

A Retrospective Observational Study Of The Use of Gastroprotection For Patients on Dual Antiplatelet Therapy In An Acute Trust In England

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Introduction

- Dual antiplatelet therapy (DAPT), a combination of aspirin and either clopidogrel, prasugrel or ticagrelor is recommended for secondary prevention of ischaemic events in people with coronary artery disease.
- Patients taking DAPT may be at high gastrointestinal (GI) bleed risk if other factors are present and should be considered for gastroprotection e.g. a Proton Pump Inhibitor (PPI) or H2 receptor antagonist.
- Patients are at high risk of GI adverse effects with antiplatelet treatment if the following risk factors are present¹:
 - High dose of aspirin (300mg once daily)
 - Older age, especially aged over 70 years.
 - History of gastroduodenal ulcer, GI bleeding, or gastroduodenal perforation.
 - Helicobacter pylori infection.
 - Concomitant use of medicines that are known to increase the risk of GI bleeds such as anticoagulants.

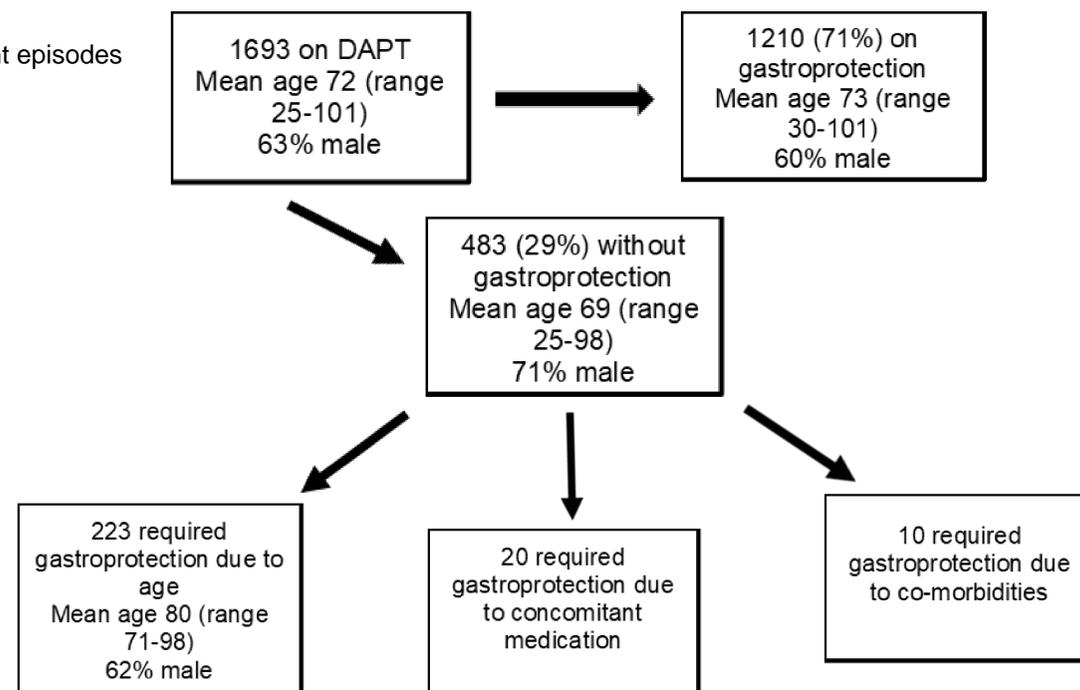
Aim: To assess whether gastroprotection was prescribed for inpatients taking DAPT in a 750-bed acute hospital in England.

Methods

- This was a retrospective analysis of existing data.
- Patient episodes involving prescription of DAPT upon discharge between April 2020 and April 2021 were extracted from the e-prescribing system.
- Electronic records of the identified patient episodes were searched for co-prescription of either PPI or H2 antagonist.
- Other risk factors such as age older than 71 and concomitant drugs that can increase bleeding risk were also checked to ascertain if patients on DAPT should have received gastroprotection.
- Ethics approval was not required for this study.

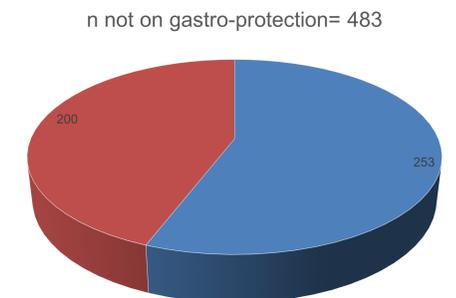
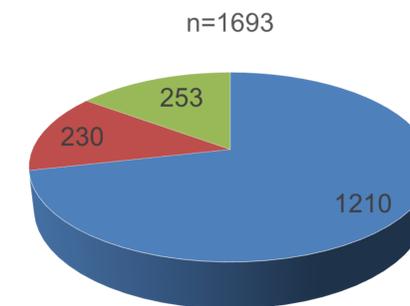
Results

Figure 1: Flow of patient episodes



Discussion

- There was a total of 1693 patient episodes on DAPT, 29% did not receive gastroprotection. 15% of these patients should be on gastroprotection due to their age (>70) and other high risk indicators such as being on concomitant medications that can increase bleeding risk and co-morbidities.
- Of the 1210 patient episodes receiving gastroprotection, a PPI was prescribed in 1171 (564 omeprazole, 10 esomeprazole and 10 rabeprazole), and an H2 antagonist was prescribed in 39 (28 famotidine and 11 ranitidine).



■ On gastro-protection ■ Not on gastro-protection (do not need it) ■ Should be on gastro-protection ■ Should be on gastro-protection ■ Do not need gastro-protection

Limitations

- It was conducted in a single centre in England; this might restrict the generalisability of our findings.
- Retrospective nature of this study may introduce bias or other uncertainties.
- We did not check the indication for DAPT nor if patients came in on these drugs as opposed to being started during their admission.
- The risk factors for gastroprotection other than age and selected concomitantly prescribed drugs were not checked.
- Significant contraindications other than electrolytes abnormalities for those not prescribed gastroprotection were not checked.

Conclusion

- Gastroprotection was potentially missing in 242 (14%) of the total DAPT patient episodes.
- Other studies have also reported that gastroprotection has been missed in the at risk group.²
- When considering newly initiated gastroprotection, various factors are considered, e.g. any possible interaction with a PPI if one of the antiplatelets is clopidogrel. Also, PPIs have adverse effects such as low sodium/ magnesium, and this may be a reason for consideration of a H2 antagonist.
- Further work is required to make our prescribers and pharmacy team aware of the importance of considering gastroprotection for this patient cohort.

References:

1. CKS, NICE., 2021. Scenario: Secondary prevention of CVD | Management | Antiplatelet treatment | CKS | NICE. [online] Cks.nice.org.uk. Available at: <https://cks.nice.org.uk/topics/antiplatelet-treatment/management/secondary-prevention-of-cvd/> [Accessed 30 May 2021].
2. Badar A, Scaife J, Yan AT, et al. Provision of gastroprotective medication and bleeding risk following acute coronary syndrome. J Invasive Cardiol 2013;25:397-401.