

# Data Science 101: Electronic Health Record (EHR) Data

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Montefiore

### **Today's Presentation**



- Logistics of Working with EHR Data
- Challenges Inherent in EHR Data
- Methodological Considerations
- Building EHR Research Team





### What Are EHR Data?



### Digital record of care delivery generated during clinical encounters

Demographics

Encounter Information

Diagnoses & Problems

Procedures & Orders

Medications

Laboratory, Test, & Vital Signs Measurements Clinical Narratives and Reports Special Tests and Reports (ECG, EEG, genomics)





## **Why EHR Data Matter**





Nearly every patient encounter generates digital data



Huge scale, real-world context, inexpensive and quick to obtain



Enables questions impractical or unethical for trials





### Case Study: Tocilizumab in COVID-19

### **Original Investigation**

FREE

October 20, 2020

# Association Between Early Treatment With Tocilizumab and Mortality Among Critically Ill Patients With COVID-19

Shruti Gupta, MD, MPH<sup>1</sup>; Wei Wang, PhD<sup>2</sup>; Salim S. Hayek, MD<sup>3</sup>; et al

> Author Affiliations | Article Information

JAMA Intern Med. 2021;181(1):41-51. doi:10.1001/jamainternmed.2020.6252

Mortality OR 0.68 (0.46-0.99)

### **Study Design and Oversight**

We emulated a hypothetical target trial in which critically ill adults with COVID-19 received or did not receive tocilizumab in the first 2 days of intensive care unit (ICU) admission. We used data from the Study of the Treatment and Outcomes in Critically Ill Patients With COVID-19 (STOP-COVID), a multicenter cohort study that enrolled consecutive adults with laboratory-confirmed COVID-19 (detected by nasopharyngeal or oropharyngeal swab) admitted to participating ICUs at 68 hospitals across the United States (eTable 1 in <u>Supplement 1</u>). Study personnel at each site collected data by detailed medical

### Case Study: Tocilizumab in COVID-19

Mortality OR: 0.65 (0.43-0.99)







### Case Study: Tocilizumab in COVID-19

Tocilizumab in patients admitted to hospital with COVID-19 (RECOVERY): a randomised, controlled, open-label, platform trial

Affiliations & Notes ✓ Article Info △ Linked Articles (7) ✓

Publication History: Published May 1, 2021

DOI: 10.1016/S0140-6736(21)00676-0 ¬ Also available on ScienceDirect ¬

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Mortality OR: 0.76 (0.66-0.88)

