Research IT Information & Exchange Series

Collaborative Opportunities and Tools in Informatics Research

April 28, 2017

Research IT Information & Exchange Series

- **Goal:** To educate pediatric researchers on the Research IT and Informatics resources and expertise available to facilitate their research and to identify areas where we can enhance IT methods to better support research.
- **Format:** One hour sessions led by subject matter experts to present information on the current services and expertise available.
- Intended audience: Researchers with an interest in capitalizing on Research IT tools to make their research better. Also, researchers who are interested in using Big Data and Healthcare Analytic approaches in their research.

Research IT Information & Exchange Series:

Learn about suitable collaborators and specific areas of expertise in Atlanta based institutions. We will also highlight and discuss best practices in these types of collaborations.

Research IT Information & Exchange Series Our presenters today



- Gari D. Clifford, PhD, Interim Chair, Associate Professor, Biomedical Informatics, Emory University, Associate Professor, Biomedical Engineering, Georgia Institute of Technology
 - gari.clifford@bme.gatech.edu



- Jon Duke, MD, Director of Health Data Analytics, Georgia Tech Research Institute
 - Jon.Duke@gatech.edu

- Dr. Gari Clifford and his team presented work that is to be published and therefore cannot be shared until it is published.
- The video from today's session does NOT include this part of the presentation, but starts with the presentation by Dr. Jon Duke.
- Dr. Jon Duke's slides follow





Applying Big Data to Big Ideas: Collaborating with Georgia Tech in Health Analytics

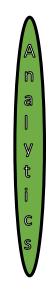
April 28th, 2017

Jon Duke, MD, MS



(Big) Data





Decision Support



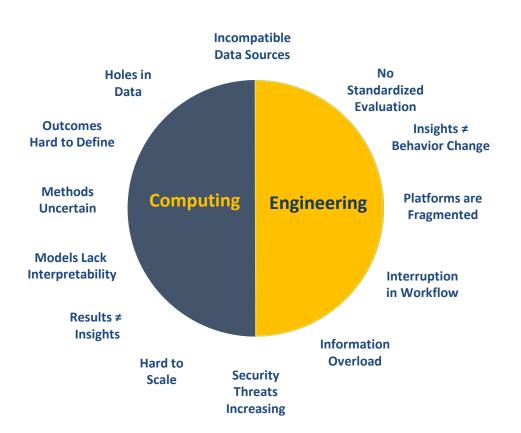




Targeted Action

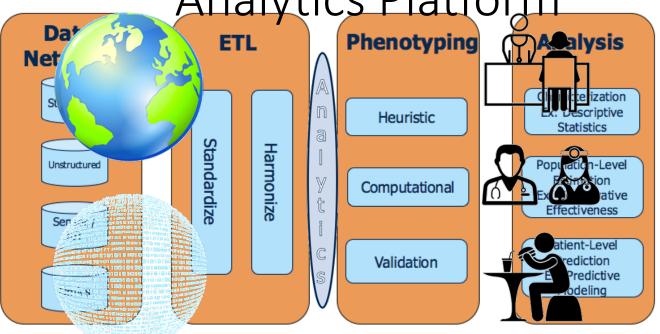


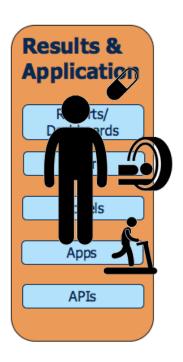




GT Health Data

Analytics Platform



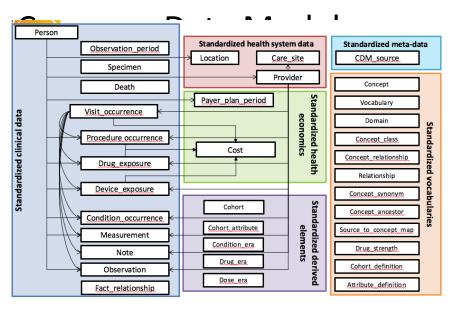


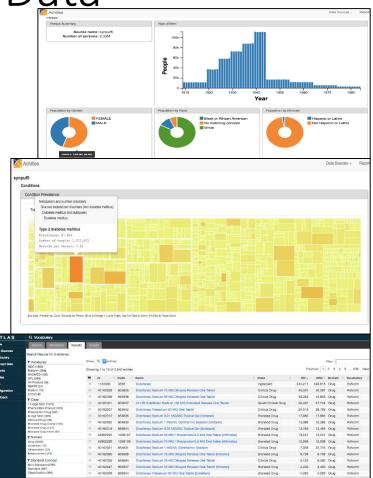
Example: C. Difficile

- Clostridium Difficile colitis is a hospital-borne infection associated with antibiotic use
- C. Diff costs the US Healthcare System \$4.8 billion annually and is a major target for infection control efforts nationally and internationally
- Predicting risk of C Diff infection can optimize prescribing and early detection

