Diagnosis and Management of Foodborne Illness

TIMOTHY L. SWITAJ, MD, *Reynolds Army Community Hospital, Fort Sill, Oklahoma* KELLY J. WINTER, DO, *William Beaumont Army Medical Center, Fort Bliss, Texas* SCOTT R. CHRISTENSEN, MD, *Reynolds Army Community Hospital, Fort Sill, Oklahoma*

The Centers for Disease Control and Prevention estimates that each year, one in six Americans will experience a foodborne illness. The most common causes in the United States are viruses, such as norovirus; bacteria, such as *Salmonella, Escherichia coli, Campylobacter*, and *Listeria*; and parasites, such as *Toxoplasma gondii* and *Giardia*. Resources are available to educate consumers on food recalls and proper handling, storage, and cooking of foods. Diagnosis and management of a foodborne illness are based on the history and physical examination. Common symptoms of foodborne illnesses include vomiting, diarrhea (with or without blood), fever, abdominal cramping, headache, dehydration, myalgia, and arthralgias. Definitive diagnosis can be made only through stool culture or more advanced laboratory testing. However, these results should not delay empiric treatment if a foodborne illness is suspected. Empiric treatment should focus on symptom management, rehydration if the patient is clinically dehydrated, and antibiotic therapy. Foodborne illnesses should be reported to local and state health agencies; reporting requirements vary among states. (*Am Fam Physician*. 2015;92(5):358-365. Copyright © 2015 American Academy of Family Physicians.)

CME This clinical content conforms to AAFP criteria for continuing medical education (CME). See CME Quiz Questions on page 336.

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▶ Patient information: A handout on this topic, written by the authors of this article, is available at http://www.aafp.org/ afp/2015/0901/p358-s1. html.

oodborne illness can be caused by a multitude of microorganisms such as viruses, bacteria, and parasites. Foodborne illness is a worldwide problem, and U.S. outbreaks often garner media attention and result in food recalls. Foodborne illnesses are becoming a greater challenge because of new and emerging microorganisms and toxins, the growth of antibiotic resistance, increasing food contamination caused by new environments and methods of food production, and an increase in multistate outbreaks.1 There are more than 250 identified pathogens that cause foodborne illness. The Centers for Disease Control and Prevention (CDC) estimates that one in six Americans (approximately 48 million) will become sick from a foodborne pathogen each year, resulting in 128,000 hospitalizations and 3,000 deaths.² Most foodborne illnesses, hospitalizations, and deaths are caused by one of eight common pathogens: norovirus, nontyphoidal Salmonella, Clostridium perfringens, Campylobacter, Staphylococcus aureus, Toxoplasma gondii, Listeria monocytogenes, and Shiga toxin-producing Escherichia coli² (Table 1³). E. coli is commonly divided into two broad types, Shiga toxin-producing-of which

E. coli O157:H7 is the best studied—and non-Shiga toxin—producing, which includes enteropathogenic, enteroinvasive, enteroag-gregative, and diffusely adherent *E. coli*. New pathogens emerge constantly, whereas others decrease in significance or disappear altogether. Predicting the emergence or disappearance of specific pathogens—other than in the setting of an identified outbreak—is difficult and has not significantly prevented or limited foodborne illnesses.

Prevention

Prevention is the first step in combatting foodborne illnesses. Consumer information on food safety is available from the http://www.cdc.gov/foodsafety/ CDC at facts.html or from the U.S. Food and Drug Administration at http://www.fda. gov/Food/FoodborneIllnessContaminants/ FoodborneIllnessesNeedToKnow/default.htm or http://www.foodsafety.gov, or by calling (888) SAFEFOOD ([888] 723-3366). Guidelines for safely handling and preparing food should be followed regardless of the setting in which food is consumed. Although these recommendations have been shown to reduce the risk of foodborne illness in the United States, the risk increases with travel.