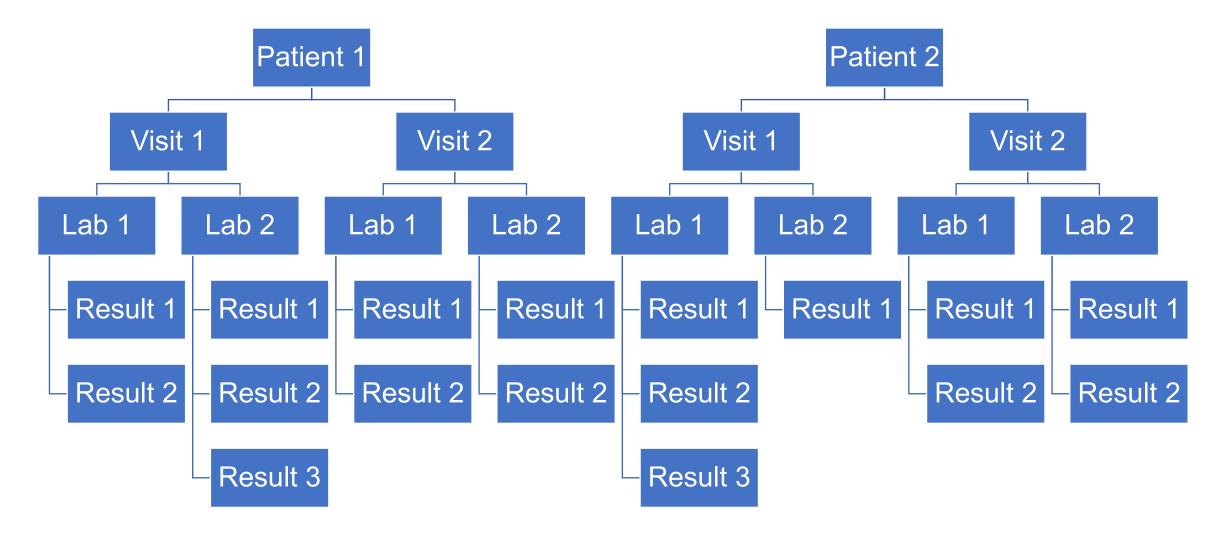




# Working with EHR Data

### **How Are EHR Data Stored?**





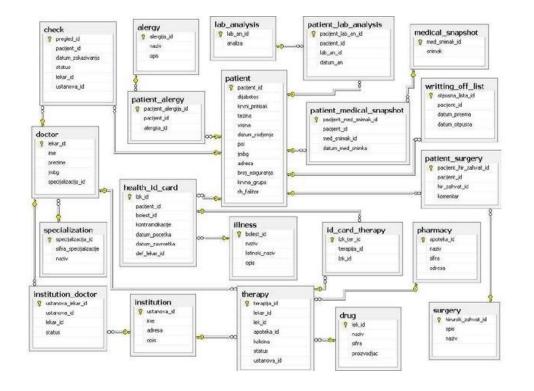




### **EHR Data Structure**



- Stored in relational tables (patients, visits, meds, labs, notes...)
- Requires joins, cleaning, reshaping
  - SQL and data wrangling skills
- Different vendors & data models
  - Epic alone has 3 data models!







### **Epic Clarity Data Dictionary**

Over 18,000 tables!



A B C D E F G H I K L M N O P Q R S T U V W

### Tables Starting with A Top of page of

- ABN\_DOCUMENT\_ID
- ABN\_FOLLOW\_UP
- ABN MEDICATIONS
- ABN\_NOTES
- ABN\_NOTE\_COMMENTS
- ABN\_NOTE\_CONTACT\_SERVICE
- ABN\_NOTE\_PROC
- ABN\_ORDERS
- ABN\_ORDER\_INFO
- ABN\_PLAN\_INFO
- ABN SERVICE INFO
- ACCCATH3\_MED\_DISCH
- ACCESSIBLE\_DOCUMENTS\_PREF
- ACCESSIBLE SERVICES
- ACCOUNT
- ACCOUNT\_2
- ACCOUNT 3
- ACCOUNT\_CONS\_SP\_SA\_BILL
- ACCOUNT\_CONTACT
- ACCOUNT CONTACT 2
- ACCOUNT\_CREATION
- ACCOUNT\_DISCON\_CVG
- ACCOUNT\_FPL\_INFO
- ACCOUNT\_MYC\_GP\_FAILED\_LOG
- ACCOUNT\_RQG\_GROUPERS

#### **ORDER PROC**

**Description:** The ORDER\_PROC table enables you to report on the procedures ordered in the clinical system. We have also included patient and contact identification information for each record.

Discontinued?

Type

VARCHAR

#### **Primary Key**

Name

Column Name Ordinal Position
ORDER PROC ID 1

The order type category number for the procedure order. May contain organization-specific values: Yes

Medications

#### **Column Information**

Category Entries:

9 PROC ID PROC NAME

1	ORDER_PROC_ID	NUMERIC	No	
	The unique ID of the order record associated with this procedure order.			
2	PAT_ID	VARCHAR	No	
	he unique ID of the patient record for this order. This column is frequently used to link to the PATIENT table.			
3	PAT_ENC_DATE_REAL	FLOAT	No	
	• •		of the number indicates the date of the contact. The digits after the decimal distinguish different contacts on the .00 is the first/only contact, .01 is the second contact, etc.	
4	PAT_ENC_CSN_ID	NUMERIC	No	
	The unique contact serial number for this contact (UCI).	. This number is uniq	ue across patients and encounters in your system. If you use IntraConnect this is the Unique Contact Identifier	
5	RESULT_LAB_ID	NUMERIC	No	
	The unique ID of the lab or other resulting agency	,, such as radiology, t	that provided the order results.	
6	RESULT_LAB_ID_LLB_NAME	VARCHAR	No	
	Interface laboratory name.			
7	ORDERING_DATE	DATETIME	No	
	The date when the procedure order was placed.			
8	ORDER_TYPE_C_NAME	VARCHAR	No	

