

Modeling Likelihood of Coverage for Narrow Spectrum Antibiotics in Patients Hospitalized with Urinary Tract Infections

Authors: Courtney Hebert, MD¹, Erinn M. Hade, PhD¹, Protiva Rahman¹, Yuan Gao¹, Mark Lustberg, MD, PhD¹, and Kurt Stevenson, MD¹, Preeti Pancholi, PhD¹
 Institutions: Department of Biomedical Informatics, Department of Internal Medicine, The Ohio State University

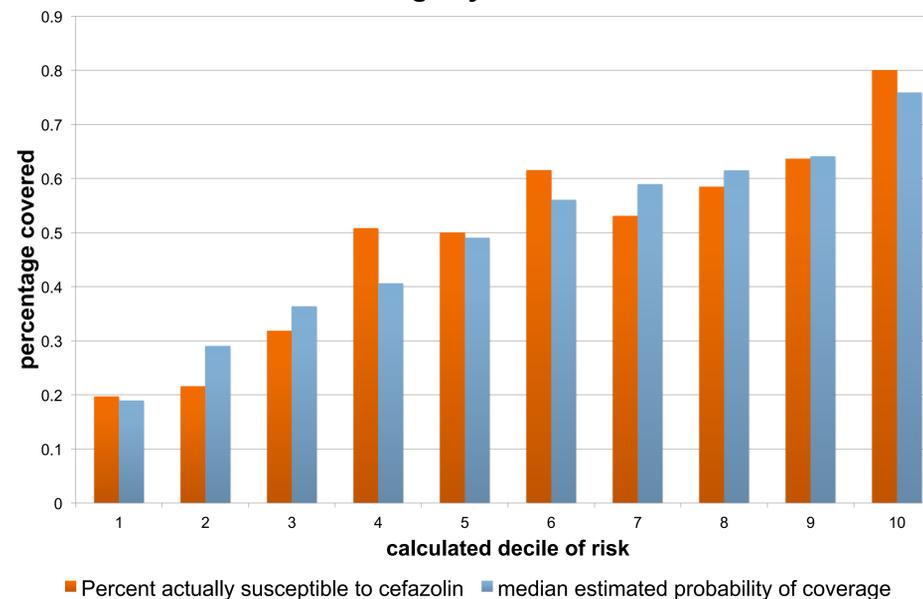
Background

- When prescribing empiric antibiotics, providers try to choose the narrowest spectrum antibiotic that will cover a patient's infection.
- To do this they must assess the likelihood of coverage of different regimens.
- As a test of feasibility, we developed a predictive model for cefazolin/cephalexin coverage for patients admitted to the hospital with urinary tract infections (UTI) to identify a group of patients with a high likelihood of coverage by this first-line, narrow spectrum antibiotic.
- We also investigated how cefazolin/cephalexin treatment would compare to the actual empiric antibiotic treatment given in terms of breadth of coverage and number of antimicrobials given

