

A Pharmacoepidemiologic Evaluation of Echinocandin Use

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BACKGROUND

- Invasive candidiasis (IC) is a devastating fungal infection and candidemia is the most common bloodstream infection with high attributable mortality rates of 30-40% in the US hospitals^{1,2}
- Rates of IC caused by drug-resistant Candida spp, designated by the CDC as a serious threat, are increasing, and Candida auris has become an urgent threat³
- Currently three available classes of systemic antifungals are echinocandin-, azole-, and amphotericin-based therapies⁴
- Comparatively, echinocandins demonstrate low minimum inhibitory concentration (MICs) against most Candida species and favorable toxicity⁴

OBJECTIVES

- 1. To perform a pharmacoepidemiologic analysis on echinocandin use at a quaternary care medical center
- 2. To review duration of therapy of echinocandins for positive Candida cultures and days to therapy initiation during hospitalization
- assess echinocandin disposition upon discharge after hospitalization

METHODS

- Echinocandin use and clinical microbiologic data between 2017 and 2019 were pooled via Theradoc
- Monthly days of therapy (DOT) per 1,000 patient days were calculated
- The proportion of echinocandin-treated patients with or without positive Candida cultures was evaluated along with echinocandin use, and hospital admission and discharge dates was also evaluated
- A subgroup analysis of the first 50 included patients was performed to evaluate echinocandin discharge disposition
- R statistical analysis (ggplot2) was used to generate visual data

CONCLUSION

- Overall, echinocandin use did not change appreciably
- Initiation of echinocandin occurred throughout the entire hospitalization time period
- A significant portion of echinocandin courses continued after hospital discharge
- Further studies evaluating potential benefits of long-acting echinocandin with an emphasis of transition of care are warranted

FUNDING

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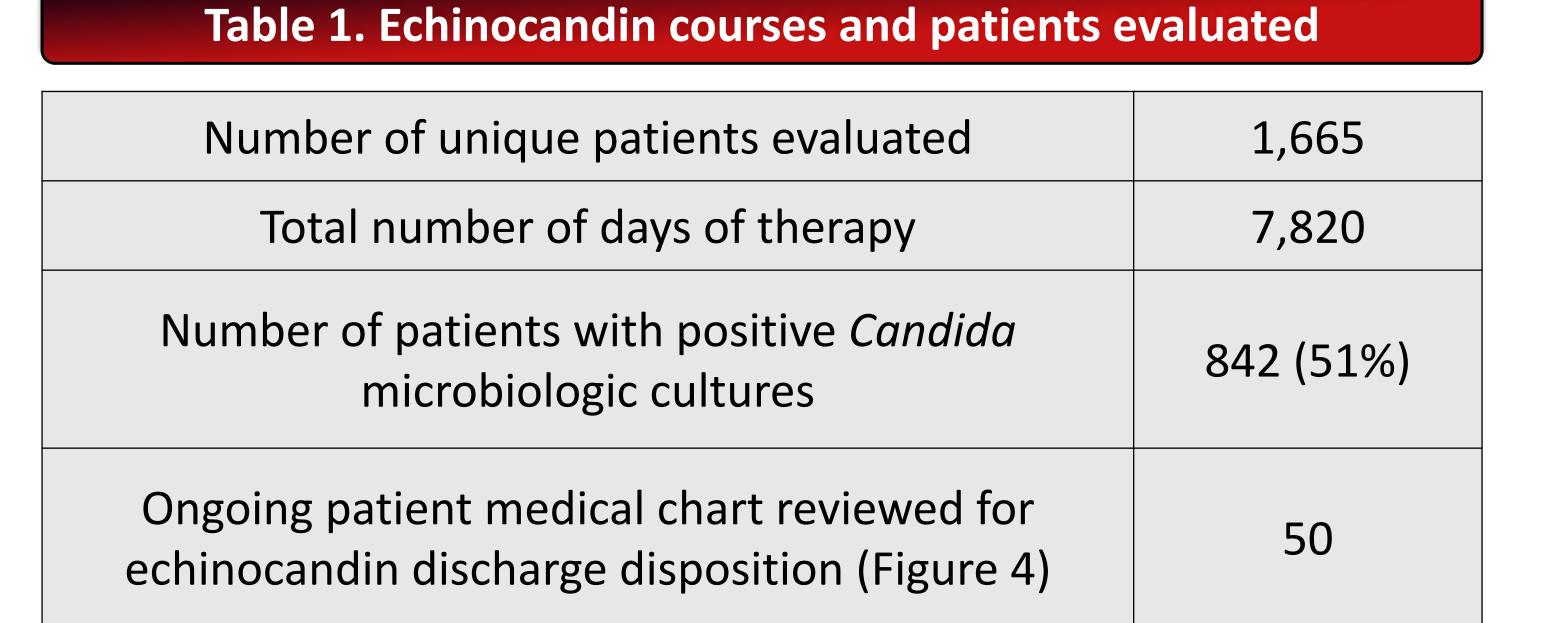