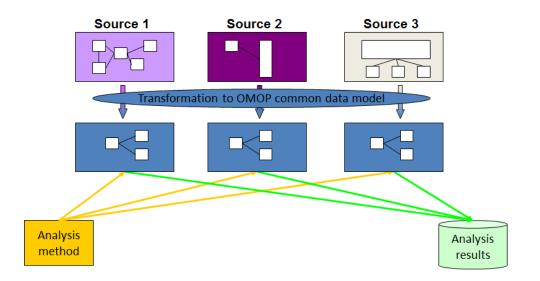


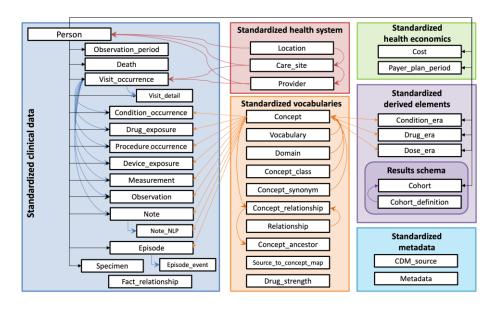


### **Common Data Models**



- OMOP, PCORnet standardize structure & vocabularies for research
- Enable multicenter studies, repositories, shared tools









### **How Are EHR Data Accessed?**



- EHR data contain protected health information (PHI)
- Use governed by IRB, HIPAA, and institutional data use policies
- Often accessed through secure enclaves or virtual data centers
  - Access restricted to IRB-approved subset of data
  - Structured Query Language (SQL) or specialized user interface tools (ATLAS, SlicerDicer, etc)
- Data rarely exported; analyses happen inside the firewall/enclave





### **EHR Data Analyses**



### Description

- What is currently happening?
- Example: what proportion of patients with heart failure with reduced ejection fraction get readmitted within 30 days?

### Prediction

- What will likely happen?
- Which patients will likely have adverse outcomes after a procedure?

### Causal Inference

- What would happen under different courses of action?
- How would outcomes differ if we used terlipressin versus current standard of care as first-line treatment for HRS?





### **Phenotyping**



- Unlike in prospective studies (including clinical trials), EHR data does not have pre-defined cohorts or study-specific variables
- Phenotyping = process of turning available data into study-specific features
  - This is a key step for working with EHR data
- Using data as a surrogate of true disease or health event
  - Example: COPD exacerbation
- Can be done using code (SQL) or specialized tools (ATLAS, TriNetX)





### **Cohort Building**



- Descriptive and feasibility analyses can often be conducted using simple queries
  - E.g. how many COPD admissions do we get per year?
- Most clinical research questions are person-centric
  - ► E.g. in patients admitted with COPD not on triple therapy, how would outcomes differ if we initiated triple therapy in the hospital versus leaving it to the outpatient provider?
- Clinical questions require cohort framework and complex queries
  - Foundation for causal inference and real-world evidence generation







## **Cohort Definition Framework**

Cohort entry event – what triggers the patient to be evaluated for your cohort?



Inclusion criteria – patients who meet them at entry event enter your cohort



Exit criteria – what triggers the patient to exit cohort?





### **Project Example: COPD**



- Study question: in patients admitted with COPD not on triple therapy, how would outcomes differ if we initiated triple therapy in the hospital versus leaving it to the outpatient provider
- Phenotyping tasks:
  - Eligible patients
    - Admitted with COPD exacerbation
    - Not on triple therapy at the time of admission
  - Prescribed triple therapy at discharge
- Phenotypes combined into complex study cohort





