

Figure 2: Nitrofurantoin utilization before and after suppression

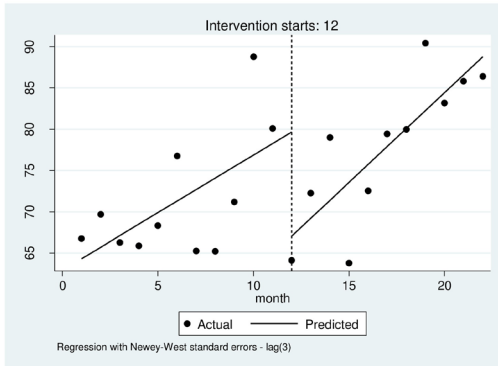


Figure 4: Ceftriaxone utilization before and after suppression

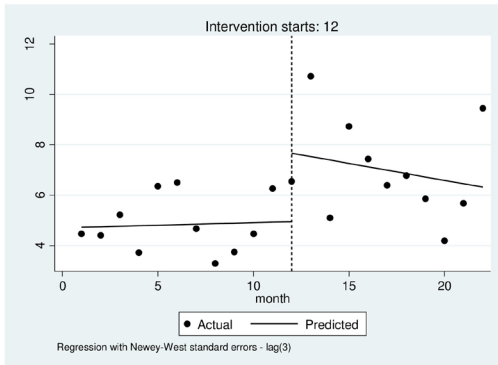


Figure 3: Cephalixin utilization before and after suppression

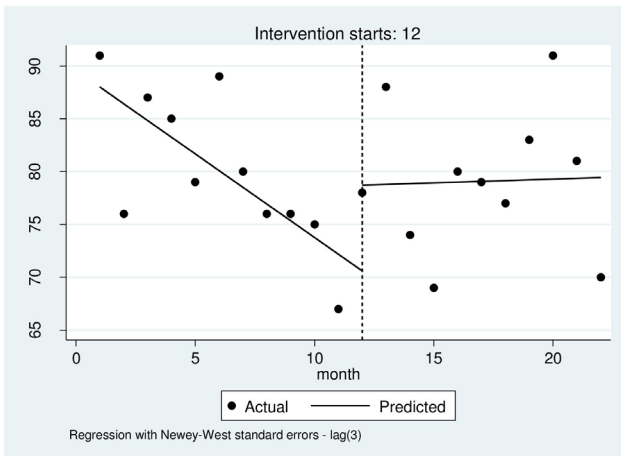


Figure 1: *P. aeruginosa* susceptibility to ciprofloxacin before and after suppression

Disclosures. All authors: No reported disclosures.

1986. Impact of Two-Step Testing on the Diagnosis and Management of *Clostridium difficile* in a Multi-Hospital Healthcare System

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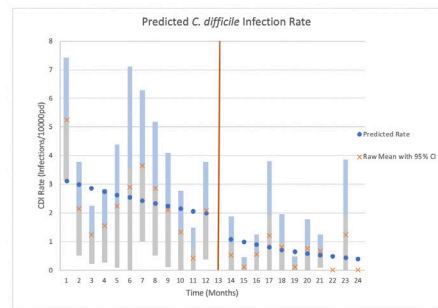
Background. Distinguishing active *C. difficile* infection (CDI) from asymptomatic colonization remains a significant challenge. A multi-step testing algorithm can improve the diagnostic accuracy of CDI and associated antibacterial prescribing. This study evaluated the impact of two-step testing on CDI rates and management in a multi-hospital community health system.

Methods. Two-step *C. difficile* testing (PCR for initial screening followed by EIA for toxin detection) was implemented in 6 acute care community hospitals in April 2018. EIA testing was automatically performed on all stool samples with a positive *C. difficile* PCR result. Prior to implementation, PCR alone was used to identify CDI. Messaging attached to the PCR laboratory report alerted prescribers of discrepant results (PCR +/EIA -). Anti-*C. difficile* therapy was at the discretion of the prescriber. We performed a retrospective cohort analysis over a 2-year period to evaluate the effect of two-step testing on system-wide hospital-onset CDI (HO-CDI) per 10,000 patient-days (PD) and anti-CDI antimicrobial use (AU) in days of therapy (DOT) per 1,000 PD. Segmented negative binomial regression with hospital clustering was used to estimate predicted HO-CDI rate for the baseline period between April 1, 2017 through March 31, 2018 and the post-intervention between May 1, 2018 through March 31, 2019. The implementation date at all sites in April 2018 was unknown; therefore, this month was removed from the analysis. Anti-CDI agents included fidaxomicin, metronidazole, and oral vancomycin, but may have included non-CDI indications for metronidazole.

Results. A total of 115 HO-CDI cases were identified; 91 (79%) before and 24 (21%) after. Prior to implementation of two-step testing, CDI rates declined at 4% per month ($P = NS$). The rate immediately dropped by 36% ($P = 0.004$) after two-step testing was implemented, but the trend did not significantly change ($P = 0.52$, Figure 1). Community-onset CDI rates also decreased during this time period. Combined facility-wide anti-CDI agent use was 824.87 before and 838.21 DOT/1,000 PD after and did not significantly change.

