

The LEGEND Initiative

OHDSI Community Call Oct. 19, 2021 • 11 am ET

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Upcoming OHDSI Community Calls

Date	Topic
Oct. 19	Focus Topic: The LEGEND Project
Oct. 26	Trick or Treat
Nov. 2	Collaboration Opportunities: Methods Res., Data Standards, Open-Source, Clinical App.
Nov. 9	Demos: Tools for Adoption of OHDSI Data Standards
Nov. 16	Open Network Studies
Nov. 23	History of OHDSI
Nov. 30	Collaborator Showcase Presentations







Three Stages of The Journey

Where Have We Been?
Where Are We Now?
Where Are We Going?







Upcoming Workgroup Calls



Date	Time (ET)	Meeting			
Tuesday	1 pm	Common Data Model			
Tuesday	2 pm	Health Equity			
Tuesday	3 pm	OMOP CDM Oncology – Outreach/Research Subgroup			
Wednesday	9 am	Vaccine Vocabulary			
Wednesday	10 am	OMOP CDM Oncology – Development Subgroup			
Wednesday	1 pm	Data Quality Dashboard			
Wednesday	7 pm	Medical Imaging			
Thursday	12 pm	HADES			
Thursday	1 pm	OMOP CDM Oncology – CDM/Vocabulary Subgroup			
Friday	1 pm	1 pm Phenotype Development and Evaluation			
Monday	10 am	GIS-Geographic Information System			
Monday	11:30 am	Pharmacovigilance Evidence Investigation (PEI)			
Tuesday	9 am	OMOP CDM Oncology – Genomic Subgroup			

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Get Access To Different Teams/WGs/Chapters

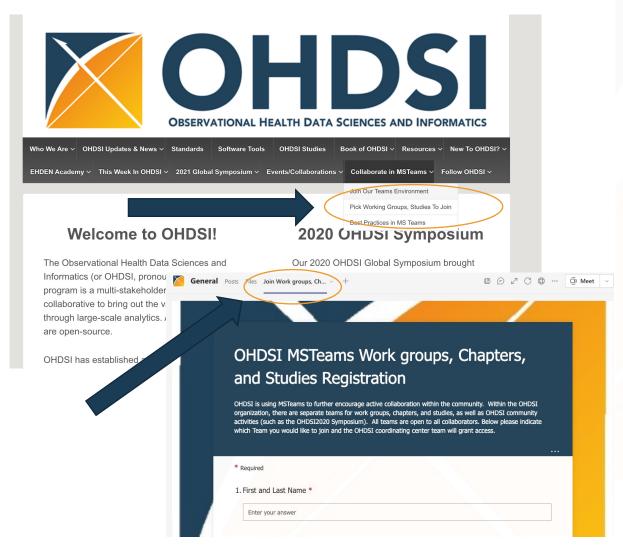


ATLAS		
Clinical Trials		
Common Data Model	Phenotype Development and Evaluation	
Data Quality Dashboard Development	Population-Level Effect Estimation / Patient-Level Prediction	
Early-stage Researchers	☐ Psychiatry	
	Registry (formerly UK Biobank)	
Education Work Group	Surgery and Perioperative Medicine	
Electronic Health Record (EHR) ETL	☐ Vaccine Safety	
Geographic Information System (GIS)	☐ Vaccine Vocabulary	
HADES Health Analytics Data-to-Evidence Suite	☐ Women of OHDSI	
Health Equity		
Latin America	6. Select the chapter(s) you want to join	
Laun America	Africa	
Medical Devices	Australia	
Natural Language Processing	China	
OHDSI APAC	Europe	
OTIDSI AFAC	Japan	
OHDSI APAC Steering Committee	☐ Korea	
OHDSI Steering Committee	Singapore	
Oncology	☐ Taiwan	
Patient-Generated Health Data		
Pharmacovigiliance Evidence Investigation	7. Select the studies you want to join	
Pharmacovigiliance Evidence Investigation	HERA-Health Equity Research Assessment	





Get Access To Different Teams/WGs/Chapters



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Oncology	Talwan	
Patient-Generated Health Data		
Pharmacovigiliance Evidence Investigation	7. Select the studies you want to join	





2021 APAC Symposium – Nov. 18

Nov 18 (APAC time zone)	Contents	Speaker(s)
Morning	OHDSI State of the Community	George Hripcsak/Patrick Ryan
	OHDSI APAC State of the Community	Mui Van Zandt
	EHDEN	Peter Rijnbeek
	FHIR and OHDSI Collaboration	Christian Reich
	APAC Chapter vision for 2022	APAC chapter leaders
Break		
Afternoon	Networking Session	All

