# The use of order-based clinical decision support alerting to increase the homogeneity of premedication regimens in patients with known contrast allergies

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## INTRODUCTION

### **lodinated Contrast Utilization**

- 62 million CT scans completed annually in the US
- 0.7-3.2%: Prevalence of adverse reactions to non-ionic iodinated contrast
- · Symptoms range from mild (e.g. urticaria) to life-threatening (e.g. anaphylactic shock)

#### Premedication

- ACR recommends corticosteroids with or without an anti-histamine
- · Ideally, premedication begins 12-13 hours prior to study
- · Less effective regimens can be used in emergent settings
- IV steroids may have no effect when given <4-6 hours prior to study

## **Objectives**

- 1. Increase homogeniety of premedication use
- 2. Assess efficacy of CDS alerting intervention

## **METHODS**

## Electronic Health Record (EHR) Changes

- Clinical decision support (CDS) alert installed into Epic 4/7/2014
- · Providers alerted of patient allergies to intravenous iodinated contrast
- CDS prompts providers to order a recommended premedication regimen
- Alert discontinued if patient has adequate premedications ordered

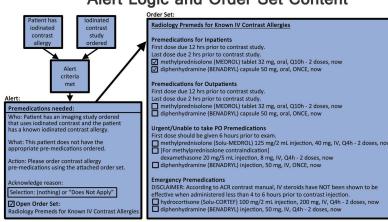
## **Premedication Analysis**

- Patients classified by premedication regimen received:
- 1. Preferred premedication regimen, based on ACR recommendations
- Corticosteroids <24 hours prior to study (i.e. not following recommendations)
- 3. No premedication with corticosteroid

## **Data Collection and Analysis**

- Retrospective analysis; 11 months pre- and post-implementation
- · Pre- and post-CDS patients compared
- Type of premedication regimen
- 2. Documented allergic reactions

## Alert Logic and Order Set Content



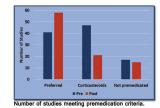
## RESULTS

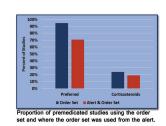
## Patient Population & Alert Firing

- 200 patients with documented allergy received IV contrast for a radiologic exam
- Alert fired appropriately for all premedication patients, only 2/3 of non-premedicated patients
- Alert did not fire inappropriately for any patients (100% specificity)
- Non-premedicated patients where alert did not fire had premedications not meeting criteria
- Overall alert sensitivity: 94.68%

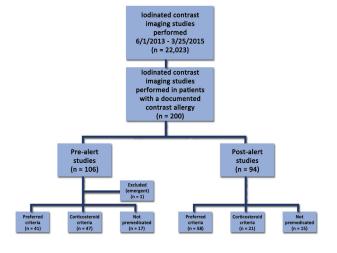
## Statistical Analysis

- Proportion of patients who received preferred regimen increased (Z-score=3.25, p=0.001)
- No difference in proportion of patients who were not premedicated (Z-score = -0.02, p=0.98)
- One patient had allergic reaction; occurred post-CDS in patient given preferred premedication





## Patient Population by Study Period and Premedication Criteria



## DISCUSSION

### Conclusion

- Homogeneity of premedication regimens significantly improved using CDS in Epic
- · Alert firing is primarily associated with increased orderset usage
- Order set usage led to increased homogeneity
- Sample size not sufficient to analyze number of contrast reactions
- Alert sensitivity suboptimal as premedications have other clinical indications, thus may cause alert to not fire inappropriately

### **Future Direction**

- · Additional analysis needed to see if CDS has lead to increased patient safety
- Further steps to ensure patient compliance with premedication
- Implement similar CDS designed for allergies to gadolinium
- Further optimize alert firing for all relevant studies

#### References

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