Table 1. Summary of Foodborne Illnesses

Organism (common name of illness)	Time from ingestion to symptom onset	Signs and symptoms	Duration	Food sources
<i>Bacillus cereus</i> (food poisoning)	10 to 16 hours	Abdominal cramps, nausea, watery diarrhea	24 to 48 hours	Gravy, meats, stews, vanilla sauce
Campylobacter jejuni (campylobacteriosis)	2 to 5 days	Abdominal cramps, diarrhea (may be bloody), fever, vomiting	2 to 10 days	Contaminated water, raw or undercooked poultry, unpasteurized milk
Clostridium botulinum (botulism)	12 to 72 hours	Blurred vision, diarrhea, difficulty swallowing, double vision, muscle weakness, vomiting; can cause respiratory failure and death	Variable	Fermented fish, improperly canned foods (especially home-canned vegetables), potatoes baked in aluminum foil
Clostridium perfringens (food poisoning)	8 to 16 hours	Intense abdominal cramps, watery diarrhea	Usually 24 hours	Dried or precooked foods, gravy, meats, poultry, undercooked foods
Cryptosporidium (cryptosporidiosis)	2 to 10 days	Abdominal cramps, diarrhea (usually watery), slight fever	May be remitting and relapsing over weeks to months	Contaminated drinking water, cooked foods that are not reheated after contact with an infected food handler, uncooked foods
Cyclospora cayetanensis (cyclosporiasis)	1 to 14 days, usually at least 1 week	Abdominal cramps, diarrhea (usually watery), fatigue, loss of appetite, nausea, substantial weight loss, vomiting	May be remitting and relapsing over weeks to months	Contaminated raw produce (e.g., basil, imported berries, lettuce)
Escherichia coli (traveler's diarrhea)	1 to 3 days	Abdominal cramps, vomiting, watery diarrhea	3 to 7 days	Food or water contaminated with human feces
E. coli O157:H7 (Shiga toxin– producing E. coli or hemorrhagic colitis)	1 to 8 days	Abdominal pain, severe diarrhea (often bloody), vomiting; can cause kidney failure	5 to 10 days	Contaminated drinking water, contaminated raw produce (e.g., sprouts), undercooked beef (especially hamburger), unpasteurized milk or juice
Hepatitis A	15 to 50 days (mean = 28 days)	Abdominal pain, dark urine, diarrhea, fever, headache, jaundice, nausea	2 weeks to 3 months	Contaminated drinking water, contaminated raw produce, cooked foods that are not reheated after contact with an infected food handler, shellfish from contaminated water, uncooked foods
Listeria monocytogenes (listeriosis)	9 to 48 hours for gastrointestinal symptoms, 2 to 6 weeks for invasive disease	Diarrhea, fever, muscle aches, nausea; pregnant women may have mild flulike illness, and infection can lead to premature delivery or stillbirth; older adults and immunocompromised patients may develop bacteremia or meningitis	Variable	Deli meats, unpasteurized milk, soft cheeses made with unpasteurized milk
Norovirus (food poisoning, viral gastroenteritis, winter diarrhea)	12 to 48 hours	Abdominal cramps, diarrhea (more common in adults), fever, headache, nausea, vomiting (more common in children)	12 to 60 hours	Contaminated drinking water, contaminated raw produce, cooked foods that are not reheated after contact with an infected food handler, shellfish from contaminated water, uncooked foods <i>continues</i>

Organism (common name of illness)	Time from ingestion to symptom onset	Signs and symptoms	Duration	Food sources
<i>Salmonella</i> (salmonellosis)	6 to 48 hours	Abdominal cramps, diarrhea, fever, vomiting	4 to 7 days	Cheese, contaminated raw produce, eggs, meat, poultry unpasteurized milk or juice
<i>Shigella</i> (shigellosis or bacillary dysentery)	4 to 7 days	Abdominal cramps, diarrhea, fever; stools may contain blood and mucus	24 to 48 hours	Contaminated drinking water, contaminated raw produce, cooked foods that are not reheated after contact with an infected food handler, uncooked foods
<i>Staphylococcus aureus</i> (food poisoning)	1 to 6 hours	Abdominal cramps, diarrhea, fever, sudden onset of severe nausea and vomiting	24 to 48 hours	Unrefrigerated or improperly refrigerated cream pastries, meats, and potato or egg salad
Vibrio parahaemolyticus	4 to 96 hours	Abdominal cramps, fever, nausea, vomiting, watery diarrhea (occasionally bloody)	2 to 5 days	Undercooked or raw seafood
Vibrio vulnificus	1 to 7 days	Abdominal pain, bleeding under the skin, bloodborne infection, diarrhea, fever, ulcers requiring surgical removal, vomiting; can be fatal to persons with liver disease or weakened immune systems	2 to 8 days	Undercooked or raw seafood (especially oysters)