## **Project Examples: COPD**



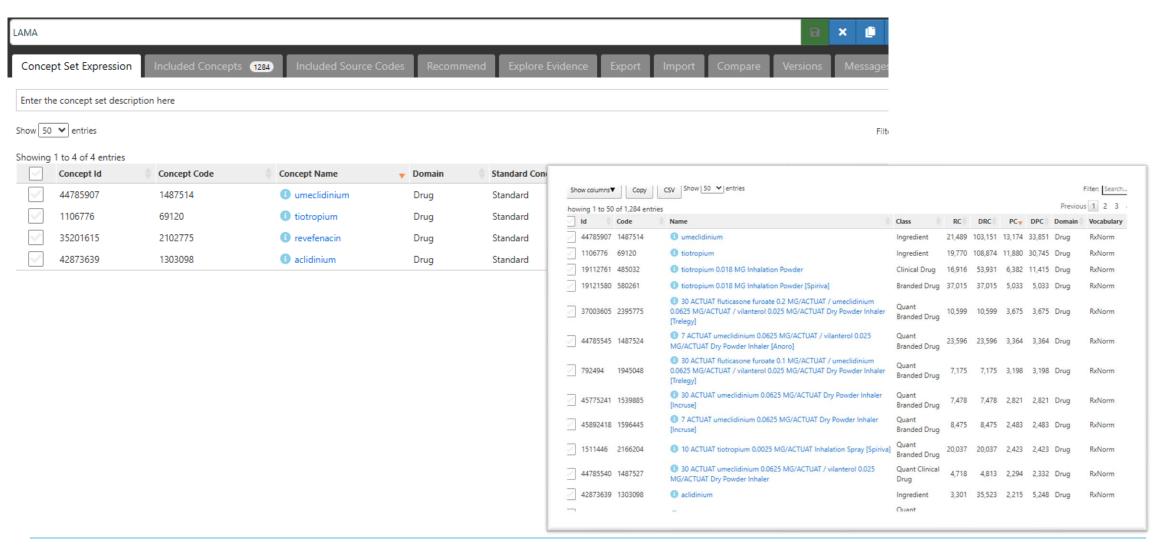
Cohort Entry Events	8
Events having any of the following criteria:	
a condition occurrence of COPD all diagnoses	0
occurrence start is: after v 2022-01-01	×
with a condition status of: Primary diagnosis Add Import	×
with a Visit occurrence of: X Inpatient Visit Add Import	×
+	
with continuous observation of at least 180 v days before and 0 v days after event index date  Limit initial events to:   all events v   per person.	
Limit initial events to: all events to:  Restrict initial events to:	
having any of the following criteria:	+
with exactly ▼ 0 ▼ using all occurrences of:	•
a drug era of LAMA - ingredient only (for dru 🔻	
+	
where event.starts between	
All v days Before v and 1 v days Before v index start date  X and event ends between 90 v days Before v and All v days After v index start date	
The index date refers to the event from the Cohort Entry criteria.	
allow events from outside observation period	
or with exactly ▼ 0 ▼ using all occurrences of:	•
a drug era of LABA - ingredient only (for dru 🔻	
+	
where event starts between	
All ▼ days Before ➤ and 1 ▼ days Before ➤ index start date  X and event ends between 90 ▼ days Before ➤ and All ▼ days After ➤ index start date	
The index date refers to the event from the Cohort Entry criteria.	
allow events from outside observation period	





### **LAMA Concept Sets**









### **SQL** Code



```
select event_id, person_id, start_date, end_date, op_start_date, op_end_date
     into #included events
       SELECT event_id, person_id, start_date, end_date, op_start_date, op_end_date, row_number() over (partition by person_id order by start_date ASC) as ordinal
         select Q.event_id, Q.person_id, Q.start_date, Q.end_date, Q.op_start_date, Q.op_end_date, SUM(coalesce(POWER(cast(2 as bigint), I.inclusion_rule_id), 0)) as inclusion_rule_mask
         from #qualified events Q
         LEFT JOIN #inclusion_events I on I.person_id = Q.person_id and I.event_id = Q.event_id
         GROUP BY Q.event id, Q.person id, Q.start date, Q.end date, Q.op start date, Q.op end date
        ) MG -- matching groups
      -- the matching group with all bits set ( POWER(2,# of inclusion rules) - 1 = inclusion_rule_mask
       WHERE (MG.inclusion_rule_mask = POWER(cast(2 as bigint),4)-1)
339 ) Results
345 select event_id, person_id,
346 case when DATEADD(day,1,end_date) > op_end_date then op_end_date else DATEADD(day,1,end_date) end as end_date
347 INTO #strategy ends
348 from #included events;
352    select person_id, start_date, end_date
353 INTO #cohort_rows
354 from ( -- first_ends
        select F.person id, F.start date, F.end date
          select I.event_id, I.person_id, I.start_date, CE.end_date, row_number() over (partition by I.person_id, I.event_id order by CE.end_date) as ordinal
           from #included_events I
361 -- End Date Strategy
```

