#### UNIVERSITY of HOUSTON COLLEGE OF PHARMACY

## P2402

## BACKGROUND

- PCR ribotyping of *Clostridioides difficile* strains is commonly used to describe the epidemiology of C. difficile infection.
- Certain ribotypes (RT) have been associated with more severe disease and clinical outcomes, such as RT 027,<sup>1</sup> while others are considered less virulent, such as RT 014-020.<sup>2</sup>
- Texas statewide surveillance identified the emergence of a rarely-described RT 255 beginning in 2015, which now represents the fifth most common ribotype in the state.<sup>3</sup>
- Here we describe clinical outcomes associated with an emergent RT 255 in Texas.

### METHODS

#### **Study design / inclusion**

- A retrospective, cohort study was conducted including patients from two tertiary care centers in Houston, Texas.
- A convenience sample of patients infected with *C*. *difficile* strains of either RT 255, 014-020, or 027 between 2016-18 were included.
- Patients demographics and outcomes were collected by trained abstractors blinded to the RT results. The primary outcome was disease severity as classified by the 2017 IDSA guidelines.

#### **Culture and ribotyping**

- C. difficile stool was plated onto cefoxitincycloserine-fructose agar plates and anaerobically incubated for 48–72 hours.
- Fluorescent ribotyping was performed as previously described.<sup>2</sup>
- This technique does not distinguish between all RTs; therefore, ribotypes 014 and 020 are reported as combined (014-020).

#### Statistical analysis

- Multivariable logistic regression analysis was done to control for other patient characteristics.
- Results were significant at P < 0.05, and all</p> statistical analyses were completed using SPSS, version 25.

## **Clinical Outcomes Associated with an Emerging Clostridioides difficile** Ribotype 255 in Texas

Gonzales-Luna AJ<sup>1</sup>, Carlson TJ<sup>1§</sup>, Lancaster C<sup>1</sup>, Miranda J<sup>1</sup>, Garey KW<sup>1</sup> <sup>1</sup>The University of Houston College of Pharmacy, Houston, TX, <sup>§</sup>Current affiliation: High Point University Fred Wilson School of Pharmacy, High Point, NC

	Ribotype				
	<b>255</b> n = 50	<b>027</b> n = 50	255 vs. 027 p-value	<b>014-020</b> n = 50	255 vs. 014-020 p-value
Demographics					
Age, mean years (±SD)	59.3 (±16.8)	69.3 (±13.6)	0.001	61 (±18.3)	0.63
CCI score, median (IQR)	2 (1-3)	3 (2-5)	0.014	2 (1-4)	0.34
Outcomes					
Initial clinical cure, no. (%)	38 (76)	30 (60)	0.09	38 (76)	Not tested
Severe/fulminant disease, no. (%)	19 (38)	40 (80)	< 0.001	20 (40)	0.83
CDI complications <sup>†</sup>	5 (10)	14 (28)	0.02	9 (18)	0.25
30d recurrence, no. (%)	2 (4)	2 (4)	Not tested	3 (6)	0.64
90d recurrence‡, no. (%)	5 (10)	10 (20)	0.17	4 (8)	0.70
All-cause 30d mortality, no (%)	5 (6)	8 (16)	0.37	6 (12)	0.75

- A total of 150 patients were included (50 patients infected with each RT). Overall, 53% of the patients had severe or fulminant disease most commonly due to RT 027 (80%) followed by RT 014-020 (40%) and RT 255 (38%).
- *C. difficile* disease severity of was similar between those infected with RT 255 and RT 014-020 (p = 0.84).



## REFERENCES

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- Aitken SL, Alam MJ, Khaleduzzaman M, et al. In the Endemic Setting, *Clostridium difficile* Ribotype 027 Is Virulent But Not Hypervirulent. Infect Control Hosp Epidemiol. 2015; 36(11): 1318-23. Gonzales-Luna AJ, Carlson TJ, Dotson KM, et al. Texas statewide surveillance of *Clostridioides difficile* infection, 2011-2018. P0275. 29th ECCMID, Amsterdam, Netherlands. April 13-16, 2019.

## DECINTC

- 95% Cl, 0.037-0.433; p = 0.001).
- recurrence between the three ribotypes.

## CONCLUSION

# •Ribotype 255 does not appear to be associated with more severe disease when compared to

•Further studies are warranted to determine contributing factors for its increasing prevalence.

**Contact Information:** Anne Gonzales-Luna University of Houston Phone: (713) 398-9051 Email: ajgonz23@central.uh.edu

> Table 1. Baseline
> characteristics and outcomes associated with an emergent ribotype 255 compared to two other endemic ribotypes in Houston, Texas; †includes ICU admission, colectomy, ileus, and toxic megacolon, ‡90-day recurrence includes those with 30-day recurrence. Abbreviations: Charlson Comorbidity Index (CCI)

In multivariable analysis, patients infected with ribotype 255 had an 87% relative reduction in the odds of severe disease compared to ribotype 027 after controlling for patient age, CCI score, and serum albumin level (OR, 0.13;

No differences were seen in the rates of 30-day mortality, or 30- or 90-day



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