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Next CBER Best Seminar Series

Topic

CBER BEST Initiative Seminar Series - Exploring Vaccine Safety Datalink COVID vaccine rapid cycle analysis (RCA) methods

Description

Background: The CBER BEST Initiative Seminar Series is designed to share and discuss recent research of relevance to ongoing and future surveillance activities of CBER regulated products, namely biologics. The series focuses on safety and effectiveness of biologics including vaccines, blood components, blood-derived products, tissues and advanced therapies. The seminars will provide information on characteristics of biologics, required infrastructure, study designs, and analytic methods utilized for pharmacovigilance and pharmacoepidemiologic studies of biologics. They will also cover information regarding potential data sources, informatics challenges and requirements, utilization of real-world data and evidence, and risk-benefit analysis for biologic products. The length of each session may vary, and the presenters will be invited from outside FDA. Please see the details below for our upcoming seminar. Anyone can register and join for free. Stay tuned for more details and additional webinars during the year.

Topic: Exploring Vaccine Safety Datalink COVID vaccine rapid cycle analysis (RCA) methods

Description: We will review statistical methods used in observational studies of the safety and effectiveness of COVID-19 vaccines. Topics will include:

- How to compare recent vaccinees with concurrent comparators (unvaccinated or less recently vaccinated) and
- with comparators who are not concurrent (historical rates or self-controls) to make inferences about outcome rates
- that would be expected among vaccinees had they not been vaccinated
- Methods for estimating risk ratios
- How to examine change in vaccine effectiveness (waning) or vaccine safety over timesince-vaccination
- Seguential tests

Presenter: Nicola P. Klein, MD, PhD

Time Oct 20, 2021 11:00 AM in Eastern Time (US and Canada)



Webinar Registration



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Prediction of early acute readmission after colorectal cancer surgery using only clinical preoperative variables.





PRESENTER: Johan Clausen

Early unplanned readmission following colorectal cancer surgery is a significant economic burden to the health care system and may delay patient recovery and adjuvant chemotherapy onset. Identifying patients at high risk of readmission when planning the surgical and oncological treatment is of high value, as preoperative training or extensive postoperative monitoring can be planned.

A CDM was built using data from the Danish colorectal cancer group's nationwide database (DCCG), containing clinical data from all colorectal cancer surgeries since 2001. DCCG-Data was enriched with information about readmission from the Danish National Patient

OHDSI's ATLAS tool and R was used to build a patient-level prediction model with acute

readmission as outcome. The target cohort was colorectal cancer patients undergoing surgery and time-at-risk was date of surgery until 30 days after. Covariates in the age, gender, measurement values, conditions, procedures and observation domain available any time prior to surgery was included in the model. Custom covariates were constructed for specific clinical scales (e.g., ASA score).





- 62.824 patients underwent colorectal cancer surgery between 2001 to 2019. The incidence of unplanned 30-day readmission was 10.423 (16.6%).
- · 96 variables were included in the model. Using only preoperative available variables, the
- prediction model had a AUC of 0.60 (95%CI 0.59:0.61) and an AUPRC of 0.22. Calibration was considered acceptable with a brier score

Preoperative clinical variables can predict early acute readmission after colorectal cancer surgery.



Accurate prediction of acute readmission may assist the multidisciplinary team in the decision-making of the patients' treatment trajectory.

CLINICAL USE OF THE PREDICTION

MODEL

The patients' treatment trajectories are often planned at a multidisciplinary team conference (MDT). Identifying natients with high risk of postoperative morbidity is crucial at MDT, and a decision support tool visualizing personalized readmission risk may prove to be of high value for identifying high-risk patients and deciding the

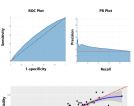
Interventions on high-risk patients may include Delay of surgery and preoperative training

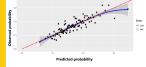
 Increased postoperative monitoring or delayed discharge

The prediction model can not be used as a stand-alone tool in the MDT-setting but models may be useful for assisting the clinicians in the decision-making process.

