



# **Gastrointestinal Pathogen Panel Guidance**

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## **Background:**

Many pathogens, including bacteria, parasites, and viruses can cause infectious diarrhea. Previously, many of the pathogens responsible could only be isolated using traditional techniques such as stool culture or ova and parasite exam that were often time consuming and lacked sensitivity. To improve the detection of intestinal pathogens, the microbiology lab has introduced multiplex PCR testing using the FilmArray Gastrointestinal (GI) panel, which detects 22 common viruses, bacteria, and parasites that cause infectious diarrhea. Results are typically available in about one hour.

# Testing:

This panel <u>will replace traditional stool culture (STOCU) and the *Giardia* and *Cryptosporidium* antigen (OVPSC) screen. Stool cultures with isolation and susceptibility will now only be obtained reflexively when *Shigella* and *Campylobacter* are detected by the GI panel. The stool Ova & Parasite microscopic test (OVPAR), used primarily for individuals with a history of travel to foreign countries, will still be available but should be reserved for those who have **both** clinical symptoms and an epidemiologic exposure strongly suggestive of a parasite not detected by the GI Panel.</u>

*C. difficile* testing is part of the panel but will not be reported. The reason for this is the microbiology lab and antimicrobial stewardship team have determined that PCR testing alone is inadequate for accurately identifying those with *C. difficile* infection (CDI) who require treatment.<sup>1,2</sup> If CDI is suspected the *C. difficile* toxin assay is recommended as it is the preferred method to diagnose CDI. In cases where the GI panel *C. difficile* PCR is positive and no *C. difficile* test has been ordered the microbiology lab will contact the clinician so they can decide if further testing for CDI is warranted.

## **Restriction:**

The GI panel may only be ordered for patients who have been hospitalized for less than 5 days; otherwise it will require approval from the microbiology director (or their designee) who will consult with the antimicrobial stewardship program or the infectious disease service when needed. The panel may also only be ordered once per admission. These restrictions will be put in place as data show that routine stool cultures from patients with diarrhea that develops after 72 hours of hospitalization are typically low yield for standard bacteria and parasites.<sup>3,4</sup>

#### Interpretation:

Results of PCR testing for stool pathogens must be taken into clinical context when making treatment decisions. Previously treatment decisions were made based upon either clinical presentation or traditional microbiologic diagnostic techniques. PCR testing is much more sensitive than traditional techniques and allows for the detection of low numbers of pathogens.<sup>5,6,7</sup> The clinical correlation of PCR results with the need for treatment and clinical outcomes has not been established. Studies evaluating stool PCR testing frequently detect more than one enteric pathogen in patient's stool and data are not available to determine the causative organism in these situations.<sup>5,7</sup> Additionally low levels of stool pathogens have been detected in healthy persons and <u>all decisions regarding need for treatment must</u> be taken into clinical context of the patient.

## **Treatment Recommendations:**

Taking all this into account most gastrointestinal infections due to common bacterial and viral causes are self-limited in nature and do not require antimicrobial therapy. Symptoms typically resolve within 7 days in a normal host and therapy should focus on providing supportive care by replacing fluid and electrolyte losses. The use of antimicrobial therapy must be carefully weighed against unintended and potentially harmful consequences, including antimicrobial-resistant infections, side effects of treatment with antimicrobial agents, super-infections when normal flora are eradicated by antimicrobial agents, the prolongation of a carrier state (particularly in *Salmonella*) and the possibility of induction of disease-producing phages by antibiotics (such as Shiga-toxin phage induced by quinolone antibiotics).

The role of antimicrobial therapy depends on the implicated pathogen. In general, antimicrobial agents are only used to treat parasitic infections as well as select bacterial infections. Misuse and overuse of antibiotics in the treatment of diarrheal illness has played an important role in the development of drug resistance, which complicates treatment of those infections in which antibiotics are indicated. Listed below are suggested criteria for treatment of specific pathogens. These recommendations apply to generally healthy persons unless otherwise noted. There is a paucity of data regarding the efficacy of antimicrobials in a number of the pathogens detected on the panel and in these cases antibiotics are generally only recommended in severe or non-resolving cases or those at risk for severe disease such as immunocompromised patients. Areas where antibiotics are always indicated have been delineated as has areas where the data are less clear. In cases where data are lacking, clinical judgment and the assessment of the risk verses benefit ration must be considered.

