index score of 1 or more, are deemed high risk.⁴ In the elective setting, physicians should use preoperative brain natriuretic peptide (BNP) level measurement, when available, to refine risk assessment; postoperative troponin is recommended when the N-terminal prohormone of BNP (NT-proBNP) is 300 ng/L or higher, or BNP 92 ng/L or higher.

Early identification allows for prompt management Patients with high-risk features such as chest pain or electrocardiography changes may require inpatient cardiology assessment and urgent cardiac catheterization.⁵

5 All patients with MINS, regardless of symptoms, should receive secondary prevention

This includes acetylsalicylic acid, statin, and lifestyle modification. In a large randomized controlled trial, long-term treatment with dabigatran decreased major arterial and venous thrombotic complications without increasing bleeding and may be considered on a case-by-case basis.³

References

- Smilowitz NR, Redel-Traub G, Hausvater A, et al. Myocardial injury after noncardiac surgery: a systematic review and meta-analysis. *Cardiol Rev* 2019;27:267-73.
- Writing Committee for the VISION Study Investigators; Devereaux PJ, Biccard BM, Sigamani A, et al. Association of postoperative high-sensitivity troponin levels with myocardial injury and 30-day mortality among patients undergoing noncardiac surgery. JAMA 2017;317:1642-51.
- Devereaux PJ, Duceppe E, Guyatt G, et al.; MANAGE Investigators. Dabigatran in patients with myocardial injury after non-cardiac surgery (MANAGE): an international, randomised, placebo-controlled trial. *Lancet* 2018;391:2325-34.
- 4. Duceppe E, Parlow J, MacDonald P, et al. Canadian Cardiovascular Society guidelines on perioperative cardiac risk assessment and management for patients who undergo noncardiac surgery. *Can J Cardiol* 2017;33:17-32.
- Borges F, Ofori S, Marcucci M. Myocardial injury after noncardiac surgery and perioperative atrial fibrillation: from evidence to clinical practice. *Can J Gen Intern Med* 2021;16:18-26.

Competing interests: Julie Nguyen reports receiving research grants from the Population Health Research Institute, McMaster Institute for Research on Aging, Juravinski Hospital and Cancer Centre Foundation, and the Hamilton Health Sciences Foundation. Maura Marcucci reports receiving, as principal investigator, research grants from the Canadian Institutes of Health Research, the Hamilton Academic Health Sciences Organization, the Population Health Research Institute, and the McMaster Institute for Research on Aging with the Michael DeGroote Institute for Pain Research and Care (MIRA-IPRC Catalyst Grant), all outside the submitted work. No other competing interests were declared. This article has been peer reviewed.

Affiliations: Division of Gynecologic Oncology (Kulkarni), Departments of Medicine, and Health Research Methods, Evidence, and Impact (Marcucci), and Division of Gynecologic Oncology (Nguyen), Department of Obstetrics and Gynecology, McMaster University, Hamilton, Ont.; Clinical Epidemiology and Research Centre (Marcucci), Department of Biomedical Sciences, Humanitas University, Milan, Italy. **Content licence:** This is an Open Access article distributed in accordance with the terms of the Creative Commons Attribution (CC BY-NC-ND 4.0) licence, which permits use, distribution and reproduction in any medium, provided that the original publication is properly cited, the use is noncommercial (i.e., research or educational use), and no modifications or adaptations are made. See: https://creativecommons.org/ licenses/by-nc-nd/4.0/

Correspondence to: Julie Nguyen, nguyenjmv@hhsc.ca