SOUTHERN ILLINOIS UNIVERSITY EDWARDSVILLE SCHOOL OF PHARMACY

### Introduction

- Proton pump inhibitors (PPIs) and hista receptor-2 antagonists (H2RAs) are wid used by patients in both the institutiona and in the community for many indication
- One such indication is reducing the risk ulcers in high risk patients, such as those taking oral corticosteroids.
- Several studies have determined PPIs inappropriately prescribed in a large nu patients.
- There is limited data on readmission ration patients who are prescribed these med

## Objective

 To determine if there is a difference in 3 readmission rates in patients on acid re therapy (PPI or H2RA) alone or in com with oral corticosteroids.

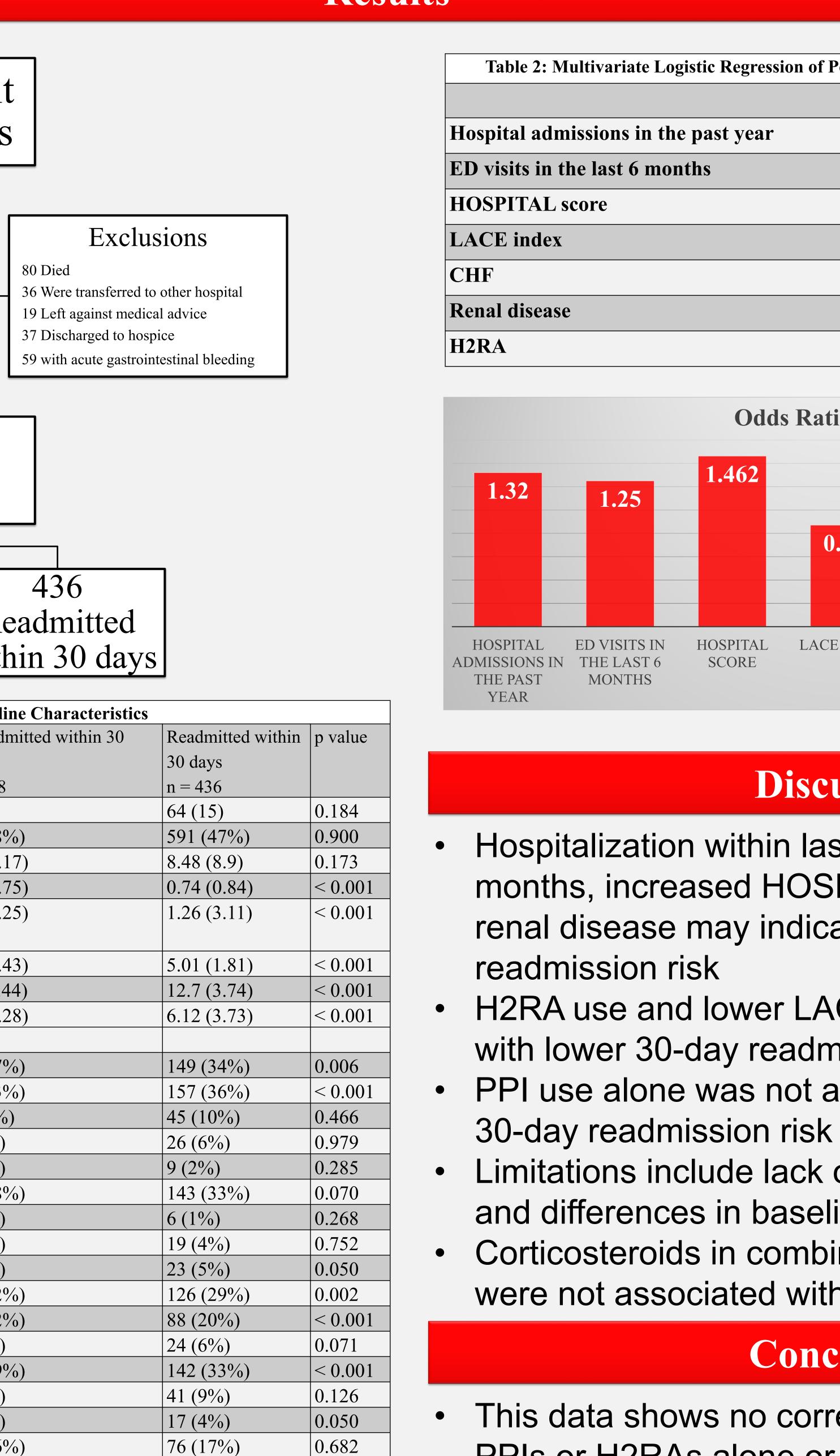
## Methods

- Retrospective chart review approved by Institutional Review Board at a 500-bec teaching hospital in Springfield, Illinois
- Inclusion Criteria:
  - Adults 18 years old or older who wer discharged on a PPI or H2RA in com with a corticosteroid as well as patien discharged with a corticosteroid alon
- Data Collected:
  - Patient demographics
  - Comorbid conditions (Prior MI, CHF,
  - Discharge medications
  - Hospital admissions within last year
  - ED visits within last 6 months
- Data Analysis: Descriptive Statistics and multivariate logistic regression