Table 1. Summary of Foodborne Illnesses (continued)

Adapted from U.S. Food and Drug Administration. Foodborne illnesses: what you need to know. http://www.fda.gov/food/resourcesforyou/consumers/ ucm103263.htm. Accessed May 7, 2015.

Diagnosis HISTORY

Foodborne illness can have various presentations, ranging from clinically mild illness that requires only outpatient care to severe illness that requires hospitalization. Most foodborne illnesses are associated with vomiting or diarrhea (more than three loose stools in 24 hours). Other common symptoms include fever, bloody diarrhea, abdominal cramping, headache, dehydration, myalgia, and arthralgias.⁴ Patients may have several symptoms or only one. The history is the most important step in evaluating a patient with diarrheal illness (*Table 2*).⁵

None of the symptoms of foodborne illness is specific, so the clinician must consider the history, epidemiologic features, and objective findings to make an accurate diagnosis. Symptoms and time of onset can narrow the differential diagnosis (*Table 3*) and help identify a likely pathogen.⁴ Early onset of vomiting and diarrhea results from ingestion of preformed toxins, most often *S. aureus* or *Bacillus cereus*. Diarrhea within 24 hours of ingestion is most likely caused by *C. perfringens* or *B. cereus*. Diarrhea within 24 to 48 hours of ingestion is most often caused by *Campylobacter jejuni* in individual cases

or *Salmonella* in outbreaks.⁴ Foodborne illnesses commonly associated with fever are caused by *Vibrio cholerae* non-O1, *Shigella*, and *C. jejuni (Table 4)*. Enterotoxins in the small bowel caused by *E. coli*, *C. perfringens*, and viruses produce excessive secretions of fluids and electrolytes that overwhelm the large bowel; therefore, they are typically associated with watery diarrhea. Bloody diarrhea with abdominal pain should prompt consideration of inflammatory damage to the intestinal mucosa or an infection (e.g., *C. jejuni, Salmonella enteritidis*, enteroinvasive *E. coli*) affecting the large bowel.

PHYSICAL EXAMINATION

The physical examination can help narrow the differential diagnosis, and vital signs can help determine the severity of volume depletion. Orthostatic pulse and blood pressure changes should be noted, and a basic general physical examination should be performed, with assessment of skin turgor, the abdomen, mucous membranes, and mental status.⁵

ANCILLARY TESTING

Watchful waiting is often the most appropriate option in the initial diagnosis and management of foodborne illness; ancillary testing is usually not necessary. If testing is performed, stool culture can provide a definitive diagnosis of infectious diarrhea and is useful for outbreak identification. In most outpatients who have self-limiting gastroenteritis, a stool culture does not affect management.⁴ Bacteria are the most common cause of non–self-limiting foodborne illness; however, stool cultures are positive in less than 40% of cases.^{4,5} Newer techniques such as polymerase chain reaction testing have become readily available and provide more rapid, reliable determination of specific pathogens. An organism-specific diagnosis can help clinicians to narrow treatment recommendations, aid public health professionals, and prevent unnecessary procedures.

Other tests that can be considered include serum chemistry (including albumin levels), C-reactive protein

levels, complete blood count, blood cultures, urinalysis, abdominal radiography, anoscopy, and endoscopy, if warranted by the severity and pattern of symptoms.⁵ In severe cases of infectious diarrhea, toxic megacolon should be considered, which can be identified on plain abdominal radiography.⁴ Severe inflammatory changes can also be seen on computed tomography. It may be reasonable to obtain blood cultures in patients with fever and diarrhea (with or without blood), because up to 1% of cases of nontyphoidal *Salmonella* infections are associated with bacteremia.^{4,5} Sigmoidoscopy or colonoscopy may be useful in hospitalized patients with bloody diarrhea to obtain tissue and histology, which could aid in the diagnosis.

