

# Local Guideline

Document number: JHH\_0449

## Perioperative management of antiplatelet agents

|  |   |
|--|---|
| <b>Sites where Local Guideline applies</b> | John Hunter Hospital  |
| <b>This Local Guideline applies to:</b>    |   |
| 1. Adults                                  | Yes   |
| 2. Children up to 16 years                 | No  |
| 3. Neonates – less than 29 days            | No  |
| <b>Target audience</b>                     | Anaesthetists, doctors, perioperative nurses, cardiologists, neurologists, surgeons   |
| <b>Description</b>                         | This guideline provides guidance on the management of antiplatelet agents in the perioperative period for patients having elective surgery. |

[Go to Guideline](#)

|  |   |
|--|---|
| <b>Keywords</b>  | Aspirin, clopidogrel, prasugrel, ticagrelor, ticlopidine, antiplatelet, antithrombotic, perioperative, surgery  |
| <b>Document registration number</b>  |   |
| <b>Replaces existing document?</b>   | No  |
| <b>Related Legislation, Australian Standard, NSW Ministry of Health Policy Directive or Guideline, National Safety and Quality Health Service Standard (NSQHSS) and/or other, HNE Health Document, Professional Guideline, Code of Practice or Ethics:</b> |   |
| • See Reference Section on page 5  |   |
| <b>Prerequisites (if required)</b>   | The patient should be booked for elective surgery at the John Hunter Hospital and be taking an antiplatelet medication.   |
| <b>Local Guideline note</b>  | This document reflects what is currently regarded as safe and appropriate practice. This guideline does not replace the need for the application of clinical judgment in respect to each individual patient. If staff believe that the guideline should not apply in a particular clinical situation they must seek advice from a Perioperative Anaesthetist, and/or the patient's antiplatelet prescriber and surgeon, and document the variance in the patient's health record.<br>If this document needs to be utilised outside of the John Hunter Hospital please liaise with the local cardiology, neurology, and surgical services to ensure the appropriateness of the information contained within the Guideline and Procedure. |
| <b>Date Initial authorisation:</b>   | 27 <sup>th</sup> April 2022   |
| <b>Authorised by:</b>  | Yes   |
| <b>This document contains advice on therapeutics:</b>  | Approval gained from John Hunter Quality Use of Medicines Committee on 27 <sup>th</sup> April 2022  |
| <b>Contact person:</b>   | Dr Paul Healey  |
| <b>Contact details:</b>  | <a href="mailto:Paul.Healey@health.nsw.gov.au">Paul.Healey@health.nsw.gov.au</a> (02) 49223018  |
| <b>Date Reviewed:</b>  | July 2022   |
| <b>Review Due Date:</b>  | April 2025  |
| <b>Version:</b>  | 1.0 27 <sup>th</sup> July 2022  |

Note: Over time links in this document may cease working. Where this occurs please source the document in the PPG Directory at: <http://ppg.hne.health.nsw.gov.au/>

## PURPOSE AND RISKS

Antiplatelet therapies are used for a variety of indications in the community. The perioperative period poses a risk to the patient of adverse vascular events due to medication changes and the proinflammatory and prothrombotic states that exist at this time. Management in the perioperative period requires consideration of patient, surgical and anaesthetic factors to reduce the risks of:

- Cardiovascular events including myocardial infarction, stroke and acute limb ischaemia.
- Surgical site bleeding

The balance between these competing interests can be complex and consultation may be required with other specialties to formulate a perioperative plan for higher risk patients.<sup>1</sup>