# Table 1 – Etiology and Treatment Recommendations<sup>8,9,10,11</sup>

Pathogen	Common Presentation	Commonly Implicated Sources and Seasonality	Treatment Recommendations	Antibiotics (If Indicated)
Bacteria	·			•
Campylobacter	Fever, abdominal cramps, and diarrhea within 6-48 hours, fecal leukocytes often present	Poultry, unpasteurized milk and dairy products Peak season – spring, summer	Most patients recover without antimicrobial therapy. Antibiotics have been shown to reduce symptom duration by 1.3 days and are recommended for severe illness (high fever, bloody, severe, or worsening diarrhea) or risk factors for complications (elderly, pregnant women, immunocompromised).	Azithromycin 500 mg daily x 3 days Fluoroquinolone x 3 days* Immunocompromised patients may require prolonged therapy (7-14 days)
Clostridium difficile (toxin A/B)	More than 3 watery, loose, or unformed stools within 24 hours; lab findings may include leukocytosis and elevated creatinine	Recent antibiotic use, especially broad spectrum agents	Test not reported on panel. If CDI is suspected, order the <i>C. difficile</i> toxin assay.	Metronidazole 500 mg TID x 10-14 days Vancomycin 125 mg QID x 10-14 days See CDI treatment algorithm Discontinue antibiotics if possible www.nebraskamed.com/asp
Plesiomonas shigelloides	Severe abdominal cramps, and diarrhea within 6-48 hours	Fresh water, shellfish, international travel	Most patients recover without antimicrobial therapy. Unclear if antibiotics shorten the duration of illness. Consider in severe diarrhea, extremes of age, and immunocompromised.	Fluoroquinolone x 3 days* Azithromycin 500 mg daily x 3 days TMP/SMX DS BID x 3 days
Salmonella	Fever, abdominal cramps, and diarrhea within 6-48 hours, fecal leukocytes often present	Poultry, eggs, dairy products, produce, reptile contact Peak season – summer, fall	Antibiotics have no significant effect on the length of illness and may prolong carriage of the organism in the stool. Antibiotics should generally be avoided but recommended for severe illness (>8 stools/day, high fever, hospitalized) or risk for complications (age <1 or > 50, immunocompromised)	Antibiotics typically not indicated Fluoroquinolone x 7 days* Azithromycin 500 mg daily x 7 days TMP/SMX DS BID x 7 days Immunocompromised patients require 14 days of therapy or longer if relapsing
Yersinia enterocolitica	Fever and abdominal cramps within 1-11 days, with or without diarrhea, fecal leukocytes often present	Unpasteurized milk, undercooked pork, chitterlings Peak season – winter	Most patients recover without antimicrobial therapy. Unclear if antibiotics shorten the duration of illness.	Antibiotics typically not indicated For immunocompromised patients, doxycycline 100 mg IV BID + tobramycin or gentamicin 5 mg/kg/day (TMP/SMX, FQs are alternatives)
Vibrio species (if positive and V. cholera negative indicates V. parahaemolyticus or V. vulnificans is present)	Fever, abdominal cramps, and diarrhea within 6-48 hours, fecal leukocytes often present	Shellfish	Most patients recover without antimicrobial therapy. Unclear if antibiotics shorten the duration of illness. Consider in severe or prolonged diarrhea.	Azithromycin 1 g x 1 dose Doxycycline 300 mg x 1 dose

Vibrio cholerae	Abdominal cramps and large volume watery diarrhea within 16-72 hours	Shellfish, travel to Haiti or other areas where cholera is endemic	Oral rehydration is the key intervention. Antibiotics shorten the duration of illness and are recommended.	Azithromycin 1 g x 1 dose Doxycycline 300 mg x 1 dose Levofloxacin 500 mg x 1 dose Ciprofloxacin 500 mg x 1 dose
Diarrheagenic E. co	bli/Shigella	I		
Enteroaggregative <i>E. coli</i> (EAEC)	Abdominal cramps and watery diarrhea within 16-72 hours, can be	International travel, infantile diarrhea in developing countries	Limited data in EAEC and EPEC. Many patients recover without antimicrobial therapy. Antibiotics have been shown to shorten the	Fluoroquinolone x 3 days* Rifaximin 200 mg TID x 3 days Azithromycin 1 g x 1 dose or 500 mg
Enteropathogenic <i>E. coli</i> (EPEC)	prolonged		duration of illness in ETEC and are generally indicated for moderate to severe diarrhea (>4 stools/day, fever, or blood or pus in stool).	daily x 3 days
Enterotoxigenic <i>E. coli</i> (ETEC) <i>lt/st</i>				
Shiga-like toxin- producing <i>E. coli</i> (STEC) <i>stx1/stx2</i> (shiga-toxin producing <i>E. coli</i> is present) <i>E. coli</i> O157 (the shiga-toxin producing <i>E. coli</i> is type O157)	Bloody diarrhea with minimal fever within 3-8 days	Unpasteurized milk, fresh produce, ground beef, petting zoos	Antibiotics and antimotility agents should be avoided. Antibiotics have no effect on duration or severity of symptoms and certain antibiotics may increase the risk for hemolytic-uremic syndrome.	Supportive care only
<i>Shigella</i> /Enteroinvasive <i>E. coli</i> (EIEC)	Fever, abdominal cramps, and diarrhea within 6-48 hours, fecal leukocytes present	Egg salad, lettuce, day care	Treatment is recommended if detected.	TMP-SMX 160-800 mg BID x 3 days Fluoroquinolone x 3 days* Immunocompromised patients with Shigella require 7-10 days of therapy