Step One: Harmonize Data

 Structured data is harmonized to the OMOP









OHDSI Collaborators:

- >200 researchers in academia, industry, government, health systems
- >20 countries
- Multi-disciplinary expertise: epidemiology, statistics, medical informatics, computer science, machine learning, clinical sciences

Databases converted to OMOP CDM within OHDSI Community:

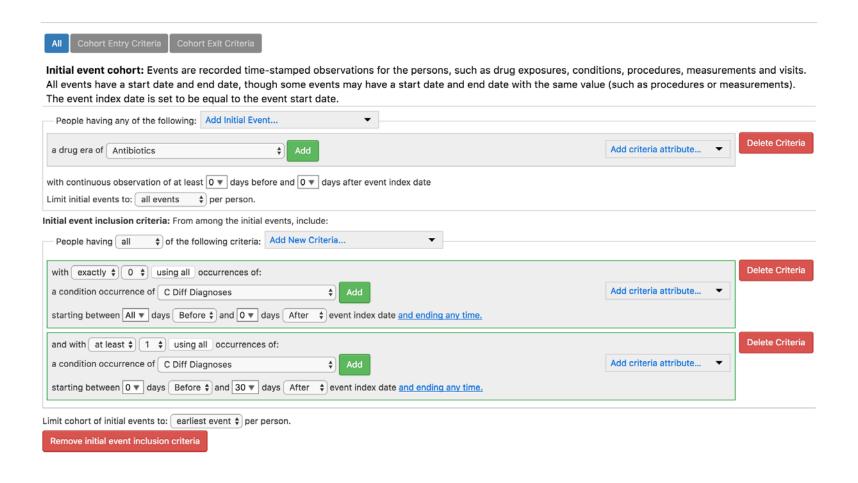
- >55 databases
- >660 million patients

Step One: Harmonize Data

 Unstructured data is analyzed via NLP tools including cTakes and GT-developed open-