Stool microscopy is rarely diagnostic, but the presence of red and white blood cells may signal a colonic source.⁴

Its primary use is identification of ova, cysts, and parasites, although antigen testing is more sensitive and specific for *Giardia*. Microscopic evaluation for fecal polymorphonuclear leukocytes or lactoferrin measurements may be useful if an inflammatory etiology is suspected. A positive stool culture is more likely when analysis indicates an inflammatory process. Compared with leukocyte examinations, lactoferrin measurements are more sensitive but more expensive, have a higher false-positive rate, and require a fresh-cup sample examined by an experienced microscopist.⁵

Management SYMPTOMATIC TREATMENT

Use of antidiarrheal medications, including antimotility agents, anticholinergics, and adsorbents, is not recommended in children, especially those younger than two years, and is discouraged if infection with Shiga toxin–producing *E. coli* is suspected.⁶⁻⁸ Symptomatic treatment with loperamide (Imodium) and bismuth subsalicylate (Pepto-Bismol) is effective and may be considered in adults with uncomplicated acute or traveler's diarrhea.⁹⁻¹¹ Although loperamide is more effective than bismuth subsalicylate, it is not recommended for patients with hematochezia and systemic symptoms because it may increase the risk of invasive disease.¹⁰

In patients with clinically significant vomiting, antiemetics can alleviate symptoms and reduce the need for hospitalization and

Table 2. Clues to the Differential Diagnosis of Diarrhea

Findings from patient history	Comment
Acute abdominal pain, fever, and vomiting	Together, these symptoms raise suspicion for infectious diarrhea
Dietary factors	Recent changes in diet and ingestion of foods included in recent recalls or undercooked foo should raise suspicion for foodborne illness
Duration of symptoms	Longer duration raises concern for dehydratio
Employment history	Persons who work at child care centers or in close contact with others are at risk of viral diarrhea
Exposure to other persons with diarrhea	Cross-contamination and transmission of pathogens are possible; may help narrow differential if cause is known in the other person
Hospitalization or nursing home admission	Raises suspicion for Clostridium difficile infection
Immunocompromise	Raises suspicion for atypical causes of diarrhe
Medical history	Can help determine possible comorbidities th suggest a cause
Recent antibiotic use	Raises suspicion for C. difficile infection
Stool characteristics (bloody, foul smelling, watery)	Bloody diarrhea raises suspicion for Salmonell Shigella, or enterohemorrhagic Escherichia coli infection (or mesenteric ischemia in at- risk populations)
	Foul-smelling stools in patients with recent hospitalization or antibiotic use raise suspicion for C. <i>difficile</i> or <i>Giardia</i> infection
	Watery stools raise suspicion for viral cause or <i>Giardia</i> infection
Travel history	Travel to foreign countries, especially non- Western countries, should raise suspicion fo infectious diarrhea

Diagnosis	Common presenting symptoms	Diagnostic studies
Acute cholecystitis	Decreased appetite, fever, jaundice, nausea, right upper-quadrant abdominal pain, vomiting	CBC, C-reactive protein level, liver function testing, right upper-quadrant ultrasonography
Acute hepatitis	Abdominal pain, arthralgias, arthritis, fever, jaundice, malaise, nausea, vomiting	Ammonia levels, hepatitis panel, liver biopsy, liver function testing, ultrasonography
Diverticular disease	Fever, left lower-quadrant abdominal pain	Abdominal CT, CBC; contrast enema and colonoscopy may be considered
Inflammatory bowel disease	Abdominal pain, chronic diarrhea, occasional bloody diarrhea, weight loss	Colonoscopy with tissue biopsy, negative stool culture
Mesenteric ischemia	Abdominal pain, diarrhea, hematochezia, melena, weight loss	Abdominal CT, arterial blood gas levels, blood chemistry panel, CBC, colonoscopy, electrocardiography, lactate levels, magnetic resonance angiography
Viral syndromes	Abdominal pain, anorexia, diarrhea, fever, nausea, vomiting	Diagnosis is generally clinical; may be confirmed by antigen- detecting enzyme immunoassay, immunofluorescence assay, microscopy, polymerase chain reaction testing, serology, or viral culture (although routine use of these tests is not necessary)