\*Levofloxacin 500 mg daily or ciprofloxacin 500 mg BID

# Table 1 (cont.) – Etiology and Treatment Recommendations<sup>8,9,10,11</sup>

Parasites				
Cryptosporidium	Prolonged watery diarrhea	Contaminated water (recreational and drinking), unpasteurized apple cider	Most patients recover without antimicrobial therapy but antibiotics may decrease the duration of illness. Immunocompromised patients often develop prolonged symptoms and respond poorly to therapy.	May use antimotility agents and/or nitazoxanide 500mg BID x 3 days for prolonged or severe illness ID consult recommended for immunocompromised patients
Cyclospora cayetanensis		Imported fresh produce	Treatment indicated if symptomatic.	TMP/SMX DS BID x 7-10 days ID consult recommended for immunocompromised patients
Entamoeba histolytica		Returning travelers	Treatment recommended if detected.	Metronidazole 500 mg TID x 7-10 days OR Tinidazole 2 g daily x 3 days OR Nitazoxanide 500 mg PO BID x 3 days followed by paromomycin 25 mg/kg/day in 3 divided doses x 7 days
Giardia lamblia		Contaminated recreational water, daycare, international travelers	Treatment indicated if symptomatic.	Tinidazole 2 g x 1 dose Nitazoxanide 500 mg PO BID x 3 days Metronidazole 500 mg TID x 5-7 days
Viruses				
Adenovirus F 40/41 Astrovirus Norovirus GI/GII Rotavirus A	Vomiting and non-bloody diarrhea within 10-51 hours	Children <2 yrs, day care Children <1 yr, day care Salads, shellfish, cruise ships, epidemic foodborne disease Peak season – winter Infants	No therapy available. Treat symptomatically.	Antibiotics not indicated
Sapovirus	-	Peak season – winter Children		

Agent	Recommended Dosing	
Azithromycin	10 mg/kg daily	
Ciprofloxacin*	20-30 mg/kg/day in 2 divided doses (max 1.5 g/day)	
Doxycycline*	≥ 8 years: 2-4 mg/kg/day divided every 12-24 hours (max 200 mg/day)	
Levofloxacin*	< 5 years: 8-10 mg/kg/dose twice daily	
	≥ 5 years: 10 mg/kg/dose once daily (max 750 mg/day)	
Metronidazole	Giardiasis: 15 mg/kg/day in divided doses every 8 hours (max 250 mg/dose)	
	C. difficile: 30 mg/kg/day in divided doses every 6 hours (max 2000 mg/day)	
Nitazoxanide	1-3 years: 100 mg every 12 hours	
	4-11 years: 200 mg every 12 hours	
	≥ 12 years: 500 mg every 12 hours	
Paromomycin	25-35 mg/kg/day divided every 8 hours	
Rifaximin	3-11 years: 100 mg four times daily (limited data)	
	≥ 12 years: 200 mg three times daily	
Tinidazole	50 mg/kg single dose or daily (max 2000 mg/day)	
TMP/SMX	≥ 2 months: 8-10 mg/kg/day (TMP component) in divided doses every 12 hours	
Vancomycin (oral)	40 mg/kg/day PO divided every 6-8 hours	

Table 2 – Pediatric Dosing Recommendations

\*Fluoroquinolones and doxycycline are not routinely used as first line therapy in pediatrics

#### **References:**

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