# Developing A Medical Question Answering Model for Duke Department of Medicine

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### Objective

#### Background

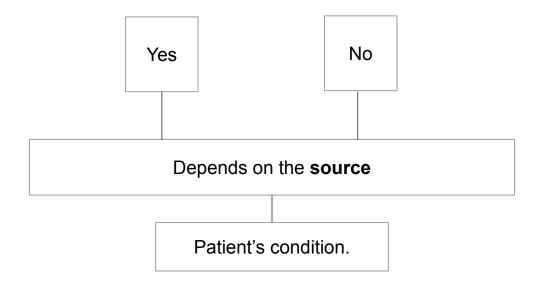
As the number of research papers and guidelines increase, it becomes a
 tremendous burden for physicians to stay up-to-date to make informed
 decisions faced with various clinical conditions.

#### Objective

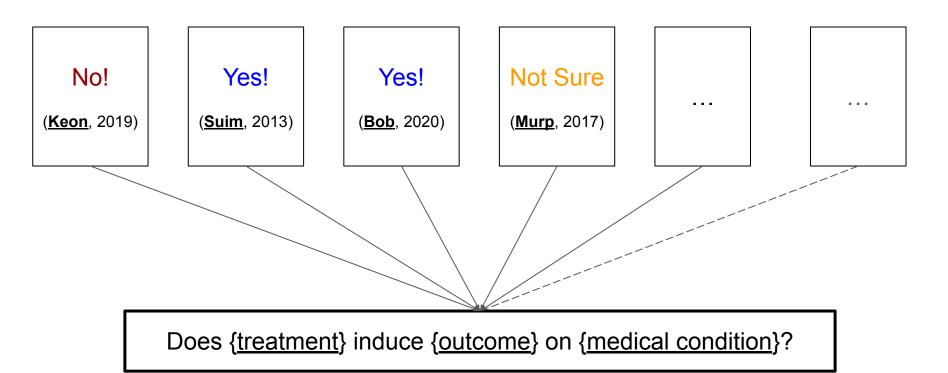
 The project aims to develop of an evidence-based medical question answering model for critical care medicine

# An Example Question

Do patients with severe **ARDS** (Acute Respiratory Distress Syndrome, a <u>medical</u> <u>condition</u>) being treated with **neuromuscular blocking** (<u>medication</u>) agents have increased **muscle weakness**? (<u>Outcome</u>)

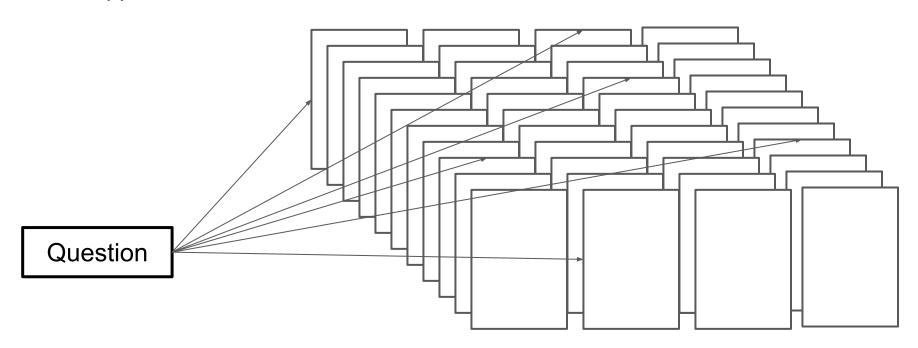


## Medical Questions Answered by Research Papers



# **Quantity Counts**

What happens when the amount of source document becomes HUGE?



# Output Format

**Papers Supporting** The Claim

Papers Against The Claim

Medical Question/ Claim:

**Supporting** the claim:

Do <u>steroids</u> improves <u>survival</u> and reversal of shock in patients with <u>septic shock</u>?