**Risk Category:** Clinical Care & Patient Safety

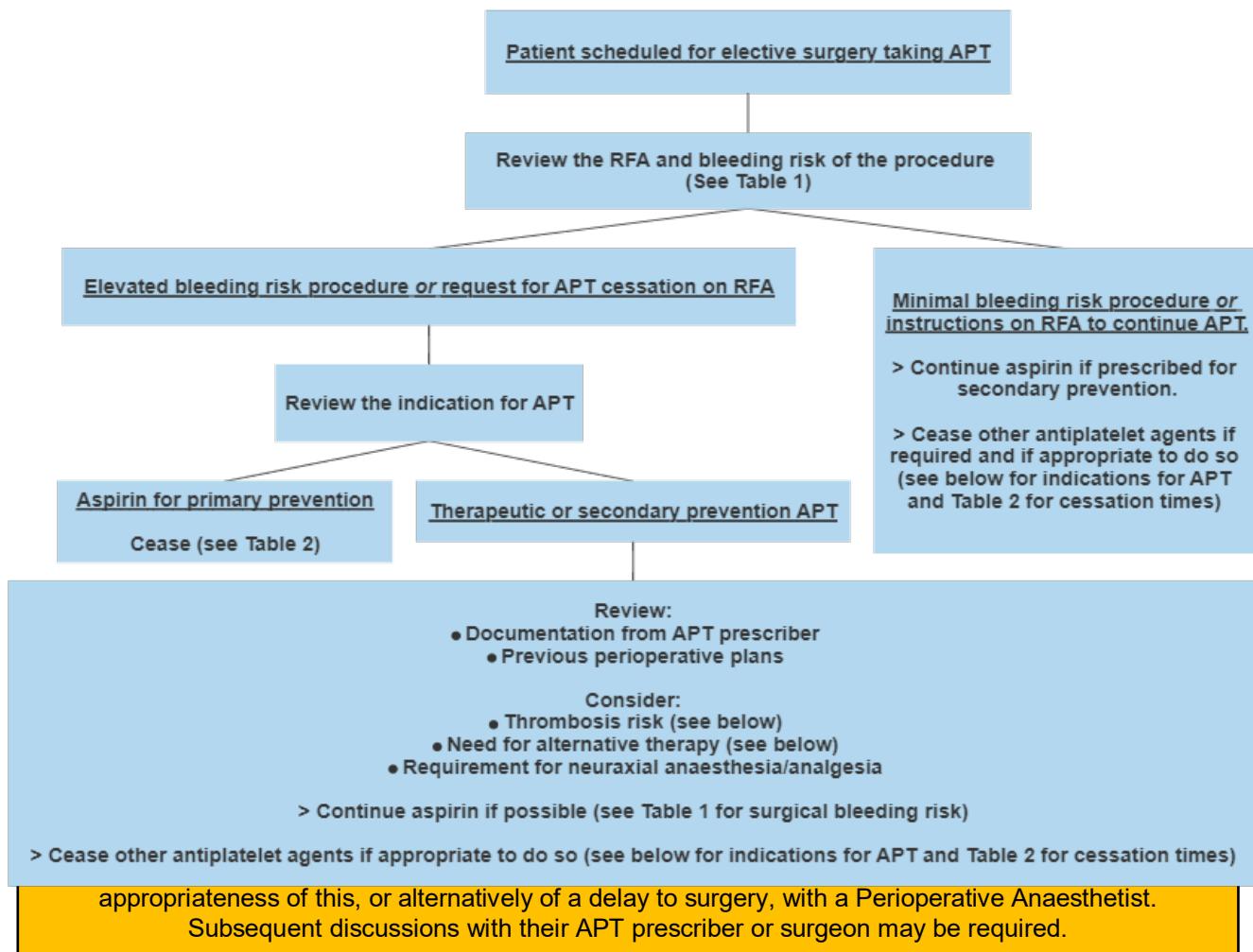
## GLOSSARY

| Acronym or Term | Definition                                |
|-----------------|---|
| ACS             | Acute coronary syndrome                   |
| APT             | Antiplatelet therapy                      |
| DAPT            | Dual antiplatelet therapy                 |
| ENT             | Ear nose and throat                       |
| ESWL            | External shock wave lithotripsy           |
| LAD             | Left anterior descending                  |
| PCI             | Percutaneous coronary intervention        |
| PCNL            | Percutaneous nephrolithotripsy            |
| RFA             | Request for admission                     |
| TURBT           | Transurethral resection of bladder tumour |
| TURP            | Transurethral resection of prostate       |

## GUIDELINE

This Guideline does not replace the need for the application of clinical judgment in respect to each individual patient.