Table 3. Differential Diagnosis of Foodborne Illness

intravenous fluid administration. Multiple studies support the use of a single dose of ondansetron (Zofran) in children with gastroenteritis-related vomiting.¹²⁻¹⁴ The use of antiemetics in adults with gastroenteritis is reasonable, but data about adverse effects are lacking.

DEHYDRATION

Many physicians are reluctant to use oral rehydration therapy, despite its proven effectiveness in the management of diarrhea-associated dehydration.^{6,15} It has been

Fever	Vomiting
Characteristically associated	Characteristically associated
Campylobacter jejuni Shiqella	Bacillus cereus (emetic syndrome)
Vibrio cholerae non-01	Norwalk virus
Often associated	Staphylococcus aureus
Norwalk virus	Often associated
Salmonella	Clostridium botulinum
Vibrio parahaemolyticus	V. cholerae O1
, ,	V. parahaemolyticus

proven to prevent and treat dehydration in patients of all ages.^{5,16-19} Guidelines that recommend the use of oral rehydration therapy for mild to moderate dehydration in infants and children have been published by the World Health Organization, the American Academy of Pediatrics, and the CDC.^{6,7,20} Oral rehydration solutions contain a blend of electrolytes, as well as carbohydrates. Sports drinks and soft drinks have a high carbohydrate-to-sodium ratio and total osmolality, and can exacerbate diarrhea.¹⁹

Clinical assessment should be used to guide rehydration therapy. Children—especially infants—are predisposed to dehydration and require more diligence in determining hydration status. CDC guidelines recommend the initial use of oral rehydration therapy with replacement of continuing fluid losses in children with mild to moderate dehydration. Children with severe dehydration should be hospitalized and given intravenous fluids.^{6,7} When oral rehydration therapy or intravenous fluid administration is used in infants, care should be taken to minimize interruptions in breastfeeding or formula feeding.^{6,7}

EMPIRIC ANTIBIOTICS

Most cases of acute infectious diarrhea are viral, and improper use of empiric antibiotics is associated with

Clinical recommendation	Evidence rating	References
Stool cultures are the diagnostic standard for bacterial foodborne illness; however, culture results are positive in less than 40% of cases.	С	4, 5
A single dose of ondansetron (Zofran) is recommended in children with clinically significant gastroenteritis-related vomiting.	А	12-14
Oral rehydration therapy is effective in preventing and treating dehydration in patients of all ages.	А	16-19
Empiric antibiotic therapy should be considered in cases of suspected foodborne illness if the patient is febrile and has signs of invasive disease, if symptoms have persisted for more than one week or are severe, or if hospitalization may be required.	С	10, 16, 17, 19

A = consistent, good-quality patient-oriented evidence; B = inconsistent or limited-quality patient-oriented evidence; C = consensus, diseaseoriented evidence, usual practice, expert opinion, or case series. For information about the SORT evidence rating system, go to http://www.aafp. org/afpsort.

increased morbidity caused by adverse effects and *Clos-tridium difficile* colitis. Empiric antibiotics should be considered in cases of suspected foodborne illness only if the patient is febrile and has signs of invasive disease (e.g., gross hematochezia, leukocytes on fecal smear), if symptoms have persisted for more than one week or are severe (i.e., more than eight liquid stools per day), or if hospitalization may be required.^{10,16,17,19} A fluoroquino-lone (or trimethoprim/sulfamethoxazole in children) is generally recommended for empiric antibiotic therapy.⁵ Stool testing should still be performed.

