

# Proton-Pump Inhibitor (PPI) Use

The FDA has issued multiple warnings on the long-term use of PPIs. These include: increased risk of *C. difficile* infection<sup>1</sup>, hypomagnesemia<sup>2</sup>, and fractures of the hip, wrist, and spine<sup>3</sup>. Therefore, prudent prescribing of PPIs is warranted. The FDA recommends use of the lowest dose and shortest duration of PPI therapy appropriate for the condition being treated<sup>1-3</sup>. Patient compliance, time of administration (prior to meals), and dietary indiscretions (i.e. alcohol or irritating foods) should be assessed prior to titration of PPI doses.

Indication	Treatment	Duration
<b>Gastroesophageal reflux disease (GERD)<sup>6</sup></b>  Symptomatic relief  Acute healing of erosive or ulcerative esophagitis  Maintenance healing of erosive or ulcerative esophagitis	Omeprazole 20 mg PO once daily <b>OR</b> Pantoprazole 40mg PO once daily	Initial 8 week course for symptom relief or esophagitis  Maintenance therapy determined by response and severity of disease  For patients that require more long-term therapy, consider a trial of a lower dose, on-demand therapy, or intermittent therapy to minimize exposure
<b>Stress ulcer prophylaxis</b> Reserve PPIs for critically ill patients <i>with increased risk of bleeding</i> : <sup>4,5</sup> At least one of the following: <ul style="list-style-type: none"> <li>- Coagulopathy (platelet count &lt;50,000 mm<sup>3</sup>, INR &gt;1.5, or aPTT &gt;2x control)</li> <li>- Mechanical ventilation &gt;48 hours</li> <li>- History of GI ulceration or bleeding within past year</li> <li>- Glasgow Coma score ≤10</li> <li>- Traumatic, severe thermal or spinal cord injury</li> <li>- Hepatic failure</li> </ul> Two or more minor risk factors: <ul style="list-style-type: none"> <li>- Sepsis</li> <li>- ICU stay ≥1 week</li> <li>- Occult GI bleeding ≥6 days</li> <li>- High-dose corticosteroids (≥250 mg/day hydrocortisone equivalent)</li> </ul>	Omeprazole 20-40 mg once daily <b>OR</b> Pantoprazole 40mg IV/PO once daily	Transition to PO when possible  Continue until resolution of underlying risk factors and/or critical illness  Recommend discontinuation at discharge, unless there is another indication for use

aPTT= activated partial thromboplastin time; GERD= Gastroesophageal reflux disease; GI= Gastrointestinal; ICU= intensive care unit; INR= international normalized ratio; IV= intravenous; PO= by mouth; PPI= proton pump inhibitor

## References

1. U.S. Food and Drug Administration (FDA). FDA Drug Safety Communication: Clostridium difficile-associated diarrhea can be associated with stomach acid drugs known as proton pump inhibitors (PPIs) [internet]. Updated May 2012 [cited 11/21/12]. Available from: <http://www.fda.gov/Drugs/DrugSafety/ucm290510.htm>
2. U.S. Food and Drug Administration (FDA). FDA Drug Safety Communication: Low magnesium levels can be associated with long-term use of Proton Pump Inhibitor drugs (PPIs) [internet]. Updated February 2012 [cited 11/21/12]. Available from: <http://www.fda.gov/Drugs/DrugSafety/ucm245011.htm>.
3. U.S. Food and Drug Administration (FDA). FDA Drug Safety Communication: Possible increased risk of fractures of the hip, wrist, and spine with the use of proton pump inhibitors [internet]. Updated March 2011 [cited 11/21/12]. Available from: <http://www.fda.gov/Drugs/DrugSafety/PostmarketDrugSafetyInformationforPatientsandProviders/ucm213206.htm>
4. ASHP Therapeutic Guidelines on Stress Ulcer Prophylaxis. ASHP Commission on Therapeutics and approved by the ASHP Board of Directors on November 14, 1998. *Am J Health Syst Pharm* 1999; 56:347.
5. Spirt MJ, Stanley S. Update on stress ulcer prophylaxis in critically ill patients. *Crit Care Nurse* 2006; 26:18.
6. Katz PO, Gerson LB, Vela MF. Guidelines for the diagnosis and management of gastroesophageal reflux disease. *Am J Gastroenterol* 2013;108:308-28.