## Antiplatelet Management in the Perioperative Period



**Table 1: Assessment of surgical bleeding risk**

|  | <b>Surgery Type</b>   | <b>APT Management</b>  |
|--|---|--|
| <b>High surgical bleeding risk</b>     | <ul style="list-style-type: none"> <li>• Intracranial surgery</li> <li>• Urology –TURP</li> <li>• Intraocular surgery</li> <li>• Major spinal surgery</li> </ul>  | Cease all APT agents   |
| <b>Moderate surgical bleeding risk</b> | <ul style="list-style-type: none"> <li>• Urology – Cystoscopy/ureteroscopy with diathermy, resection of tumour, stone therapy, prostatectomy, ESWL, PCNL, scrotal surgery, TURBT (large tumours), major open surgery</li> <li>• Major ENT including adenotonsillectomy, septorhinoplasty, middle ear surgery.</li> <li>• Major Head and Neck cancer surgery</li> <li>• Spinal surgery</li> <li>• Thyroidectomy and similar</li> <li>• Facio-maxillary surgery</li> <li>• Primary and revision arthroplasty</li> </ul> | <p>Consider ceasing aspirin depending on indication for use.</p> <p>Cease clopidogrel (+/- replace with aspirin, if indicated)</p> |
| <b>Minimal or low bleeding</b>         | <ul style="list-style-type: none"> <li>• All other surgeries <i>unless requested by</i></li> </ul>  | Continue aspirin.  |

|             |                     |   |
|-------------|---------------------|---|
| <b>risk</b> | <i>the surgeon.</i> | Cease clopidogrel (+/- replace with aspirin if indicated) |
|-------------|---------------------|---|

**Table 2: Duration of cessation of APT<sup>1</sup>**

| Antiplatelet agent | When to cease antiplatelet therapy (if required) |
|--------------------|--|
| aspirin            | At least 5 days prior                            |
| clopidogrel        | At least 7 days prior                            |
| prasugrel          | At least 7 days prior                            |
| ticagrelor         | At least 5 days prior                            |
| ticlopidine        | At least 14 days prior                           |

**Assessment of thrombotic risk and consequences**

Factors which contribute to the risk associated with stopping antiplatelet therapy:

- Duration of time since coronary or neurovascular event.
- Location (left main or proximal LAD stent), length, number, branching, difficulty of placement, clinical context of insertion (e.g. myocardial infarction v. symptomatic coronary artery disease), calibre and possibly type of stent.<sup>2</sup>
- Multi-vessel or multi-lesion coronary disease or stents<sup>2,3</sup>
- Multiple anatomical sites of vascular disease<sup>3</sup>
- A history of stent thrombosis
- Patient characteristics such as prothrombotic states, smoking, chronic kidney disease, diabetes mellitus, advanced age.<sup>2</sup>

*Patients with high thrombotic risk may require longer periods of DAPT after stenting, combinations of antiplatelet and anticoagulant medications, and longer delays to non-urgent surgery after a vascular event. If in doubt, discuss with a Perioperative Anaesthetist. A discussion with the patient's antiplatelet prescriber and their surgeon may be required for complex cases.<sup>2,3</sup>*

**For patients with coronary artery disease:**

- Where possible, aspirin should be continued perioperatively in patients with coronary stents.<sup>4</sup>
- *Urgent surgery may be undertaken 1-3 months after coronary stenting with bare metal stents or drug eluting stents if the risk of surgical delay outweighs the risk of stent thrombosis. This requires discussion with a Perioperative Anaesthetist prior to discussion with the patient's cardiologist and surgeon.<sup>5,6</sup>*
- Where possible, non-urgent surgery should be delayed until 6 months after PCI.
- With PCI after ACS, perioperative MI rates remain elevated for 12 months, thus DAPT should continue and non-urgent elective surgery should be delayed until this time.<sup>2,7,8</sup>

**For patients with cerebrovascular disease**

- Non-urgent surgery should be delayed until 9 months post CVA when the risk of perioperative stroke reaches its nadir (see the local guideline for Elective Surgery after Stroke)
- There is limited evidence regarding best practice regarding antiplatelet therapies in this population.
- In patients with existing cerebrovascular stents, aspirin is usually continued perioperatively where possible.<sup>4,7</sup> Dual antiplatelet therapy, or an antiplatelet agent and an anticoagulant, is often used for 3-6 months after intracranial stenting. Temporary cessation (of agents other than aspirin) after 6 months, when required for surgery, is usually appropriate.