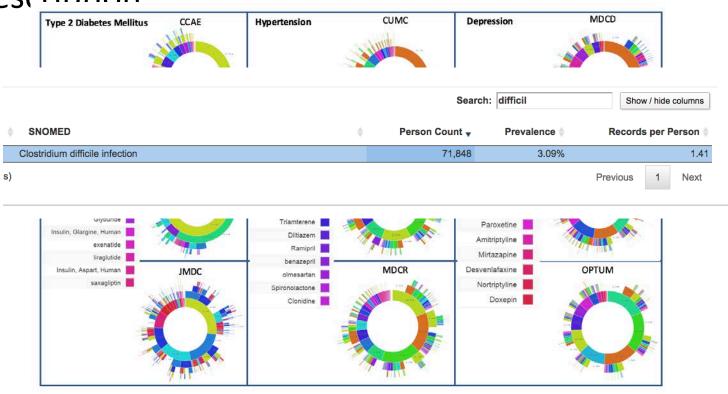


Step Two: Create Phenotypes



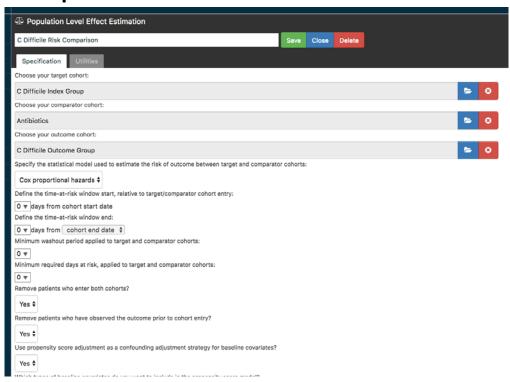
Step Three: Run Analyses

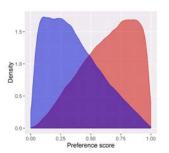
Description

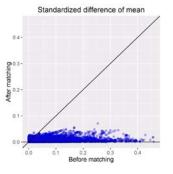


Step Three: Run Analyses

Comparison

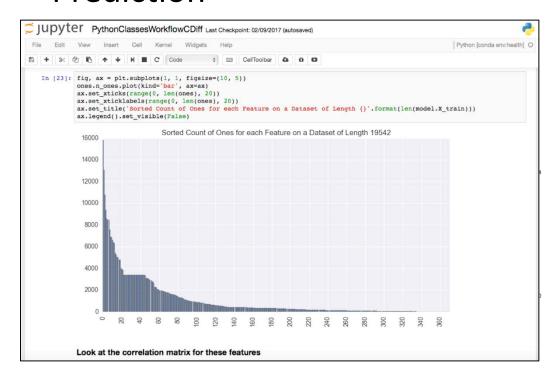




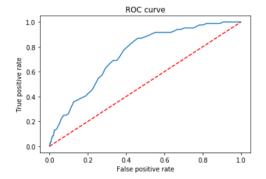


Step Three: Run Analyses

Prediction

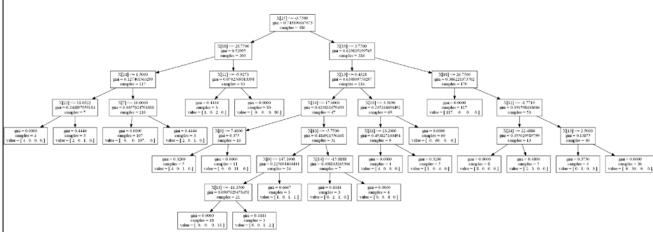


Accuracy
0.797697620875
Confusion matrix
[[5162 1269]
[49 35]]
Normalized confusion matrix
[[0.80267455 0.19732545]
[0.58333333 0.41666667]]
Precision
0.0268404907975
F1 score
0.0504322766571
AUC
0.740352718603

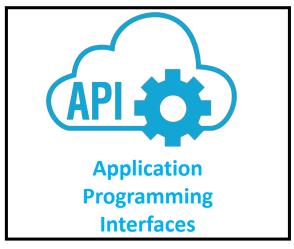


Long Way from Predictive Models to Decision Support

Classification Tree	P(Target)	# Inst
▼ <root></root>	0.090	2071
▼ MAX_POT <=5.050	0.050	1655
▼ MAX_CALCIUM <=10.150	0.048	1467
▼ PRE_HYPERK = 0	0.038	1416
▶ AGE <=65.500	0.032	753
▶ AGE >65.500	0.045	663
▼ PRE_HYPERK = 1	0.333	51
DM = 0	0.500	24
▶ DM = 1	0.185	27
▼ MAX_CALCIUM > 10.150	0.059	188
HTN = 0	0.167	6
▼ HTN = 1	0.055	182
Max_CR <=3.700	0.050	179
Max_CR > 3.700	0.333	3
▼ MAX_POT >5.050	0.250	416
▼ PRE_HYPERK = 0	0.138	327
AGE <=55.500	0.098	51
▼ AGE >55.500	0.145	276
▶ AGE <=65.500	0.148	88
▶ AGE >65.500	0.144	188
▼ PRE_HYPERK = 1	0.663	89
▼ MAX_POT <=6.250	0.519	52
DM = 0	0.933	15
▶ DM = 1	0.351	37
▼ MAX_POT >6.250	0.865	37
▶ CHF = 0	0.778	9
CHF = 1	0.893	28



Step Four: Connect to the World with APIs





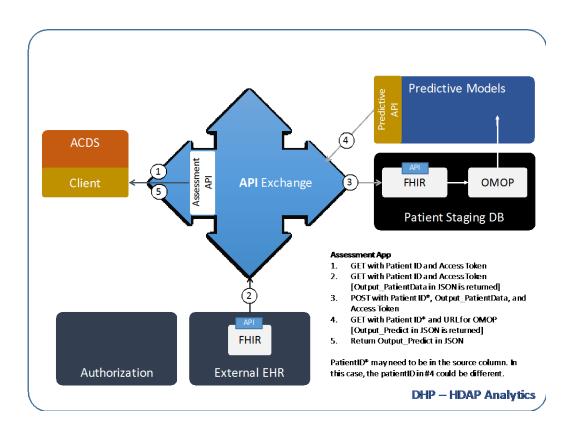
Fast Healthcare Interoperabilty Resources