Empiric antibiotic therapy decreases the duration of symptoms in patients with traveler's diarrhea.^{5,9} Enterotoxigenic *E. coli* is the most common cause of traveler's diarrhea worldwide and is generally susceptible to ciprofloxacin, but azithromycin (Zithromax) is equally effective and a better choice in areas where fluoroquinolone-resistant *C. jejuni* is present.^{9,21} Patients with diarrhea of more than 10 days' duration that is associated with fatty or foul-smelling stools, cramps, bloating, and weight loss can be treated empirically for *Giardia* infection.²² Because of an increased risk of hemolytic uremic syndrome, patients receiving empiric antibiotic therapy should be monitored closely if Shiga toxin–producing *E. coli* infection is suspected.^{4,23}

TARGETED ANTIBIOTICS

If empiric treatment has not been initiated, antibiotic therapy may be indicated once stool culture, bacterial toxin, or microscopy results are available. Antibiotic therapy can shorten the duration of symptoms and may prevent bacteremia in older adults, newborns, and immunocompromised patients.⁴ Recommended treatment regimens for foodborne pathogens are outlined in *Table 5.*^{8,18,19,21-24} Ciprofloxacin is no longer recommended for treatment of *Campylobacter* infection; a macrolide, such as erythromycin, is recommended instead.²⁴

Outbreak Surveillance and Reporting

Although reporting requirements vary, most states require physicians to report cases of hepatitis A and botulism, and Salmonella, Shiga toxin-producing E. coli, Listeria, Shigella, and Vibrio infections. A definitive diagnosis is not necessary to report a suspected foodborne illness. Outbreaks can be declared by local, county, state, or national agencies; reporting outbreaks to the CDC is left to the discretion of the agency. Physicians should avoid attributing cases of foodborne illness to specific food sources without definitive testing and reporting. In cases of suspected foodborne illness or in the setting of an outbreak, stool and vomitus samples should be sent for testing. All reports of outbreaks are entered into the CDC's Foodborne Disease Outbreak Surveillance System, after which the data are analyzed to monitor and identify the root cause of the outbreak. Once an outbreak has occurred, prevention strategies should focus on educating the public through the news media and correcting the underlying cause, such as through food product recalls. Information on outbreaks, outbreak surveillance, and identified trends can be found at http://www.cdc.gov/foodsafety/fdoss/ index.html or by calling the CDC at (800) CDC-INFO ([800] 232-4636).

Data Sources: A PubMed search was completed using the key terms foodborne illness, infectious diarrhea, food poisoning, salmonella, *Shigella, Escherichia coli, Campylobacter*, food safety, hemolytic-uremic syndrome, oral rehydration solution, oral rehydration therapy, traveler's diarrhea, and bloody diarrhea. We also searched Essential

Pathogen	Recommended regimen for adults	Recommended regimen for children
Bacteria Campylobacter	Erythromycin, 500 mg 2 times per day for 5 days or Azithromycin (Zithromax), 500 mg on day 1, then 250 mg on days 2 through 5	Azithromycin, 10 mg per kg per day for 3 to 7 days
Escherichia coli (non-Shiga toxin–producing)	Ciprofloxacin, 500 mg 2 times per day for 3 days or TMP/SMX, 160/800 mg 2 times per day for 3 to 7 days* or Azithromycin, 500 mg on day 1, then 250 mg on days 2 through 5	TMP-SMX, 5/25 mg per kg 2 times per day for 3 days* <i>or</i> Azithromycin, 10 mg per kg per day for 3 to 7 days
E. coli (Shiga toxin–producing)	Not recommended	Not recommended
Salmonella (nontyphoidal)	Generally not recommended† <i>If indicated:</i> Ciprofloxacin, 500 mg 2 times per day for 5 to 7 days <i>or</i> Azithromycin, 500 mg on day 1, then 250 mg on days 2 through 5‡ <i>or</i> Ceftriaxone, 1 to 2 g per day intramuscularly or intravenously for 5 to 7 days	Generally not recommended † <i>If indicated:</i> TMP-SMX, 5/25 mg per kg 2 times per day for 5 to 7 days* <i>or</i> Azithromycin, 10 mg per kg per day for 3 to 7 days <i>or</i> Ceftriaxone, 50 to 100 mg per kg per day intramuscularly or intravenously for 5 to 7 days
Shigella	Ciprofloxacin, 500 mg 2 times per day for 3 days or Azithromycin, 500 mg on day 1, then 250 mg on days 2 through 5	TMP-SMX, 5/25 mg per kg 2 times per day for 3 days† or Azithromycin, 10 mg per kg per day for 3 to 7 days
Parasites		
Cryptosporidium	Generally not recommended§	Generally not recommended§
Entamoeba histolytica	Metronidazole (Flagyl), 750 mg 3 times per day for 5 to 10 days <i>plus</i> Paromomycin, 500 mg 3 times per day for 7 days, or iodoquinol (Yodoxin), 650 mg 3 times per day for 7 days	Metronidazole, 30 to 50 mg per kg per day in 3 divided doses for 7 to 10 days <i>plus</i> Paromomycin, 25 to 35 mg per kg per day in 3 divided doses for 5 to 10 days
Giardia	Metronidazole, 250 to 750 mg 3 times per day for 7 to 10 days	Metronidazole, 30 to 50 mg per kg per day in 3 divided doses for 7 to 10 days

