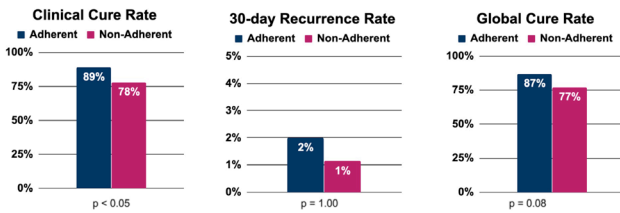


Figure 1. Rates of Clinical Cure, 30-day Recurrence, and Global Cure of *C. difficile* Infection



Disclosures. All authors: No reported disclosures.

1984. A Multi-Disciplinary Team-based Quality Improvement Initiative to Reduce *Clostridioides difficile* Rates and Promote Antimicrobial Stewardship in Targeted Surgical Wards

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Background. At Stanford, two surgical wards, E3 and F3, were responsible for 1/5 of hospital-acquired *Clostridioides difficile* infection (HO CDI) cases in the fiscal year 2018 (FY2018). We used a quality improvement framework with a goal to reduce yearly HO CDI episodes by 1/2 on these wards.

Methods. A multidisciplinary quality improvement team was created with front-line nursing leaders and representatives from colorectal surgery, gynecology oncology, antimicrobial stewardship (ASP), infection prevention, and pharmacy. Coaching and instruction on quality improvement were provided as part of Stanford's "Realizing Improvement through Team Empowerment" (RITE) program. Using A3 problem solving, root cause analysis identified key drivers, and interventions were performed. Cumulative HO CDI cases in FY2019 and weekly antibiotic days of therapy (DOT) on E3/F3 were monitored.

Results. Review of FY2018 HO CDI cases ($n = 14$) revealed the most common key driver as inappropriate antibiotic prescribing (8 cases, 57%). Multiple interventions were instituted (Figure 1). Three ASP interventions began February 2019: nursing questioned antibiotic choice/duration on daily interdisciplinary rounds (Figure 2), automatic infectious disease consultation for > 72 hours of piperacillin/tazobactam on gynecology/oncology patients, and twice-weekly rounds between ASP and a colorectal attending. Data from ASP/colorectal rounds from March 19, 2019 to April 16, 2019 showed means of 18.2 minutes taken for chart review and 4.4 minutes for discussion. 25 charts reviewed led to 16 (64%) ASP recommendations and 14/16 (87.5%) of recommendations accepted. Common interventions included: appropriate duration of antibiotics, clarification of the team's planned duration, and review of microbiology data to narrow therapy. Mean DOT decreased from 35.28 to 21.61 (39%) since July 2018 (Figure 3). Patient volume and case mix index remained stable throughout, suggesting no impact on DOT. Though CDI cases did not decrease, interventions were in place for only 2 months (Figure 4).

Conclusion. While too early to determine its impact on HO CDI rates, a multi-disciplinary team approach utilizing A3 problem solving was successful in implementing effective ASP measures including nursing-led ASP and structured antibiotic timeouts.

Figure 1. Key Drivers and Interventions Targeting HO CDI

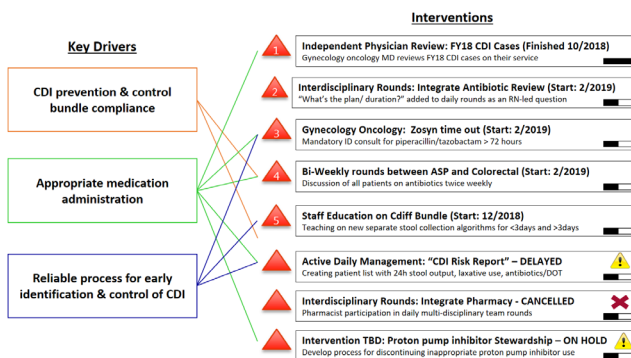


Figure 2. Nursing-led ASP: Integration of "What's the plan/duration" of antibiotics into Interdisciplinary Rounds

