# The Mount Sinai Health System Divisions of Infectious Diseases and Gastroenterology Consensus Guidelines to Diagnosing and Treating Clostridium difficile Infection (CDI)

## Testing:

- Do NOT send stool for *C. difficile* testing without diarrhea (≥ 3 loose bowel movements in 24 hours).
- Do not send stool if there has been recent use of laxatives. Discontinue laxatives and consider testing if diarrhea continues beyond 48 hours and there is no alternative explanation for the diarrhea.
- The Clinical Microbiology Laboratory will reject stool that are not diarrheal (Bristol Stool Chart 1-4).
- Repeat samples are not necessary as the negative predictive value of a single test is >99%.
- Do not send repeat specimens to document "test of cure." The test can remain positive for weeks after treatment. Do not repeat testing in patients who have undergone recent fecal microbiota transplantation (FMT) for the same reason.
- The Clinical Microbiology Laboratory will not accept stool specimens from patients with a negative test within the past 7 days or a positive test within 14 days.
- Downtime forms will only be accepted in the setting of a known EPIC downtime or with ID/Clinical Microbiology Approval.

#### **Treatment:**

STOP ALL ANTIBIOTICS WHEN POSSIBLE

Severity	Clinical Manifestations	Treatment
Asymptomatic	Positive <i>C. difficile</i> test without diarrhea,	No treatment necessary
colonization	ileus, or colitis	TVO treatment necessary
Mild to moderate	Positive <i>C. difficile</i> test with diarrhea	Metronidazole 500 mg PO/NGT every 8 hours for 10
	and no manifestations of severe disease	days
		OR
		Vancomycin 125 mg PO/NGT every 6 hours for 10 days
		<ul> <li>No response to PO metronidazole after 5 days of therapy</li> </ul>
		Intolerance to PO metronidazole
		Pregnant or breastfeeding
		<ul> <li>Underlying inflammatory bowel disease (IBD)</li> </ul>
Severe	Positive C. difficile test with diarrhea	Vancomycin 125 mg* PO/NGT every 6 hours for 10-
	and ≥ 1 of the following attributable to CDI	14 days
	• WBC ≥ 15,000	Consider GI consultation for Fecal Microbiota
	Increase in serum Cr >50% from	Transplantation (FMT) in patients without
	baseline	improvement on 5 days of therapy
Severe	Criteria as above with ≥ 1 of the	Vancomycin 500 mg PO/NGT every 6 hours
Complicated*	following attributable to CDI	AND
	Hypotension	Metronidazole 500 mg IV every 8 hours
	Toxic megacolon	
	• Lactate ≥ 4	If unable to tolerate oral therapy can consider
	ICU admission for severe disease	Vancomycin retention enema (500 mg in 100 mL
		Normal Saline every 6 hours)
		Please consult ID and Surgery
		Consider GI consultation for Fecal Microbiota
		Transplantation (FMT)

- \* There is no evidence for increased doses for oral vancomycin for CDI. Higher doses may be considered in the setting of severe complicated CDI and require ID approval.
- Avoid use of anti-motility agents in patients with CDI.
- Avoid use of binding agents (e.g. cholestyramine) as they can bind oral vancomycin.
- Routine prophylactic use of metronidazole or oral vancomycin is not recommended.

### **Recurrent CDI**

- Resistance to either metronidazole or vancomycin has not been described
- Recurrence occurs in approximately 25% of patients and can be due to failure to eradicate spores or acquisition of a new strain. The risk for recurrence increases with every bout of CDI.

Episode	Treatment
First recurrence	Same regimen as first episode.
Second recurrence	Tapered PO Vancomycin Dose
	125 mg four times a day x 10 days
	125 mg twice a day x 7 days
	125 mg daily x 7 days
	125 mg daily every other day for 7-14 days
	Consider FMT
Third recurrence	Consider FMT

- Avoiding use of anti-motility agents in patients with CDI
- Avoiding use of binding agents (e.g. cholestyramine) as they can bind oral vancomycin
- Routine prophylactic use of metronidazole or oral vancomycin is not recommended

## References

Cohen SH, Gerding DN, Johnson S *et al.* Clinical Practice Guidelines for *Clostridium difficile* Infection in Adults: 2010 Update by the Society for Healthcare Epidemiology of America (SHEA) and the Infectious Diseases Society of America (IDSA). *Infect Control Hosp Epidemiol*. 2010: 31(5); 431-55.

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