Paper 1

Study Design and Methodology:

Study Population: Interventions:

Comparator:

Outcomes:

Strengths and Weaknesses:

Key Findings and Conclusion:

Paper 2

**Against** the claim:

Paper 3

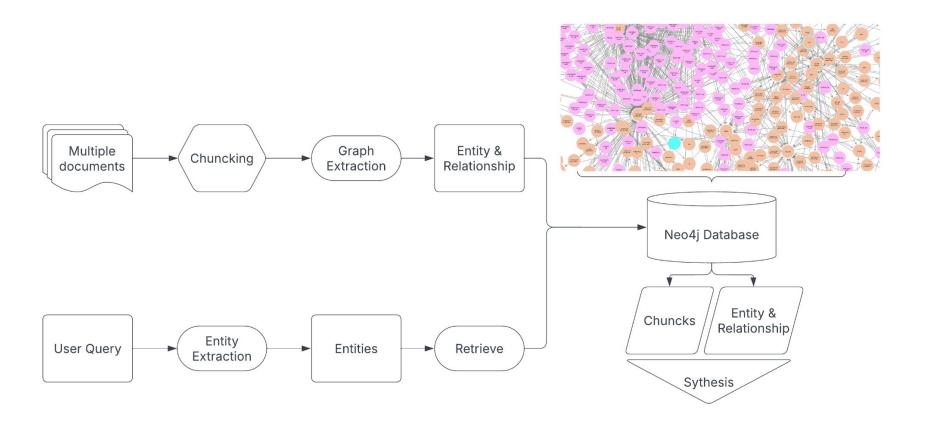
Study Design and Methodology:

Study Population:

A Summary of each paper

**User Question** 

### Retrieval Augmented Generation Plus Knowledge Graph



### **Dataset**

#### Research Papers

- <u>112 research papers</u> related to any medical conditions are collected for model build up and evaluation. (34 papers from WikiJournal, 86 from PubMed, Google Scholar)

#### Medical Questions

- <u>36 yes-no medical questions</u> (12 for ARDS, 10 for sepsis, 6 for cardiac arrest, 7 for delirium, 1 for Sepsis *and* delirium).

### **Ground Truth Annotation**

 <u>Each</u> research paper was annotated in terms of relevance and stance against <u>all</u> in medical questions (yes-no) or claim.

#### **Relevance Annotation**

	Q1	Q2	
Paper 1	1 (relevant)	0 (irrelevant)	
Paper 2		1 (relevant)	

#### **Stance Annotation**

	Q1	Q2	
Paper 1	1 (supporting)	0 (neutral)	
Paper 2		-1 (against)	

### **Ground Truth Annotation II**

 For <u>each research paper</u>, we leveraged large language model (Claude from AWS Bedrock), to generate summaries. After editing and proofreading by physicians (Keon), the summaries serve as ground truth.

	Study Design and Methodology	Study Population	Interventions	•••	Key Findings and Conclusion
Paper 1					
Paper 2					

### **Evaluation**

#### Relevance

 Relevance of retrieved papers measured measured by <u>accuracy</u>, <u>precision</u>, <u>recall</u>, and <u>F-1 score</u>.

### Factuality

- Summaries are evaluated on <u>factual correctness</u> and <u>semantic similarities</u> against <u>ground truth</u> using <u>Ragas</u>.

### Results - Retrieval

	Recall	Precision	F-1	Accuracy
Naïve RAG (Suim)	.476	.560	.444	.896

<sup>\*</sup> Mean value of all metrics from 36 questions

### Issue - Hallucination

#### Example Question/ Claim:

#### ← NON EXISTING PAPER

Does early administration of neuromuscular blocking agents increases the ventilator free days?