PERSPECTIVES

Enriching the CDM with further phenomics from other nationwide data sources may improve the performance of the prediction model significantly







MONDAY

Prediction of early acute readmission after colorectal cancer surgery using only clinical preoperative variables Authors: Johan Clausen, Andreas Weinberger Rosen, Karoline Bendix Bräuner, Mikail Gögenur, Viviane Annabelle Lin, Eldar Allakhverdiiev, Julie Sparholt Walbech, Ismail Gögenur











Detecting PTSD and self-harm among US Veterans using positive unlabeled learning

Christophe G. Lambert, PhD

Center for Global Health, Division of Translational Informatics, Department of Internal Medicine, University of New Mexico Health Sciences Center,

Albuquerque, New Mexico, USA

Co-Authors

Praveen Kumar; Nicolas R. Lauve; Sharon E. Davis; Sharidan K. Parr; Daniel Park; Michael E. Matheny; Gerardo Villarreal; George Uhl; Yiliang Zhu; Mauricio Tohen; Douglas J. Perkins; Christophe G. Lambert

TUESDAY

Detecting PTSD and self-harm among US Veterans using positive unlabeled Learning Authors: Praveen Kumar, Nicolas R. Lauve, Sharon E. Davis, Sharidan K. Parr, Daniel Park, Michael E. Matheny, Gerardo Villarreal, George Uhl, Yiliang Zhu, Mauricio Tohen, Douglas J. Perkins, Christophe G. Lambert (presenter)



TreatmentPatterns: An R package to analyze treatment patterns of a study population of interest

△ PRESENTER: Aniek Markus

INTRO

· There is no R package available to analyze treatment patterns of a study population of interest and the standard analytics tool available in ATLAS has the disadvantage that it is not customizable to specific research needs.

- · We defined the process of constructing pathways following earlier work, highlighting key decisions in the process that need to be made (see Figure 1).
- · We demonstrate the functionalities of the package and outputs by analyzing treatment patterns of three common chronic diseases (type 2 diabetes, hypertension, and depression) in the **Dutch Integrated Primary Care** Information (IPCI) database (see online Shiny application).

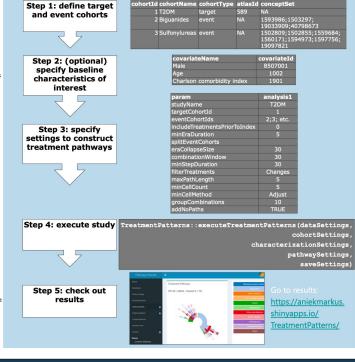
RESULTS

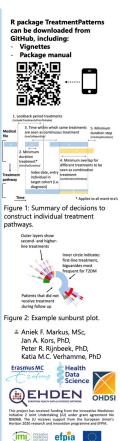
- · The R package TreatmentPatterns creates sunburst plots (see Figure 2), Sankey diagrams, and various other outputs (e.g. percentage of people treated, average duration of event cohorts) to give insight in first-, secondand higher line treatments
- · The results can be explored in an interactive Shiny application: https://aniekmarkus.shinvapps.io **TreatmentPatterns**

CONCLUSION

- · This tool is intended to make the analysis of treatment patterns more accessible, more standardized, and more interpretation friendly
- · We hope it thereby contributes to the accumulation of knowledge on real-

How to perform a treatment patterns study in 5 steps:





WEDNESDAY

Treatment Patterns: An R package to analyze treatment patterns of a study population of interest

Authors: Aniek F. Markus, Peter R. Rijnbeek, Jan A. Kors, Katia Verhamme



Higher-Level Radiation Oncology Treatment Events derived from Lower-Level CPT Codes

PRESENTER: Michael Gurley¹, Asieh Golozar¹, Rimma Belenkaya¹, Tatyana Sandler¹

1 OHDSI Oncology Workgroup

INTRO:

data for radiation oncology treatment. In OMOP, a radiation oncology treatment can be comprised of 72 lower-level clinical patient entries in the PROCEDURE_OCCURRENCE table across 11 CPT codes. Through an aggregation to a higher-level event that is one record in the EPISODE table, several use cases can be addressed such as, more intuitive treatment representation to an oncology professional and data more amenable to analytics. For the representation of higher-level events, the level of granularity of modality and technique recommended by the American Society for Radiation Oncology (ASTRO) in the minimum data elements for radiation oncology consensus paper was assessed against what can be derived and aggregated

CPT codes are a common structured source of patient

from CPT codes.

- 1. Curate relevant CPT codes
- Reconcile past efforts of modality and techniques assignments to the lower-level CPT codes²³⁴.
- Identified ASTRO's minimum relevant defined data standards for modality and technique of a radiation oncology treatment⁵.
- Assessed CPT codes assignments of higher-level treatment in step 2 against the modalities and techniques defined by ASTRO in step 3 to determine extent of coverage using CPT codes.

