

BACKGROUND

- A common inpatient drug-drug interaction or interest arises with daptomycin and hyroxymethylglutaryl-coenzyme A (HMG Cc inhibitors, or statins, due to the enhanced ri damage with concomitant administration.
- The mechanism of action for this interaction
- Statins are a commonly prescribed medical the primary and secondary prevention of AS
- Daptomycin is a cyclic lipopeptide antibiotic indicated for severe gram-positive infection
- The current daptomycin package insert reco clinicians consider temporarily suspending statins while daptomycin is being administe

OBJECTIVE

 To evaluate if patients admitted on statin the co-administration with daptomycin and the effects on creatine phosphokinase (CPK) la

- Study Design
- Retrospective chart review
- **IRB** Approval
- Springfield Committee for Research Involv Subjects Institutional Review Board
- Data Source
- 500 bed teaching hospital in Springfield, I Study Population
- Age 40 years old or older
- Admitted on a home medication of a statir
- Received at least 5 consecutive days of in daptomycin therapy
- Study Measures
- Primary Endpoint
- Home medication statin status while receiving inpatient daptomycin therapy

Management of Statins with Daptomycin Therapy Brian Batchelder, PharmD Candidate and Carrie Vogler, PharmD, BCPS

METH

of	 Secondary Endpoint CPK lab values and incide
oA) reductase	 Other Data Collected for A
risk of muscle	 Statin intensity Decomposition discontinuing
	 Reason for discontinuing ASCVD status (clinical c
n is debated tion class for	Data Analysis
SCVD events ¹	 Descriptive statistics
c typically ns ²	RES
commends	 82 participants included in f
the use of ered ³	<u>Table 1</u> Overall Demograph
	Age
	White/Caucasian Race
erapy receive resulting ab values	Male Sex
	Number of CPK Lab Values
ab values	Table 2 Statin Status
	Continued
	Discontinued
vina Human	Continued -> discontinued
ving Human	\Box :
	Figure 3 Reason for Statin
llinois	6%
า าpatient	13%
	75%

HODS	RESULTS				
	Table 4 Incidence of CPK Elevation				
idence of elevations	CPK Elevation (>200 IU/L)				
Analysis			Yes	No	
a any atudy drug	Statin	Continued	1	65	
ng any study drug or risk)	Status	Discontinued	3	10	
	Table 5 Statin Intensity Evaluation				
			High Intensity Statin*		
ULTS			Yes	No	
thic ctudy	Statin	Continued	29	37	
this study	Status	Discontinued	3	10	
hics Mean (SD) or % (N)	*Defined as atorvastatin 40 mg, atorvastatin 80 mg, rosuvastatin 20 mg, or rosuvastatin 40 mg daily.				
63.9 years (<u>+</u> 11.04)	Table 6 Clinical ASCVD Evaluation Clinical ASCVD				
82.93% (68)					
56.10% (46)			Yes	No	
1.59 (<u>+</u> 1.00)	Statin	Continued	23	43	
	Status	Discontinued	7	6	
% (N)					
70 (IN) 80.49% (66)	LIMITATIONS				
15.85% (13)	 Retrospective design Inappropriate CPK monitoring in 69.51% (57) participants Short duration of inpatient stay, limiting number of CPK lab values assessed 				
3.66% (3)					
Discontinuation (N = 16)		CONCLUSIONS			
 Unknown (12) Interaction with daptomycin (2) Elevated CPK (1) 	 Statins are often administered concomitantly with daptomycin in the inpatient setting Few incidences of CPK elevations occur when statins are administered with daptomycin Co-administration of statins with daptomycin may be a safe option for some patients with appropriate CPK 				

Patient NPO (1)

<u>References</u> 2018 AHA/ACC/AACVPR/AAPA/ABC/ACPM/ADA/ AGS/APhA/ASPC/NLA/PCNA Guideline on the Management of Blood Cholesterol Bland CM, Bookstaver PB, Lu ZK, et al; Southeastern Research Group Endeavor (SERGE-45). Musculoskeletal safety outcomes of patients receiving daptomycin with HMG-CoA reductase inhibitors. Antimicrob Agents Chemother. 2014;58(10):5726-5731.

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monitoring

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