**Patients with peripheral vascular disease**

- DAPT may be continued in the setting of vascular surgery.<sup>7</sup> Check RFA for surgical instructions.

- For patients with severe PVD or peripheral vascular stents having non-vascular surgery, check for surgical instructions and continue APT where possible.
- For carotid endarterectomy (CEA), DAPT is continued perioperatively.<sup>7</sup>

### **Alternative therapies**

- Note that heparin bridging does not reduce the risk of coronary stent thrombosis.<sup>7</sup>
- Tirofiban and eptifibatide are infrequently used for patients at extremely high thrombotic risk.<sup>3</sup>

### **Recommencement of APT**

- This should occur as soon as thought safe from a surgical bleeding risk and neuraxial anaesthesia/analgesia perspective. See the American Society of Regional Anesthesia (ASRA) guideline for guidance on specific agents.<sup>9</sup>
- A loading dose may be requested by the prescriber for patients at high thrombotic risk.<sup>7</sup>

## **IMPLEMENTATION, MONITORING COMPLIANCE AND AUDIT**

This guideline will be communicated to the Perioperative Department via their continuing medical education meetings.

## **REFERENCES**

1. Clinical Excellence Commission, 2018, [Guidelines on Perioperative Management of Anticoagulant and Antiplatelet Agents](#) Sydney: Clinical Excellence Commission
2. Mehta, S. 2018 Canadian Cardiovascular Society/Canadian Association of Interventional Cardiology Focused Update of the Guidelines for the Use of Antiplatelet Therapy. *Can J Cardiology* Volume 34, Issue 3, March 2018, Pages 214-233  
<https://doi.org/10.1016/j.cjca.2017.12.012>
3. The Task Force for dual antiplatelet therapy in coronary artery disease of the European Society of Cardiology (ESC) and of the European Association for Cardio-Thoracic Surgery (EACTS). 2017 ESC focused update on dual antiplatelet therapy in coronary artery disease developed in collaboration with EACTS. *Eur. Heart J.* (2018) 39, 213–254 ESC GUIDELINES doi:10.1093/eurheartj/ehx419
4. Muluk V, et al. Perioperative medication management. In: UpToDate (2021) (Accessed on Nov 10, 2021)
5. Egholm G, et al. Risk Associated With Surgery Within 12 Months After Coronary Drug-Eluting Stent Implantation. *J. Am. Coll. Cardiol.* 68, Issue 24, 20 December 2016, Pages 2633-2636 <https://doi.org/10.1016/j.jacc.2016.09.967>
6. Holcomb C, et al. The Incremental Risk of Coronary Stents on Postoperative Adverse Events. A matched cohort study. *Ann Surg* 2016;263:924–930. DOI: 10.1097/SLA.0000000000001246
7. Devereaux P, et al. Aspirin in Patients Undergoing Noncardiac Surgery *N Engl J Med* 2014;370:1494-503. DOI: 10.1056/NEJMoa1401105
8. Livhits M, et al. Risk of Surgery Following Recent Myocardial Infarction. *Ann Surg* 2011;253:857–864
9. Horlocker T, et al. Regional anesthesia in the patient receiving antithrombotic or thrombolytic therapy. American Society of Regional Anesthesia and Pain Medicine Evidence-Based Guidelines (fourth edition). *Reg Anesth Pain Med.* 2018;43:263-309. doi: 10.1097/AAP.0000000000000763

## **FEEDBACK**

Any feedback on this document should be sent to the Contact Officer listed on the front page.