Table 5. Recommended Treatment Regimens for Foodborne Pathogens

TMP/SMX = trimethoprim/sulfamethoxazole.

*-If susceptible.

+—Antibiotic therapy is not recommended except in severe infections, patients younger than 6 months or older than 50 years, immunocompromised patients, and patients with prostheses, valvular heart disease, severe atherosclerosis, malignancy, or decreased renal function.

+—Alternative dosing regimen for azithromycin in adults with salmonellosis: 1,000 mg on day 1, then 500 mg per day for 2 to 6 days.

§—Treatment for cryptosporidiosis is generally not recommended in immunocompetent patients. For persistent symptoms, some clinicians prescribe nitazoxanide (Alinia), although it has not been approved for this use and has not been proven superior to placebo. Dosing for adults and children 12 years and older is 500 mg orally every 12 hours for 3 days. Dosing for children 4 to 11 years of age is 200 mg of oral suspension every 12 hours for 3 days. Dosing for children 1 to 3 years of age is 100 mg of oral suspension every 12 hours for 3 days.

Information from references 8, 18, 19, and 21 through 24.

The views expressed in this article are those of the authors and do not necessarily reflect the official policy of the Department of Defense, the Department of the Army, or the U.S. Army Medical Department.

The Authors

TIMOTHY L. SWITAJ, MD, is deputy commander for clinical services at Reynolds Army Community Hospital, Fort Sill, Okla.

KELLY J. WINTER, DO, is a staff family physician at William Beaumont Army Medical Center, Fort Bliss, Tex.

SCOTT R. CHRISTENSEN, MD, is officer-in-charge at the family medicine clinic #1 at Reynolds Army Community Hospital.

Address correspondence to Timothy L. Switaj, MD, at timothy.l.switaj. mil@mail.mil. Reprints are not available from the authors.

REFERENCES

- Centers for Disease Control and Prevention. CDC and food safety. http://www.cdc.gov/foodsafety/cdc-and-food-safety.html. Accessed May 15, 2014.
- Centers for Disease Control and Prevention. Food safety: foodborne illness, foodborne disease, (sometimes called "food poisoning"). http:// www.cdc.gov/foodsafety/facts.html. Accessed May 15, 2014.
- U.S. Food and Drug Administration. Foodborne illnesses: what you need to know. http://www.fda.gov/food/resourcesforyou/consumers/ ucm103263.htm. Accessed May 7, 2015.
- 4. Conlon C. Food-borne diarrheal illness. In: Cohen J, ed. *Infectious Diseases*. 3rd ed. St. Louis, Mo.: Mosby; 2010.
- Guerrant RL, Van Gilder T, Steiner TS, et al.; Infectious Diseases Society of America. Practice guidelines for the management of infectious diarrhea. *Clin Infect Dis.* 2001;32(3):331-351.
- Duggan C, Santosham M, Glass RI; Centers for Disease Control and Prevention. The management of acute diarrhea in children: oral rehydration, maintenance, and nutritional therapy. *MMWR Recomm Rep.* 1992;41(RR-16):1-20.
- American Academy of Pediatrics, Provisional Committee on Quality Improvement, Subcommittee on Acute Gastroenteritis. Practice parameter: the management of acute gastroenteritis in young children. *Pediatrics*. 1996;97(3):424-435.
- Cimolai N, Carter JE, Morrison BJ, Anderson JD. Risk factors for the progression of *Escherichia coli* O157:H7 enteritis to hemolytic-uremic syndrome [published correction appears in *J Pediatr.* 1990;116(6):1008]. *J Pediatr.* 1990;116(4):589-592.