Conclusion. Use of a two-step approach for CDI testing reduced HO-CDI rates, but did not have a significant impact on anti-CDI antibiotic use in a multi-hospital community health system.

Figure 1:



LABEL	MEAN ESTIMATE	CONFIDENCE LIMITS	P VALUE
TREND BEFORE INTERVENTION	0.096	0.8750 1.0522	0.3832
IMMEDIATE CHANGE IN APRIL 2018	0.6366	0.4673 0.8646	0.0039
CHANGE IN TREND AFTER INTERVENTION	0.9398	0.7763 1.1378	0.5245
TREND AFTER INTERVENTION	0.9023	0.8155 0.9963	0.0463

Table 1

Year	Month	HO-CDI	Rate	95% CI	P-value
2017	1	1	1.0	0.1-10.0	0.999
2017	2	1	1.0	0.1-10.0	0.999
2017	3	1	1.0	0.1-10.0	0.999
2017	4	1	1.0	0.1-10.0	0.999
2017	5	1	1.0	0.1-10.0	0.999
2017	6	1	1.0	0.1-10.0	0.999
2017	7	1	1.0	0.1-10.0	0.999
2017	8	1	1.0	0.1-10.0	0.999
2017	9	1	1.0	0.1-10.0	0.999
2017	10	1	1.0	0.1-10.0	0.999
2017	11	1	1.0	0.1-10.0	0.999
2017	12	1	1.0	0.1-10.0	0.999
2018	1	1	1.0	0.1-10.0	0.999
2018	2	1	1.0	0.1-10.0	0.999
2018	3	1	1.0	0.1-10.0	0.999
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2018	5	1	1.0	0.1-10.0	0.999
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2020	9	1	1.0	0.1-10.0	0.999
2020	10	1	1.0	0.1-10.0	0.999
2020	11	1	1.0	0.1-10.0	0.999
2020	12	1	1.0	0.1-10.0	0.999
Total		115	1.0	0.1-10.0	0.999

Figure 2.

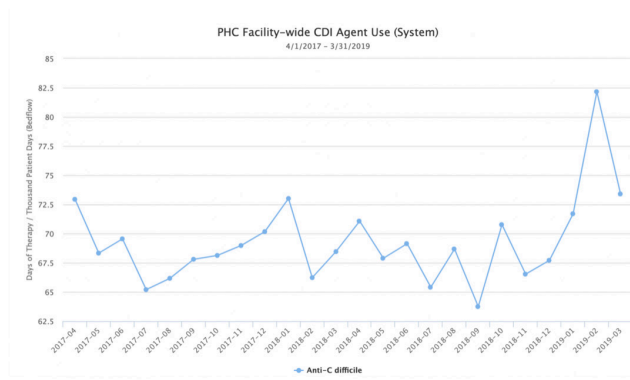


Table 2. Monthly Anti-C. difficile Agent Use (System):

Date	Days of Therapy / Thousand Patient Days (Bedflow) - Anti-C difficile	Fidaxomicin	Metronidazole	Vancomycin (Digestive Tract)
Apr-17	72.94	0.7	62.57	9.67
May-17	68.33	0.6	72.94	9.33
Jun-17	69.55	0.48	57.59	11.49
Jul-17	65.2	0.78	56.66	7.77
Aug-17	66.17	0.33	54.87	10.97
Sep-17	67.8	0.49	55.92	11.38
Oct-17	68.13	0.37	57.76	10
Nov-17	68.98	0.26	58.50	10.22
Dec-17	70.17	0.1	58.46	11.61
Jan-18	73.01	0	60.82	12.18
Feb-18	66.23	0	54.77	11.46
Mar-18	68.46	0	56.63	11.83
Apr-18	71.07	0.52	57.52	13.03
May-18	67.89	0.04	53.39	14.47
Jun-18	69.14	0.44	52.43	16.27
Jul-18	65.41	0.72	53.93	10.76
Aug-18	68.68	0.07	55.82	12.79
Sep-18	63.75	0.22	52.09	11.44
Oct-18	70.77	0.4	55.80	14.57
Nov-18	66.53	0.15	54.91	11.48
Dec-18	67.71	0.23	54.72	12.75
Jan-19	71.7	0.13	57.65	13.92
Feb-19	82.16	0	66.83	15.33
Mar-19	73.4	0.47	58.69	14.24

Disclosures. All authors: No reported disclosures.

1987. Impact of Updated IDSA Clostridium difficile Guidelines on the use of Fidaxomicin in a Large Health System

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Sesson: 234. Antibiotic Stewardship: C. difficile

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Background. Fidaxomicin (fidax) is approved for treatment of *Clostridioides difficile* infection. In February 2018 IDSA/SHEA released updated guidelines suggesting expanded use of fidax, recommending it or oral vancomycin (po vanc) in severe or non-severe initial episodes or for most recurrences. In April 2018, University of Pittsburgh Medical Center (UPMC) relaxed system-wide guidelines to allow for fidax use in the first recurrence of *C. difficile* or later, with earlier use allowed by ID or GI specialists or with local Pharmacy and Therapeutics Chair approval. Hospitals could continue to be more restrictive if desired. We reviewed changes in fidax, po vanc, and IV/PO metronidazole (metro) use at UPMC hospitals after guideline changes.