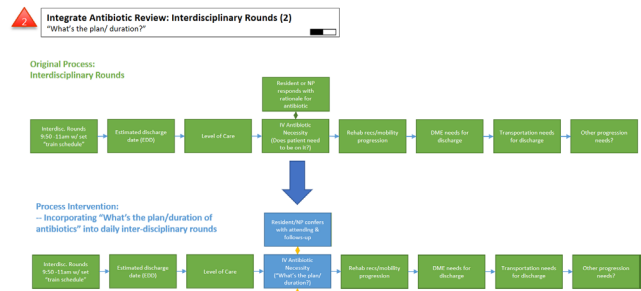


Figure 3. Total Antibiotic DOT Change Over Time: Before and After ASP Interventions

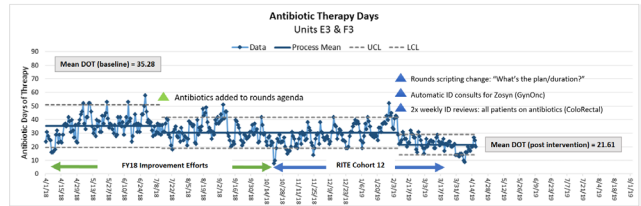
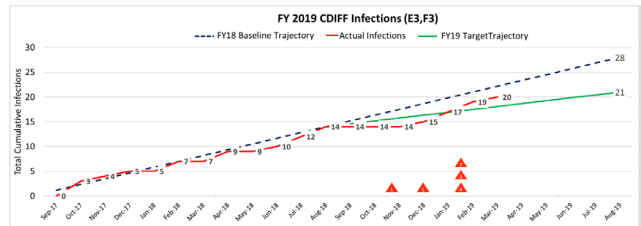


Figure 4. Baseline vs. Goal Trajectory of Cumulative HO CDI Rates on E3/F3 From FY2018 to FY2019



Disclosures. All authors: No reported disclosures.

1985. Impact of Suppressing Ciprofloxacin Susceptibility Results on Antibiotic Utilization and Hospital-acquired *Clostridioides difficile* Infection

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Background. Fluoroquinolones (FQs) are broad-spectrum antibiotics associated with multiple adverse effects and an increased risk of *Clostridioides difficile* infections (CDI). Previous data suggest that suppression of FQ susceptibility results decreased FQ use. The purpose of this study was to examine the impact of suppressing ciprofloxacin susceptibility on antibiotic use, susceptibility, and CDI.

Methods. This was a single-center quasi-experimental study of the effect of the suppression of ciprofloxacin susceptibility on pan susceptible urine isolates for *Klebsiella* sp. and *E. coli* starting in March 2018 in the 11 months before and after the intervention. Monthly antibiotic utilization in days of therapy (DOT)/1,000 patient-days for levofloxacin, ciprofloxacin, ceftriaxone, trimethoprim/sulfamethoxazole (TMP/SMZ), fosfomycin, and nitrofurantoin, hospital-acquired CDI (HA-CDI) rates as defined by CDC, and *Pseudomonas aeruginosa* susceptibility was compared with interrupted time series analysis using Stata MP 12.1 before and after the intervention to compare the level, intercept, and rate, slope, of a trend line.

Results. There was no change in the level or rate of ciprofloxacin DOT (0.27, 95% CI: -0.94 to 1.48-3.49; 95% CI: -10.89 to 3.90) and levofloxacin DOT (-5.87, 95% CI: -17.79 to 6.06; -0.98, 95% CI: -2.86 to 0.90) with the intervention, respectively. Level of *P. aeruginosa* susceptibility to ciprofloxacin level (8.13, 95% CI: 0.00 to 16.26) had a trend toward increasing and rate (1.65, 95% CI: 0.44 to 2.87) increased after the intervention. Ceftriaxone DOT level decreased after the intervention ($P = 0.01$), but the rate did not change. Cephalexin ($P = 0.01$) and nitrofurantoin ($P = 0.01$) DOT levels increased after the intervention without changes in rates. There was no change in the level or rate of HA-CDI, fosfomycin, or TMP/SMZ DOTs.

Conclusion. Suppressing ciprofloxacin susceptibility results on pan susceptible *Klebsiella* sp. and *E. coli* urine isolates was associated with increased *P. aeruginosa* susceptibility to ciprofloxacin and increased cephalexin and nitrofurantoin DOTs. No changes were seen in FQ use or HA-CDI rates.