445 lines







# Case Study: Local EHR Data Access at Montefiore Einstein

- Epic EHR + specialized data warehouse mapped to OMOP common data model
- Accessible via ATLAS for cohort building and analytics + Epic tools for basic reporting (SlicerDicer)
  - SQL access only for informatics analysis or deidentified data
- Extraction possible to a Montefiore-managed device for local analysis
  - Built-in IRB check to ensure extraction approved
- Larger extractions and ECG/DICOM require work on HPC
  - Has SQL/R/python available





### **Case Study: Epic Cosmos**



- 208 institutions across U.S. contribute data
- Access via participating institution (institutional data use agreements)
- SlicerDicer-like interface for basic summary-level queries
- For patient level data, must undergo Epic certification as Data Scientist or Data Architect
- Project review and approval by Epic + local governance
- Data access fees
- Remote repository and virtual analytic workspace (SQL, R, python)
  - All data and code stay on remote workspace and are audited





# Data-Specific Challenges

## **EHR: Not Designed for Research**





Built for care delivery, billing, regulatory compliance



Documentation priorities do not align with research priorities



Results: incomplete, inconsistent, biased





### **Data Completeness & Bias**



- Data present when patients seek care at the institution
- Health status influences follow-up & testing
  - Unlike in trials where regular follow up is built-in
- Coding practices largely driven by reimbursement incentives
- Insurance, access, and location drive representativeness





### **Data Quality & Structure**



- Not all data structured (notes, images, PDFs)
- Structured fields inconsistently filled
- Standard codes (LOINC, RxNorm) incomplete + local quirks
- Solutions:
  - Mapping concepts to standardized vocabularies
  - Extraction of structured concepts from unstructured data





## **Example: CAM-ICU**



- Used to indicate delirium
- Theoretically two options: delirium present/absent
- Actual values

	MEAS_VALUE	★ TOTAL_OCCURRENCES
1	Negative - not Delirious	943862
2	Unable to Assess	722439
3	Delirious	44327
4	Incomplete	23170
5	(INCOMPLETE)	20278
6	Unable to assess	19549
7	Not delirious	11515
8	(null)	3681
9	Negative	2907
10	Positive	1781





## Irregular/Nonrandom Measurement



- Labs/vitals not done on fixed schedule
- Healthy patients less likely to return → nonrandom missing data
  - Can bias finding if inappropriately handled



## Irregular Measurements: Example



You're comparing two cohorts – those prescribed SPS versus patiromer (two potassium binding agents) You're interested in checking their potassium lowering trends over next 6 months Challenges Some never return at 6 months Some will only return at month 1 or 6 Some will only return at months 2, 3, and 5 What are some potential solutions?





## Irregular/Nonrandom Measurement



- Labs/vitals not done on fixed schedule
- Healthy patients less likely to return → nonrandom missing data
  - Can bias finding if inappropriately handled
- Possible solution: require evidence of regular care pre-baseline





# Nonstandard & Complex Data Types

- Imaging, ECG waveforms, pathology slides
- Require specialized methods (NLP, AI/ML)
- Integration with structured data is difficult
- Potential solutions:
  - Signals/images into structured features (e.g. QT interval, digital pathology quantified features)
  - Natural language processing to extract distinct findings