Figure 1. Echinocandin DOT per 1,000 patient days (2017 -2019)

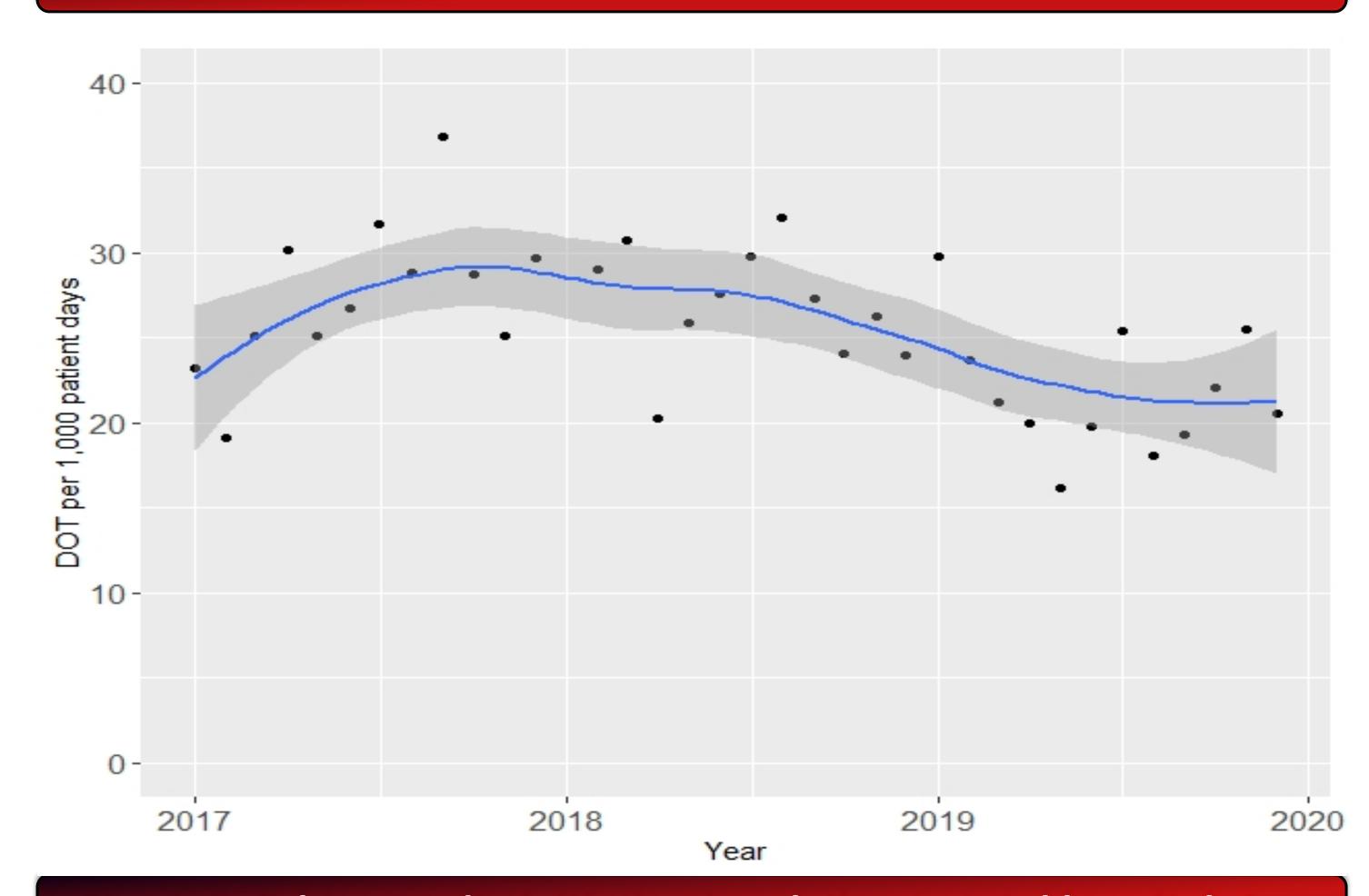
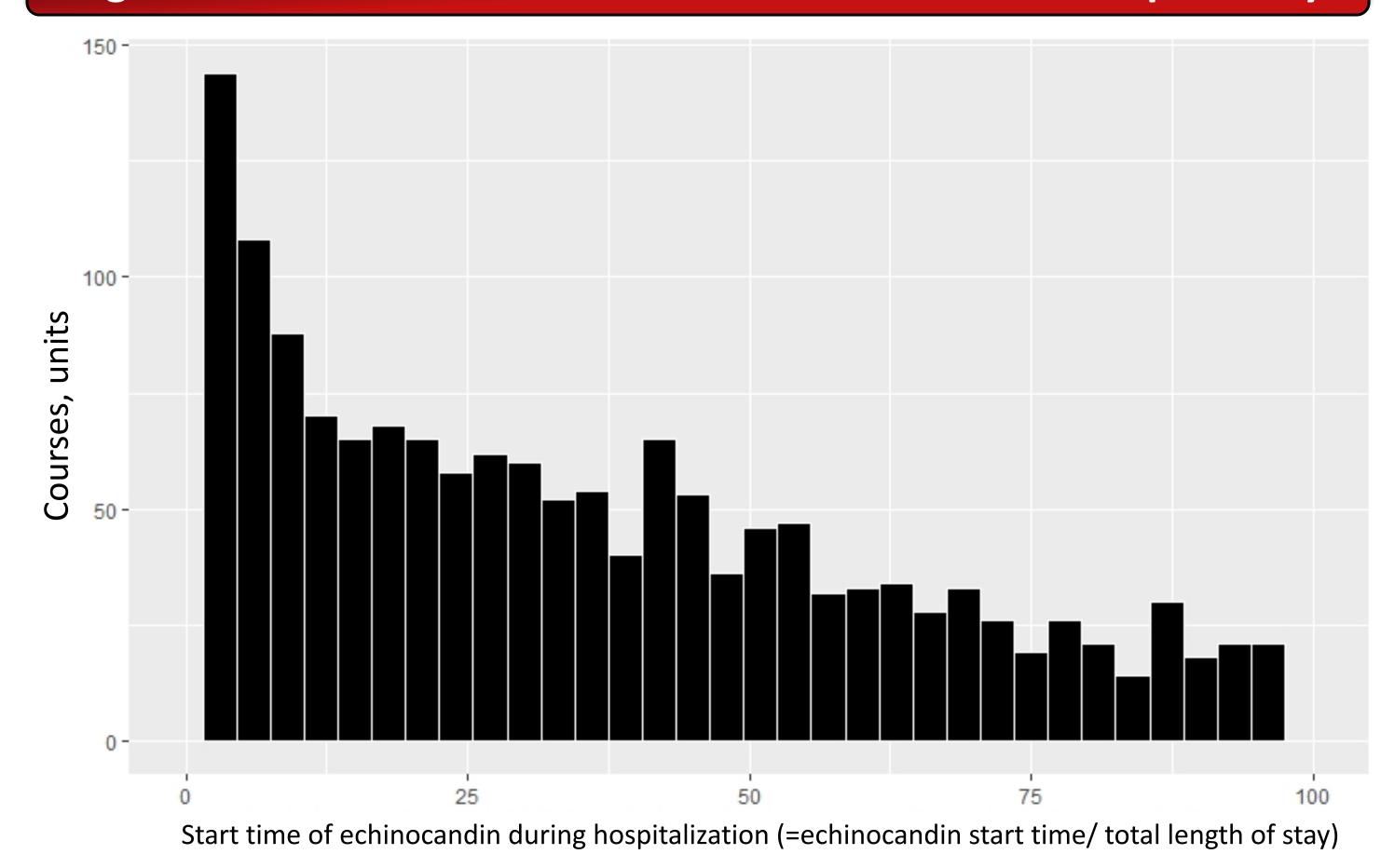