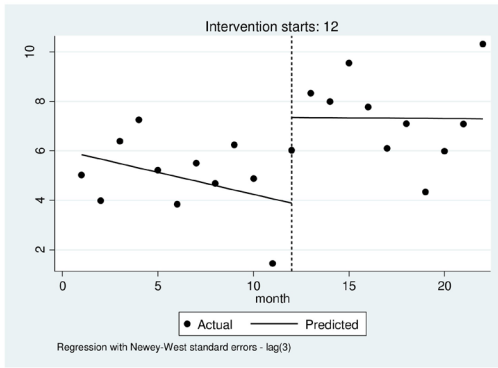


Figure 2: Nitrofurantoin utilization before and after suppression

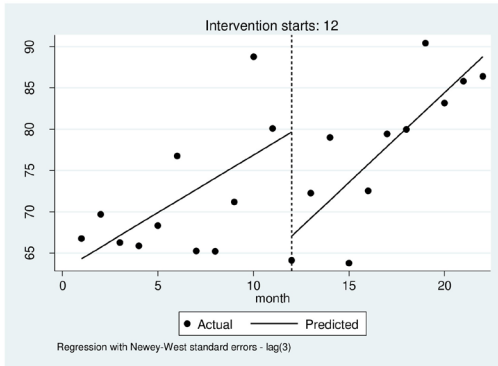


Figure 4: Ceftriaxone utilization before and after suppression

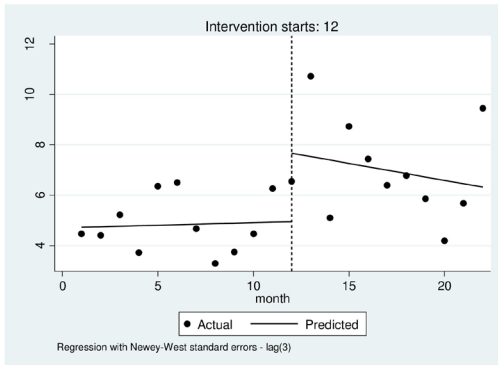


Figure 3: Cephalixin utilization before and after suppression

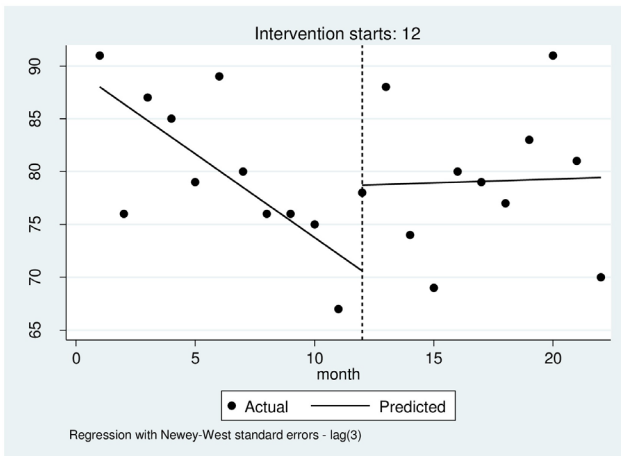


Figure 1: *P. aeruginosa* susceptibility to ciprofloxacin before and after suppression

