

## Resource 8-3: Diagnostic Testing and Treatment Options for H. pylori

| Diagnostic Testing for <i>Helicobacter pylori</i><br>NB: The sensitivity of nonendoscopic tests that identify active <i>H. pylori</i> infection<br>is reduced by use of PPIs, bismuth, or antibiotics. |  |  |  |  |
|--|--|--|--|--|
| Invasive/Endoscopic testing  | Advantages   | Disadvantages  |  |  |
| Histology  | Excellent sensitivity and specificity  | Expensive<br>Requires infrastructure and trained<br>personnel  |  |  |
| Rapid urease testing   | Inexpensive and provides rapid results<br>Excellent specificity, very good sensitivity<br>in properly selected patients  | Sensitivity significantly reduced in the posttreatment setting   |  |  |
| Culture  | Excellent specificity<br>Allows determination of antibiotics<br>sensitivities  | Expensive<br>Difficult to perform<br>Not widely available<br>Only marginal sensitivity   |  |  |
| Polymerase chain reaction<br>(PCR)   | Excellent sensitivity and specificity<br>Allows determination of antibiotics<br>sensitivities  | Methodology not standardized across laboratories and not widely available  |  |  |
| Noninvasive/Nonendoscopic<br>testing   | Advantages   | Disadvantages  |  |  |
| Serologic antibody testing (quantitative and qualitative)  | Ease of specimen collection<br>Widely available<br>Very good negative predictive value<br>(NPV)  | Poor predictive value in populations with<br>low <i>H. pylori</i> prevalence<br>Cannot distinguish between active<br>infection and prior <i>H. pylori</i> exposure<br>Not recommended as a first-line test<br>Reimbursement not consistent |  |  |
| Urea breath tests ( <sup>13</sup> C and <sup>14</sup> C)   | Identifies active <i>H. pylori</i> infection<br>Excellent positive predictive value (PPV)<br>and NPV, regardless of <i>H. pylori</i><br>prevalence<br>Useful before and after <i>H. pylori</i> therapy | Higher cost<br>Need to discontinue antibiotics or proton<br>pump inhibitors at least 2 weeks prior to<br>testing   |  |  |
| Fecal antigen test   | Identifies active <i>H. pylori</i> infection<br>Excellent positive and negative<br>predictive values, regardless of <i>H. pylori</i><br>prevalence<br>Useful before and after <i>H. pylori</i> therapy | Polyclonal test less well validated than the<br>UBT in the posttreatment setting<br>Monoclonal test appears reliable before<br>and after antibiotic therapy<br>Unpleasantness associated with<br>collecting stool                          |  |  |

Abbreviations: PPI, proton pump inhibitor; PPV, positive predictive value; NPV, negative predictive value; UBT, urea breath tests.

Source: Chey WD, Wong BC; Practice Parameters Committee of the American Journal of Gastroenterology: American College of Gastroenterology Guideline on the Management of *Helicobacter pylori* Infection. *Am J Gastroenterol.* 2007;102(8):1808-1825. Available at http://s3.gi.org/physicians/guidelines/ManagementofHpylori.pdf

Theel ES. *Helicobacter pylori:* An update on diagnostic testing. Mayo Clinic Mayo Medical Laboratories Website. https://news.mayocliniclabs.com/2016/02/01/helicobacter-pylori-an-update-on-diagnostic-testing-hot-topic/

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| Regimen   |   |  |  |
|---|---|--|--|
|   | Duration  | Eradication rates  | Comments   |
| Standard-dose PPI* PO<br>BID (esomeprazole is<br>QD), clarithromycin 500<br>mg PO BID, amoxicillin<br>1000 mg PO BID  | 10–14 d   | 70%–85%  | Consider in non–<br>penicillin-allergic<br>patients who have not<br>previously received a<br>macrolide   |
| Standard-dose PPI* PO<br>BID, clarithromycin 500<br>mg PO BID,<br>metronidazole 500 mg<br>PO BID  | 10–14 d   | 70%–85%  | Consider in penicillin-<br>allergic patients who<br>have not previously<br>received a macrolide or<br>are unable to tolerate<br>bismuth quadruple<br>therapy |
| Bismuth subsalicylate<br>525 mg PO QID,<br>metronidazole 250 mg<br>PO QID, tetracycline<br>500 mg PO QID,<br>ranitidine 150 mg PO<br>BID <b>or</b> standard-dose<br>PPI* QD to BID  | 10–14 d   | 75%–90%  | Consider in penicillin-<br>allergic patients   |
| PPI + amoxicillin 1 g PO<br>BID followed by: PPI,<br>clarithromycin 500 mg<br>PO, tinidazole 500 mg<br>PO BID   | 5 d<br>5 d  | >90%   | Requires validation in<br>North America  |
| Standard dosages for PPIs are<br>ansoprazole, 30 mg PO; omepi<br>Note: The above recommended<br>1. Bismuth 525 mg PO QID, + n<br>2. Lansoprazole 30 mg PO BID<br>3. Omeprazole 20 mg PO BID +<br>4. Esomeprazole 40 mg PO QD<br>5. Rabeprazole 20 mg PO BID - | o inhibitor; PO, orally; QD, once da<br>as follows:<br>razole, 20 mg PO; pantoprazole, 4<br>treatments are not all FDA approv<br>netronidazole 250 mg PO QID + te<br>+ clarithromycin 500 mg PO BID +<br>clarithromycin 500 mg PO BID +<br>+ clarithromycin 500 mg PO BID<br>+ clarithromycin 500 mg PO BID +<br>clarithromycin 500 mg PO BID + | 0 mg PO; rabeprazole, 20 mg PO<br>red. The FDA-approved regimens<br>etracycline 500 mg PO QID × 2 wk<br>+ amoxicillin 1 g PO BID × 10 days<br>amoxicillin 1 g PO BID × 10 days.<br>+ amoxicillin 1 g PO BID × 10 day<br>amoxicillin 1 g PO BID × 7 days. | ; esomeprazole, 40 mg PO.<br>e are as follows:<br>$x + H_2RA$ as directed × 4 wk.<br>s.  |