### **Natural Language Processing**



FLT-ITD mutated AML with monocytic differentiation





# Methodological Challenges





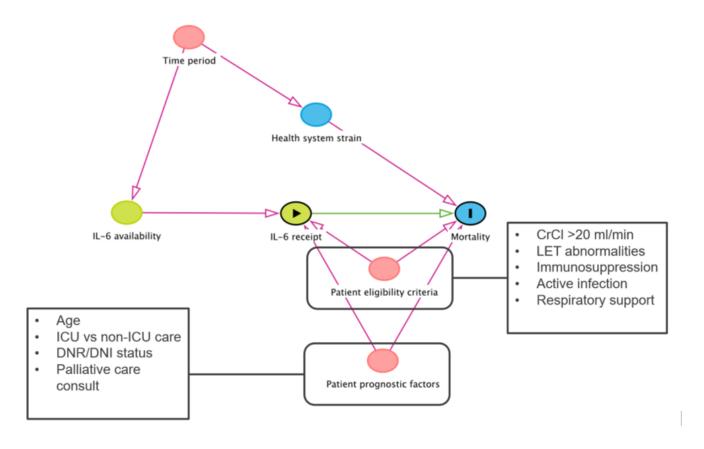
- Real-world treatment selection is not random.
- Indication, severity, provider preference drive choice
- Need to measure & adjust for these factors















### **Defining Time Zero**



- No natural randomization/enrollment date
- Must define cohort entry & follow-up start carefully
- Mis-specification of time zero leads to bias





### **Time Zero Bias in Practice**





#### Current Problems in Cardiology

Volume 44, Issue 10, October 2019, 100407



### Heart Failure Postdischarge Clinic: A Pharmacist-led Approach to Reduce Readmissions

Rasha Al-Bawardy MD, Angela Cheng-Lai PharmD, Lendita Prlesi PharmD, Manaf Assafin MD, Shuo Xu ScM, Kiana Chen, Samvit Tandan MD, PhD, Chino S. Aneke MD, PhD, Sandhya Murthy MD, Ileana L. Piña MD, MPH









https://doi.org/10.1016/j.cpcardiol.2018.12.004 7

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#### Introduction

Heart failure (HF) is a major cardiovascular syndrome that affects approximately 5.7 million adults in the United States.<sup>1</sup> Although the survival rate after HF diagnosis has improved over time, the death rate from this disease remains high. It is estimated that 50% of those diagnosed with HF will die within 5 years after the first diagnosis. The cost



#### Current Problems in Cardiology

Volume 48, Issue 2, February 2023, 101507



### **Evaluating Pharmacist-Led Heart Failure** Transitions of Care Clinic: Impact of Analytic Approach on Readmission Rate Endpoints

Angela Cheng-Lai a b c ス In Lendita Prlesi A, Sandhya Murthy b c, Eran Y. Bellin b c d, Mark J. Sinnett a, Pavel Goriacko a c d

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https://doi.org/10.1016/j.cpcardiol.2022.101507 7

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#### Abstract

Studies evaluating pharmacist-led transitions of care (TOC) services for heart failure patients reported profound decreases in hospital readmissions, Most studies restricted their analysis to clinic attendees (as-treated analysis), which can introduce selection and immortal time bias. In this study, we evaluated the impact of including only clinic attendees vs all clinic referrals in assessing the effectiveness of a pharmacist-led heart failure transitions of care (PharmD HF TOC) clinic program on 30-day readmissions. This





### **Loss of Longitudinally**



- Detailed encounter data but fragmented over time
- Patients move between systems (especially in NYC)
- Leads to misclassification and missing data bias
- Possible solutions: repositories + linkage
  - Claims, registries, mortality → fill longitudinal gaps
- Analytic solution: restrict cohort to those "engaged with health system"
  - Based on regular previous visits





### **Key Takeaways**



- EHR data is not research ready, needs cleaning & careful design
- Bias, missingness, and access hurdles are real
- Multidisciplinary expertise is essential (esp. BERD and Informatics)
- When done well: faster, cheaper, ethically feasible studies





### **Building Your EHR Research Team**



### **Clinical Expertise**

Frame meaningful questions, interpret clinical context, validate findings

## Informatics and Data Science

Extract, transform, and manage complex EHR data with institutional and regulatory compliance

## Epidemiology and Biostatistics

Design studies, address bias, and analyze results appropriately







# Electronic Health Record (EHR) Data in Clinical Research

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