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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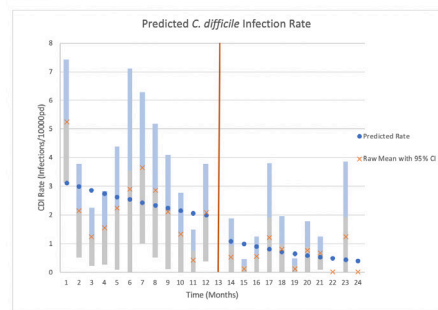
Background. Distinguishing active *C. difficile* infection (CDI) from asymptomatic colonization remains a significant challenge. A multi-step testing algorithm can improve the diagnostic accuracy of CDI and associated antibacterial prescribing. This study evaluated the impact of two-step testing on CDI 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 | CDI Rate | 95% CI | P Value |
|-------|-------|----------|-----------|---------|
| 2017 | 1 | 4.5 | 3.5 - 5.5 | 0.001 |
| 2017 | 2 | 4.2 | 3.2 - 5.2 | 0.002 |
| 2017 | 3 | 4.8 | 3.8 - 5.8 | 0.003 |
| 2017 | 4 | 4.5 | 3.5 - 5.5 | 0.004 |
| 2017 | 5 | 4.2 | 3.2 - 5.2 | 0.005 |
| 2017 | 6 | 4.8 | 3.8 - 5.8 | 0.006 |
| 2017 | 7 | 4.5 | 3.5 - 5.5 | 0.007 |
| 2017 | 8 | 4.2 | 3.2 - 5.2 | 0.008 |
| 2017 | 9 | 4.8 | 3.8 - 5.8 | 0.009 |
| 2017 | 10 | 4.5 | 3.5 - 5.5 | 0.010 |
| 2017 | 11 | 4.2 | 3.2 - 5.2 | 0.011 |
| 2017 | 12 | 4.8 | 3.8 - 5.8 | 0.012 |
| 2018 | 1 | 3.2 | 2.2 - 4.2 | 0.013 |
| 2018 | 2 | 3.5 | 2.5 - 4.5 | 0.014 |
| 2018 | 3 | 3.8 | 2.8 - 4.8 | 0.015 |
| 2018 | 4 | 3.5 | 2.5 - 4.5 | 0.016 |
| 2018 | 5 | 3.2 | 2.2 - 4.2 | 0.017 |
| 2018 | 6 | 3.5 | 2.5 - 4.5 | 0.018 |
| 2018 | 7 | 3.8 | 2.8 - 4.8 | 0.019 |
| 2018 | 8 | 3.5 | 2.5 - 4.5 | 0.020 |
| 2018 | 9 | 3.2 | 2.2 - 4.2 | 0.021 |
| 2018 | 10 | 3.5 | 2.5 - 4.5 | 0.022 |
| 2018 | 11 | 3.8 | 2.8 - 4.8 | 0.023 |
| 2018 | 12 | 3.5 | 2.5 - 4.5 | 0.024 |
| 2019 | 1 | 3.2 | 2.2 - 4.2 | 0.025 |
| 2019 | 2 | 3.5 | 2.5 - 4.5 | 0.026 |
| 2019 | 3 | 3.8 | 2.8 - 4.8 | 0.027 |
| 2019 | 4 | 3.5 | 2.5 - 4.5 | 0.028 |
| 2019 | 5 | 3.2 | 2.2 - 4.2 | 0.029 |
| 2019 | 6 | 3.5 | 2.5 - 4.5 | 0.030 |
| 2019 | 7 | 3.8 | 2.8 - 4.8 | 0.031 |
| 2019 | 8 | 3.5 | 2.5 - 4.5 | 0.032 |
| 2019 | 9 | 3.2 | 2.2 - 4.2 | 0.033 |
| 2019 | 10 | 3.5 | 2.5 - 4.5 | 0.034 |
| 2019 | 11 | 3.8 | 2.8 - 4.8 | 0.035 |
| 2019 | 12 | 3.5 | 2.5 - 4.5 | 0.036 |
| 2020 | 1 | 3.2 | 2.2 - 4.2 | 0.037 |
| 2020 | 2 | 3.5 | 2.5 - 4.5 | 0.038 |
| 2020 | 3 | 3.8 | 2.8 - 4.8 | 0.039 |
| 2020 | 4 | 3.5 | 2.5 - 4.5 | 0.040 |
| 2020 | 5 | 3.2 | 2.2 - 4.2 | 0.041 |
| 2020 | 6 | 3.5 | 2.5 - 4.5 | 0.042 |
| 2020 | 7 | 3.8 | 2.8 - 4.8 | 0.043 |
| 2020 | 8 | 3.5 | 2.5 - 4.5 | 0.044 |
| 2020 | 9 | 3.2 | 2.2 - 4.2 | 0.045 |
| 2020 | 10 | 3.5 | 2.5 - 4.5 | 0.046 |
| 2020 | 11 | 3.8 | 2.8 - 4.8 | 0.047 |
| 2020 | 12 | 3.5 | 2.5 - 4.5 | 0.048 |
| Total | | 3.5 | 2.5 - 4.5 | 0.049 |