Step Four: Connect to the World with APIs



Step Five: Build Applications

	has been performed for Byron Test1.				10.234.122.170 Change Password	Logout	
Filter out the following se Enzyme-Inducing D	Sec. 10.00 March 1	s Barbituates Reset			DUKE, JON DAVID at Fast MED Center	¥	
LEVETIRACETAM	93				Notifications	^ X	
TOPIRAMATE	74				Personalized Epilepsy Drug Ana	abusis	
LAMOTRIGINE	72.					S7 (2.0)	
ZONISAMIDE	69			Problems manage	What anti-epileptic drug will work best for Byron? The Eluminate Module provides a		
VALPROATE	21			Epilepsy (disorder)	personalized analysis of 15 common epilepsy medications and their likelihood of		
•		Show All		Epileptic seizure (finding)	success in controlling Byron's seizures.		
The probability of patient outcome improvement for modified treatment protocols list that you are viewing is intended to be one substantial reference point to be used in making.			Pain in lower limb (finding) Seizure (finding)				
decisions regarding treatment protocol changes. Physicians and those recommending and prescribing treatment regimen changes for epilepsy patients are naturally advised to weight this predictive analysis in concert with their own first-hand knowledge and / or all available information on the patient's comprehensive health in reaching any clinical decision on prescribed treatment.		d AX					
		Tinnitus (finding)	Relevant Orders Valproic Ac	0 00			
	Traumatic brain injury (disorder) 🛇	Valproate Level CBC2 Comprehensive Metabolic SGOT (AST) Show Order Menu					
			Chart Search	^ X			
			Enter search term	Search			
				Filter options			
					Labs, Meds, Reports T	J	
				Dx 8. Complaints Quantum Aug 23, 2016 17:50 — Visit Note Author: DUKE, JON DAVID (RESEARCH) Primary Care Dx Quantum Aug 23, 2016 17:50 — Visit Note Author: DUKE, JON DAVID (RESEARCH)			
				Visit Note Outpaient Aug 23, 2016 17:50			

Demonstration





Billy Bob Thornton

ZOCOR

AMIODARONE

CEFUROXIME

LEVAQUIN

CEFPODOXIME

LOSARTAN

LASIX

TOPAMAX

LISINOPRIL

Amiodarone



Dosage

BECAUSE OF THE UNIQUE
PHARMACOKINETIC PROPERTIES,
DIFFICULT DOSING SCHEDULE, AND
SEVERITY OF THE SIDE EFFECTS IF
PATIENTS ARE IMPROPERLY MONITORED,
AMIODARONE SHOULD BE ADMINISTERED
ONLY BY PHYSICIANS WHO ARE
EXPERIENCED IN THE TREATMENT OF

Interactions

Methotrimeprazine

Simvastatin

Vandetanib

cilostazol

Warnings

Amiodarone is intended for use only in patients with the indicated life-threatening arrhythmias because its use is accompanied by substantial toxicity.

Amiodarone has several potentially fatal toxicities, the most important of which is pulmonary toxicity (hypersensitivity pneumonitis or interstitial/alveolar

Journals

Interactions of digitalis and class-III antiarrhythmic drugs: Amiodarone versus dronedarone. International journal of cardiology, 2017 Feb 1

Acute hospital administration of amiodarone and/or lidocaine in shockable patients presenting with out-of-hospital cardiac arrest: A nationwide cohort study. International journal of cardiology, 2017 Jan 1

High-resolution sub-cellular imaging by correlative NanoSIMS and electron

News

With A-Fib Rhythms, Higher Odds of Stroke 2013-12-30

Muscle Aches From Statins? Drug Interactions May Play a Role 2013-12-04

Heart Warning Added to Label on Popular Antipsychotic Drug 2011-07-19

Fix a Health Problem or Live With It? 2009-08-17

Sanofi Drug Found Promising for Heart Ailment 2009-02-12

Huge Thanks To...

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Questions