RESULTS

Overall, the results show that CPT codes support a small subset of modalities and techniques represented in ASTRO's data elements. For a larger set, there are varying levels of details.

To use ASTRO's data elements for higher-level event representation, expanding the value sets of modality and technique to cover use cases with less information available about a treatment event. For example, when there is a CPT code for an external beam radiotherapy" without additional details, addition of a higher-level event to capture "external beam radiotherapy" is needed.

There are **challenges** in **deriving** higher-level

Radiation Oncology

treatment

Events from

CPT codes
with the level
of detail

recommended

by **ASTRO**.

Radiation oncology procedure	Derivable from CPT		
30-1.38-00-0-1.0-000-00-0.0-0.0-0.0-0.0-0.0-0.0	Yes	Partial	No
Modality	375		
External beam radiation therapy (EBRT)			
Protons	X		
Electrons			X
Photons (LINAC)		X	
Photons (isotope source)		X	
Neutrons	X		
Carbon			X
Brachytherapy			
Low dose rate	×		
High dose rate	X		
Pulse dose rate			X
Radiopharmaceuticals	X		
Electronic brachytherapy	×		
kV x-rays			
Intraoperative radiation therapy			X
Superficial		X	
Orthovoltage		X	
Technique		7	**
Passive scattering			×
Scanning beam intensity modulated proton therapy			×
Scanning beam multi-field optimization			X
Scanning beam single-field optimization			×
2-dimensional (2D)			×
Intraoperative radiation therapy	X		
3-dimensional (3D)	×		
Intensity modulated radiation therapy (IMRT)	X		
Intracranial stereotactic	X		
Interstitial permanent		X	
Interstitial temporary		×	×
Intracavitary permanent		X	
Intracavitary temporary		X	
Sealed			×
Unsealed			X
Intracavitary	X		



References/Citations

. Gurley Michael, Belenkaya Rimma Extending OMOP CDM to Support Observational Cancer Research OHDSI.2018; 2. Cancer Therapy Lookup Tables -Cancer Research Network (CRN) CRN. March 2018; 3. Daily Practice Reimbursement Practice Management Resources - Basics of RO Coding -American Society for Radiation Oncology (ASTRO) American Society for Radiation Oncology (ASTRO) ASTRO 2016: 4. CanMED and the Oncology Toolbox. . James A. Hayman, Andre Dekker, Mary Feng, Randi Kudner Samantha Dawes, James B. Yu. Minimum Data Elements for Radiation Oncology An American Society for Radiation Oncology Consensus Paper. Practical Radiation Oncology ASTRO 2019 VOLUME 9;ISSUE 6;P395-401

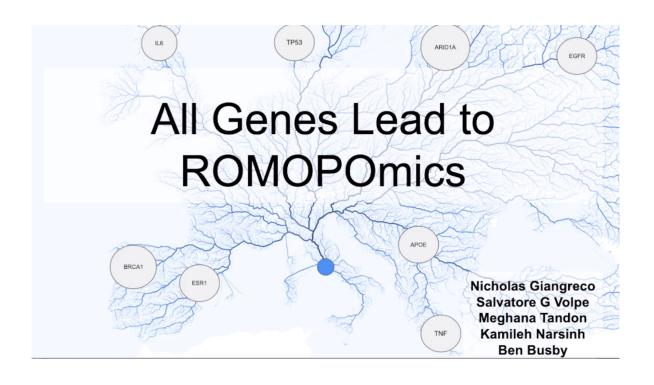
THURSDAY

Representation of High-Level Radiation Oncology Treatment Events from CPT Codes Authors: Michael Gurley, Asieh Golozar, Rimma Belenkaya, Tatyana Sandler



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FRIDAY

All Genes Lead to ROMOPomics
Authors: Nicholas Giangreco, Salvatore G Volpe, Meghana Tandon, Kamileh
Narsinh, Ben Busby







Where Are We Going?

Any other announcements of upcoming work, events, deadlines, etc?

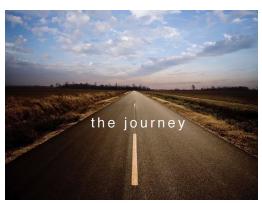






Three Stages of The Journey

Where Have We Been? Where Are We Now? Where Are We Going?







Oct. 19 Community Call: The LEGEND Initiative



Fan Bu



RuiJun "Ray" Chen



Rohan Khera



Yuan Lu



Anna Ostropolets



Aki Nishimura



Marc Suchard

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