Figure 2. Echinocandin initiation in relation to total hospital stay



RESULTS

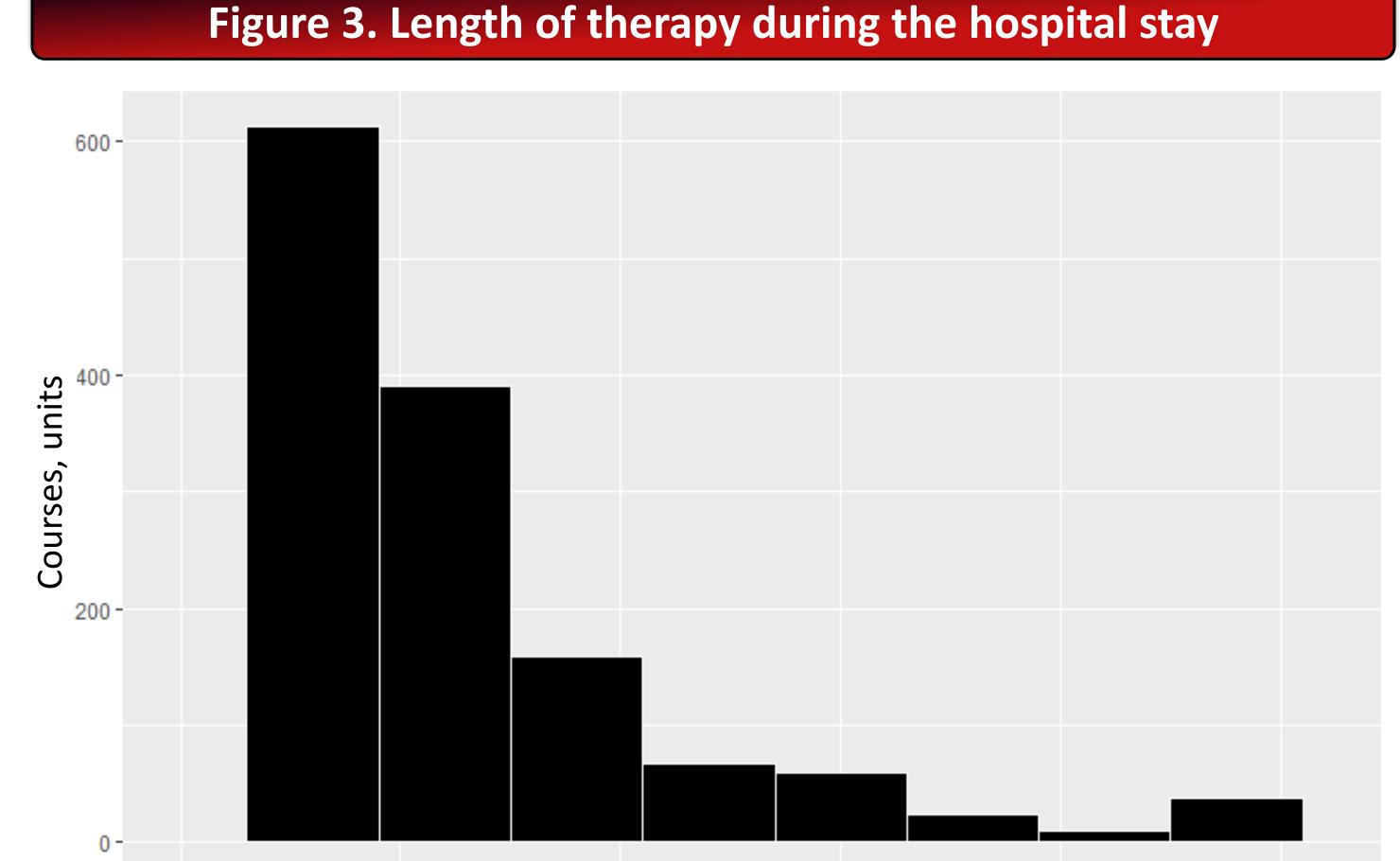
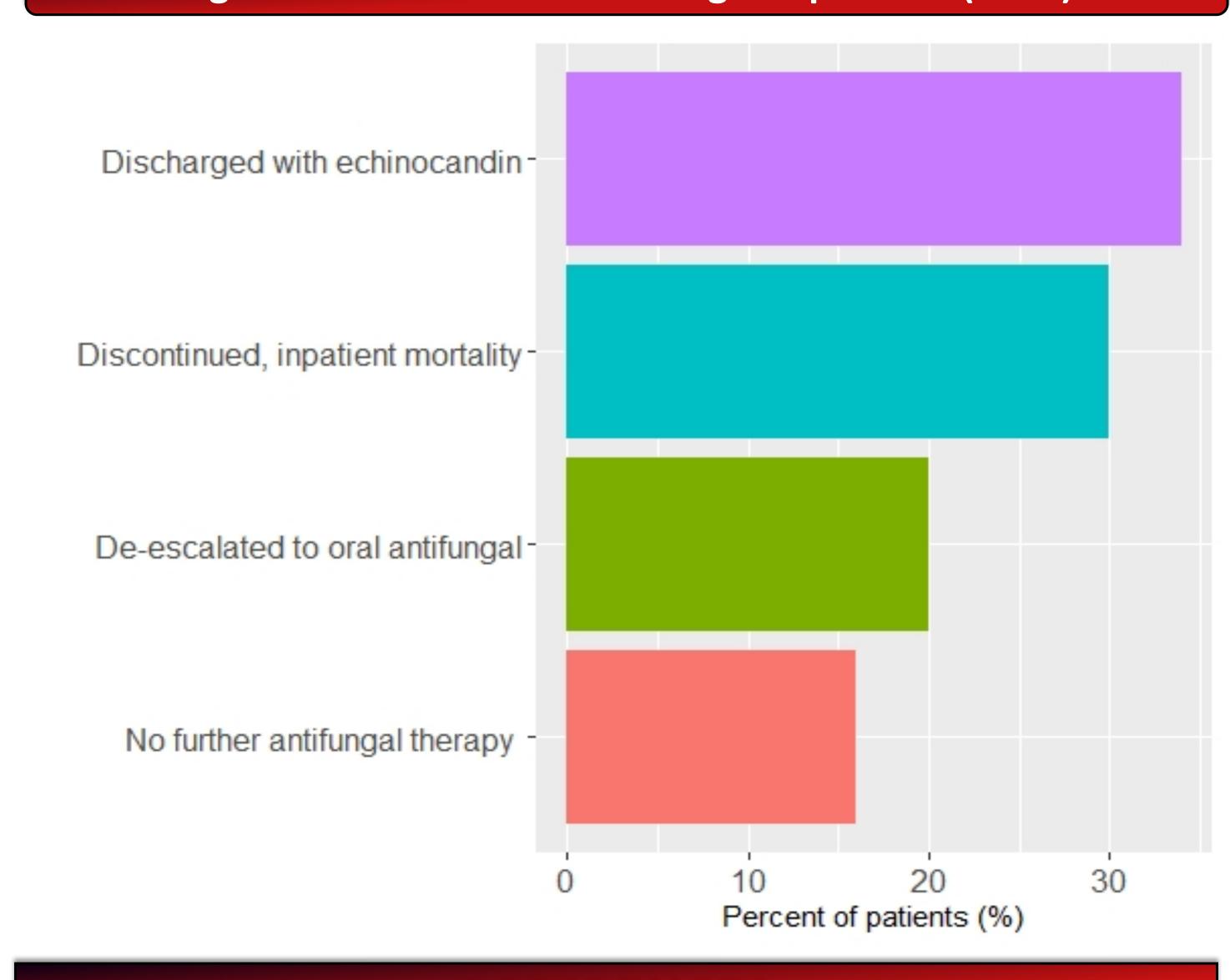


Figure 4. Echinocandin discharge disposition (n=50)

Length of echinocandin therapy, days



REFERENCES

- . Guery BP, Arendrup MC, Auzinger G, et al. Management of invasive candidiasis and candidemia in adult nonneutropenic intensive care unit patients: Part I. Epidemiology and diagnosis. Intensive Care Med. 2009;35:55-62
- 2. Strollo S, Lionakis MS, Adjemian J, et al. Epidemiology of hospitalizations associated with invasive candidiasis, United States, 2002-2012. *Emerging Infect Dis*. 2017;23:7-13
- 3. Centers for Disease Control and Prevention. Antibiotic resistance threats in the United States, 2019. Atlanta, GA
- 4. Pappas PG, Kauffman CA, Andes DR, et al. Clinical practice guideline for the management of candidiasis: 2016 update by the Infectious Diseases Society of America. Clin Infect Dis. 2016;62(4)e1-50