Methods. For the reviewed antibiotics, hospital-level usage was evaluated at 15 UPMC hospitals before/after system-level changes. Usage was measured as days of therapy per 1,000 patient-days (DOT/1,000 PD). Sites were further grouped by the level of restrictions: Standard (following new system guidelines) or more restrictive (additional restrictions remained in place locally). Hospitals were also grouped by type of local stewardship programs (ASP): Robust (included an Infectious Diseases trained clinical pharmacist or ID physician with specific time dedicated to antibiotic review) or Non-Robust.

Results. Figure 1 shows before/after changes in usage at all hospitals. Figure 2 shows changes in Standard vs. More Restrictive hospitals, and Figure 3 shows changes in Robust vs. Non-Robust hospitals.

Conclusion. Fidax use remained low, but an increase was seen after the release of the guidelines and relaxation of system restrictions, mainly in hospitals without additional restrictions in place. PO vanc also increased across the system, possibly indicating better adherence to updated guidelines regarding less metro use for *C. difficile*

treatment. Although minimal decrease, if any, was seen with metro itself. This could have been compounded by the recent fluid shortage as well as other common uses for metro. Dissemination of new guidelines to providers should be a key function of ASPs as well as monitoring for changes in usage after implementation of local changes. Further studies are needed to define any differences in practice patterns and clinical outcomes related to changes in guidelines.

Figure 1: Antibiotic Use at 15 UPMC Hospitals

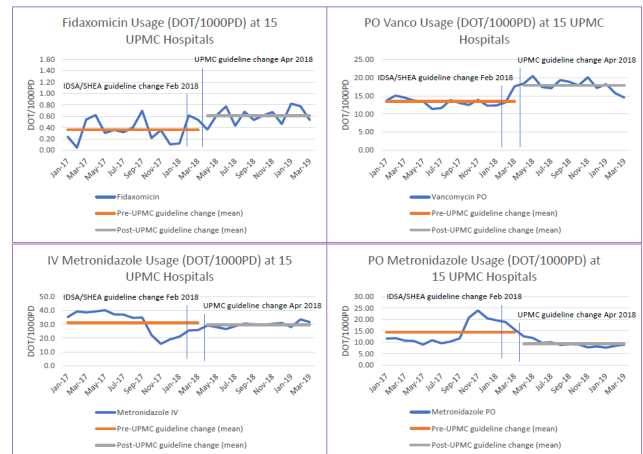


Figure 2: Antibiotic Usage by Level of Local Restriction

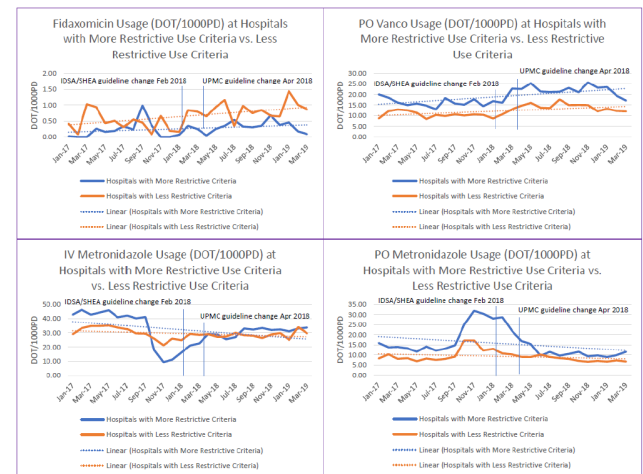
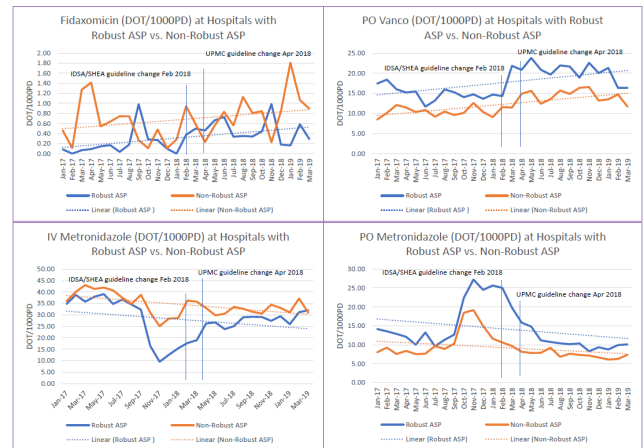


Figure 3: Antibiotic Use Based on Level of Local Stewardship Program



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1988. Impact of a Novel Pharmacist Practice Model on Antimicrobial Usage and Hospital-acquired Clostridium difficile (HACDI) Rates

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