#### **Supporting** the claim:

#### Paper 2: Impact of Early Neuromuscular Blockade on ARDS Mortality

- a. Authors: Moss et al.
- b. Year: 2016
- c. Journal: Critical Care Medicine
- d. Research Question: Does early NMBA administration affect mortality in ARDS patients?
- e. **Study Design and Methodology**: Cohort study analyzing data from a large database.
- f. **Study Population**: 1,000+ ARDS patients.
- g. **Interventions & Comparators**: Early NMBA administration vs. standard care.
- h. **Outcomes**: Decreased mortality and increased ventilator-free days.
- i. **Strengths and Weaknesses**: Large sample size is a strength; retrospective design is a weakness.
- j. **Key Findings and Conclusion**: Supports the claim with significant findings on mortality and ventilator-free days.

### Issue - Incorrect Stance

### Equivocal or Inconclusive:

- 1. \*\*Study 1\*\*
  - \*\*Authors:\*\* Annane et al.
  - \*\*Year:\*\* Not specified
  - \*\*Title:\*\* "Effect of Treatment With Low Doses of Hydrocortisone and Fludrocortisone on Mortality in Patients With Septic Shock"
  - \*\*Journal:\*\* Not specified
- \*\*Research Question:\*\* Examined the impact of low-dose corticosteroids on 28-day survival in patients with septic shock and adrenal insufficiency.
- \*\*Type of Study:\*\* Placebo-controlled, randomized trial
- \*\*Methodology:\*\*
- Randomized patients to hydrocortisone and fludrocortisone vs. placebos.
- Main outcome: 28-day survival in nonresponders to corticotropin.
- \*\*Key Findings:\*\*
- No significant difference in 28-day mortality between hydrocortisone and placebo.
- No difference in mortality between responders and nonresponders to corticotropin.
- \*\*Conclusion:\*\* Low-dose corticosteroids did not improve 28-day survival in septic shock patients with adrenal insufficiency.

**Question**: Does early application of high frequency oscillatory ventilation compared with ventilation strategy of low tidal volume decrease mortality?

**Ground Truth**: Supporting (1) / **Synthesis**: Equivocal or Inconclusive (0)

# **Synthesis Evaluation**

	Study Design and Methodology
Ground Truth (ACURASYS)	<ul> <li>Multicenter, randomized, double-blind, placebo-controlled trial</li> <li>March 2006 through March 2008</li> <li>20 Intensive Care Units (ICUs) in France</li> <li>Patients randomly assigned to cisatracurium or placebo groups</li> <li>Double-blind design</li> <li>Independent data and safety monitoring board</li> </ul>
Generated Summary	- Multicenter, double-blind trial - Randomly assigned patients to receive cisatracurium besylate or placebo.
Score	Factual Correctness (Recall): 0.25 Factual Correctness (Precision): 0.67 Semantic Similarity: 0.86

	Study Population
Ground Truth (ACURASYS)	The study enrolled <b>340 patients</b> from 20 ICUs in <b>France</b> . The patients were randomly assigned to either receive cisatracurium besylate (178 patients) or a placebo (162 patients) for 48 hours. The study population consisted of patients with severe ARDS, defined as a ratio of partial pressure of arterial oxygen (PaO2) to the fraction of inspired oxygen (FIO2) of less than 150, with a positive end-expiratory pressure of 5 cm or more of water, and a tidal volume of 6 to 8 ml per kilogram of predicted body weight Total Participants: 340 patients - Inclusion Criteria: - Receiving endotracheal <b>mechanical ventilation</b> - Acute hypoxemic respiratory failure - <b>Severe ARDS</b> (PaO2:FIO2 < 150) - ARDS onset within previous 48 hours - Cisatracurium Group: 178 patients - Placebo Group: 162 patients
Generated Summary	- 340 patients
Score	Factual Correctness (Recall): 0.08 Factual Correctness (Precision): 1.0 Semantic Similarity: 0.45

# Wrapping up

 We developed a knowledge-graph-based RAG model able to retrieve relevant document regarding medical questions in critical care medicine (accuracy ~ 90%) and provide paper summaries to facilitate clinical decision support.

#### Next steps

- Improve performance (retrieval, synthesis, reduce hallucination)
- Publish the work

# Backup