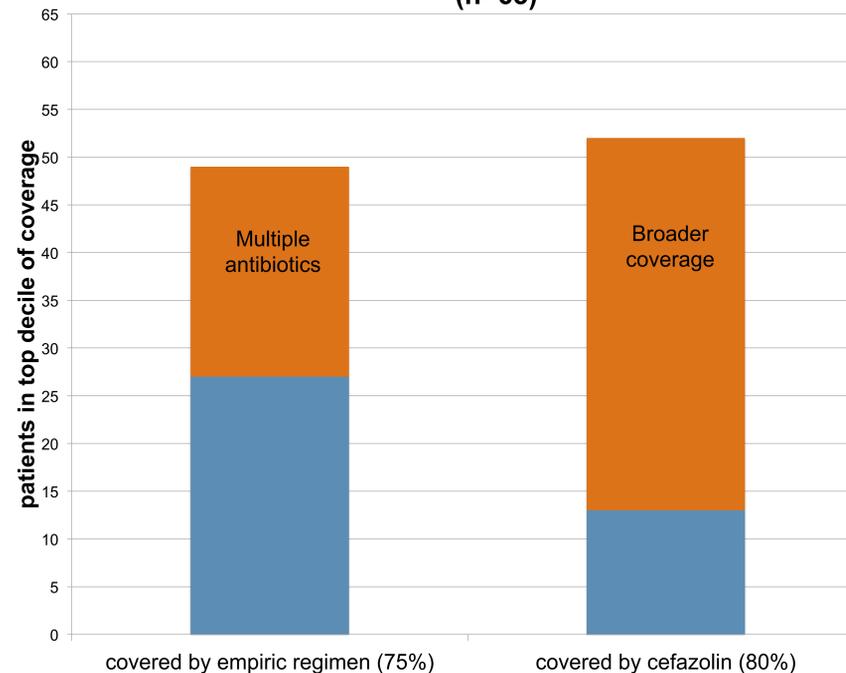
Methods

- Patients admitted from 11/1/11 to 1/1/14 with a positive urine culture in the first 48 hours and a discharge diagnosis of UTI were included.
- Data extracted from our information warehouse included empiric antibiotic administration data, demographics, comorbidities, and past antibiotic use.
- Only the first eligible admission for each patient was included.
- A 20% random sample of patients was selected as the test/validation set.
- Logistic regression models estimated the predicted probability of cefazolin coverage
- Models were built on the training set using two methods
 - Stepwise variable selection
 - 10 fold cross validation with LASSO penalized logistic regression, with and without interactions included

Comparison of actual % covered to estimated probability of coverage by decile of risk

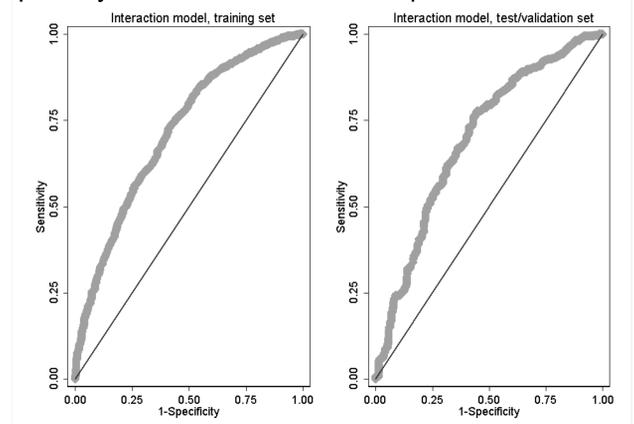


Patients in top decile of probability of cefazolin coverage (n=65)



Results

- 3,456 patients with an eligible UTI were included. 691 were held out for validation. 49% of the UTIs were covered by cefazolin.
- The final stepwise model had an area under the receiver operating curve (AUC) of 69% (95% CI: 67%, 71%) in the training and 70%, (66%, 74%) in the test/validation set.
- The final penalized model (with interaction effects included) had an AUC 72% (70%, 74%) in the training and 70% (66%, 74%) in the test/validation set.
- 80% (52/65) in the highest estimated decile of cefazolin coverage had a UTI that would have been covered by cefazolin; only 13/66 (20%) in the lowest decile would have been covered by cefazolin.
- 49/65 (75%) of patients in the highest decile of cefazolin coverage were covered by the actual empiric regimen given, however 22/49 (45%) of those regimens consisted of multiple antibiotics.
- Of those that would have been covered by cefazolin 39/52(75%) were empirically treated with a broader spectrum antibiotic.



Conclusion

- This preliminary model can reasonably identify patients whose infections would be likely to be covered by cefazolin.
- If this model were used to guide clinical decision making, our analysis suggests that a majority of patients would have been covered by more narrow spectrum antibiotics than what they received.

Research reported in this publication was supported by the National Institute Of Allergy And Infectious Diseases of the NIH under Award Number R01AI116975. The project described was supported by Award Number Grant UL1TR001070 from the National Center For Advancing Translational Sciences. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Center For Advancing Translational Sciences or the National Institutes of Health