- Löscher T, Alberer M. Clinical presentation and management of travelers' diarrhea. In: Keystone JS, ed. *Travel Medicine*. 3rd ed. Philadelphia, Pa.: Elsevier/Saunders; 2012.
- 10. Steffen R. Worldwide efficacy of bismuth subsalicylate in the treatment of travelers' diarrhea. *Rev Infect Dis.* 1990;12(suppl 1):S80-S86.
- DuPont HL, Flores Sanchez J, Ericsson CD, et al. Comparative efficacy of loperamide hydrochloride and bismuth subsalicylate in the management of acute diarrhea. Am J Med. 1990;88(6A):155-195.
- DeCamp LR, Byerley JS, Doshi N, Steiner MJ. Use of antiemetic agents in acute gastroenteritis: a systematic review and meta-analysis. Arch Pediatr Adolesc Med. 2008;162(9):858-865.
- 13. Levine DA. Antiemetics for acute gastroenteritis in children. *Curr Opin Pediatr.* 2009;21(3):294-298.
- Ramsook C, Sahagun-Carreon I, Kozinetz CA, Moro-Sutherland D. A randomized clinical trial comparing oral ondansetron with placebo in children with vomiting from acute gastroenteritis. *Ann Emerg Med.* 2002;39(4):397-403.
- Snyder JD. Use and misuse of oral therapy for diarrhea: comparison of US practices with American Academy of Pediatrics recommendations. *Pediatrics*. 1991;87(1):28-33.
- Santosham M, Daum RS, Dillman L, et al. Oral rehydration therapy of infantile diarrhea: a controlled study of well-nourished children hospitalized in the United States and Panama. N Engl J Med. 1982; 306(18):1070-1076.
- Santosham M, Burns B, Nadkarni V, et al. Oral rehydration therapy for acute diarrhea in ambulatory children in the United States: a doubleblind comparison of four different solutions. *Pediatrics*. 1985;76(2): 159-166.
- Tamer AM, Friedman LB, Maxwell SR, Cynamon HA, Perez HN, Cleveland WW. Oral rehydration of infants in a large urban U.S. medical center [published correction appears in *J Pediatr.* 1986;108(1):160]. *J Pediatr.* 1985;107(1):14-19.
- 19. Avery ME, Snyder JD. Oral therapy for acute diarrhea. The underused simple solution. *N Engl J Med*. 1990;323(13):891-894.
- World Health Organization; UNICEF. Clinical management of acute diarrhoea: WHO/UNICEF joint statement. http://www.who.int/iris/ handle/10665/68627. Accessed July 7, 2015.
- Adachi JA, Ericsson CD, Jiang ZD, et al. Azithromycin found to be comparable to levofloxacin for the treatment of US travelers with acute diarrhea acquired in Mexico. *Clin Infect Dis.* 2003;37(9):1165-1171.
- 22. DuPont HL; Practice Parameters Committee of the American College of Gastroenterology. Guidelines on acute infectious diarrhea in adults. *Am J Gastroenterol.* 1997;92(11):1962-1975.
- Wong CS, Jelacic S, Habeeb RL, Watkins SL, Tarr PI. The risk of the hemolytic-uremic syndrome after antibiotic treatment of *Escherichia coli* 0157:H7 infections. *N Engl J Med*. 2000;342(26):1930-1936.
- Smith KE, Besser JM, Hedberg CW, et al.; Investigation Team. Quinolone-resistant *Campylobacter jejuni* infections in Minnesota, 1992-1998. N Engl J Med. 1999;340(20):1525-1532.