Effects of Proton Pump Inhibitors vs Histamine Type-2 **Receptor Antagonists in Combination with Oral Corticosteroids on Hospital Readmission Rates** Hunter Ragan, Pharm.D. Candidate; Carrie Vogler, Pharm.D. **Robert Robinson, MD** 

amine idely al setting		1916 adult discharges	
ions. k of ose			
to be umber of	16	Q 5	
ates for dications.	Inclu		
	1248 Not readmitted	Rea	
30-day	within 30 days	withi	
eduction	Table 1: Study populat	tion Baseline	
bination		Not readmi days	
	Age, mean (SD)	n = 1248 63 (16)	
	Female (%)	208 (48%)	
	Length of stay (SD)	7.74 (7.17)	
	Hospital admissions in the last year (SD)		
by the	Emergency department visits in last 6 months (SD)	0.40 (1.25)	
d	HOSPITAL Score (SD)	3.85 (1.43)	
U	LACE Index (SD)	11.4 (2.44)	
	Charlson Comorbidity Score (SD)	4.97 (3.28)	
	<b>Medical Comorbidities (%)</b> <b>Myocardial infarction</b>	340 (27%)	
	Congestive heart failure	291 (23%)	
ere	Peripheral artery disease	114 (9%)	
nbination	Stroke	74(6%)	
ents	Dementia Chronic lung disease	38 (3%) 352 (28%)	
	Connective tissue disease	28 (2%)	
ne	Peptic ulcer disease	59 (5%)	
	Cirrhosis Dick stor with out converting tions	40(3%)	
	Diabetes without complications Diabetes with complications	268 (22%) 143 (12%)	
-	Paralysis	44 (4%)	
, etc.)	Renal disease	237 (19%)	
	Cancer Materia company	89 (7%)	
-	Metastatic cancer Oral Corticosteroid Therapy (%)	27 (2%) 208 (16%)	
	Gastric Acid Secretion Inhibitor (%)		
	Proton pump inhibitor	534 (43%)	
nd	H2 Receptor Antagonist	139 (11%)	
Ĩ	<b>Drug combinations (%)</b> <b>PPI + Oral Steroids</b>	96 (7%)	
	H2RA + Oral Steroids	21 (2%)	

### Results



76 (17%)

188 (43%)

34 (8%)

41 (9%)

9 (2%)

0.904

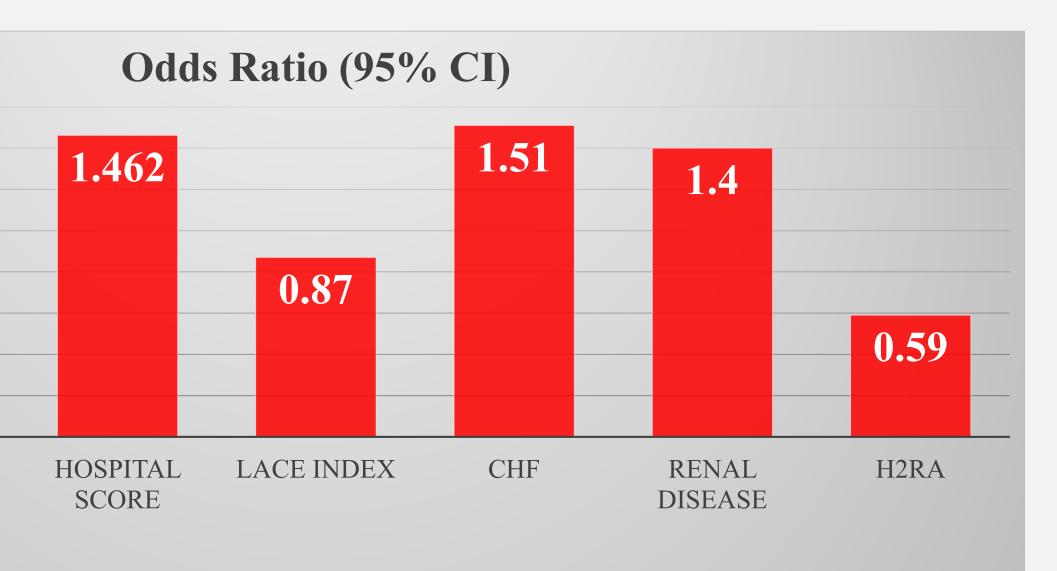
0.048

0.249

0.595



istic Regression of Potential Risk Factors with 30 Day Readmissions			
Odds Ratio (95% CI)	P value		
1.32 (1.15-1.50)	< 0.001		
1.25 (1.13-1.39)	< 0.001		
1.462 (1.31-1.63)	< 0.001		
0.87 (0.80-0.94)	< 0.001		
1.51 (1.12-2.04)	0.007		
1.40 (1.03-1.90)	0.031		
0.59 (0.36-0.93)	0.035		
	Odds Ratio (95% CI)   1.32 (1.15-1.50)   1.25 (1.13-1.39)   1.462 (1.31-1.63)   0.87 (0.80-0.94)   1.51 (1.12-2.04)   1.40 (1.03-1.90)		



# Discussion

Hospitalization within last year, ED visit within last 6 months, increased HOSPITAL score, CHF, and renal disease may indicate increased 30-day

• H2RA use and lower LACE index score associated with lower 30-day readmission risk

• PPI use alone was not associated with increased

Limitations include lack of standard dosing protocol and differences in baseline disease states

Corticosteroids in combination with PPIs or H2RAs were not associated with higher 30-day readmission

## Conclusion

This data shows no correlation between the use of PPIs or H2RAs alone or with concurrent oral corticosteroid use and 30-day readmission rates. Further investigation warranted before any clinical significance can be determined