

Rapid Fire Presentations of Collaborators

Katia Verhamme, MD, PhD
Associate Professor of Use and Analysis of
Observational Data



FEEDERNET (FEDERATED E-HEALTH BIG DATA FOR EVIDENCE RENOVATION NETWORK) PLATFORM IN KOREA

Seongwon Lee, Chungsoo Kim, Junhyuk Chang, Rae Woong Park

FeederNet (Federated E-Health Big Data for Evidence Renovation Network) platform in Korea

2022.06.24

Seongwon Lee, Chungsoo Kim, Junhyuk Chang, Rae Woong Park

Department of Biomedical Informatics, Ajou University School of Medicine, Suwon, Korea Department of Biomedical Sciences, Ajou University Graduate School of Medicine, Suwon, Korea

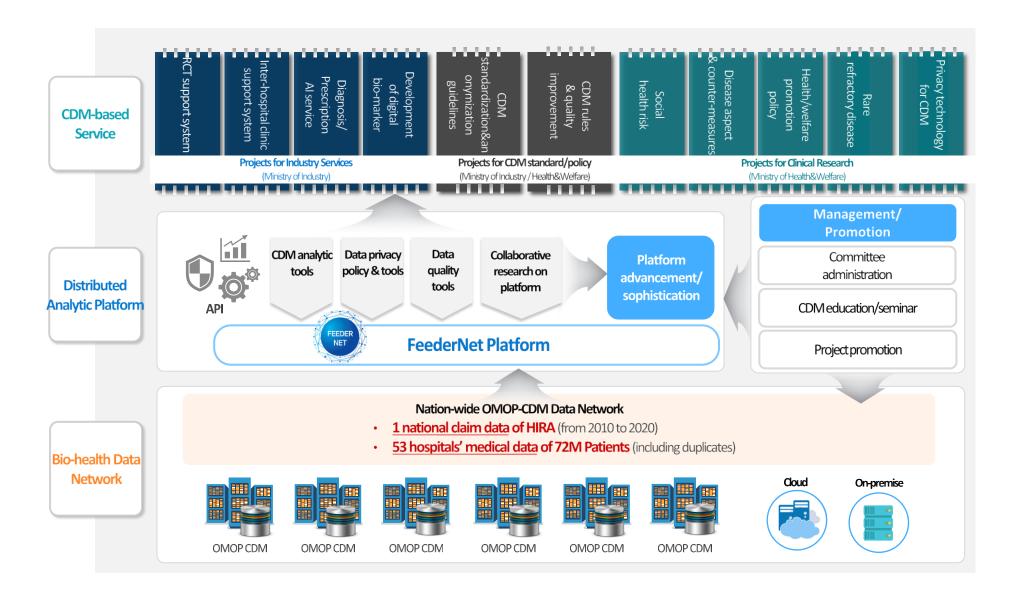
* Presenting author: Chungsoo Kim, Pharm D







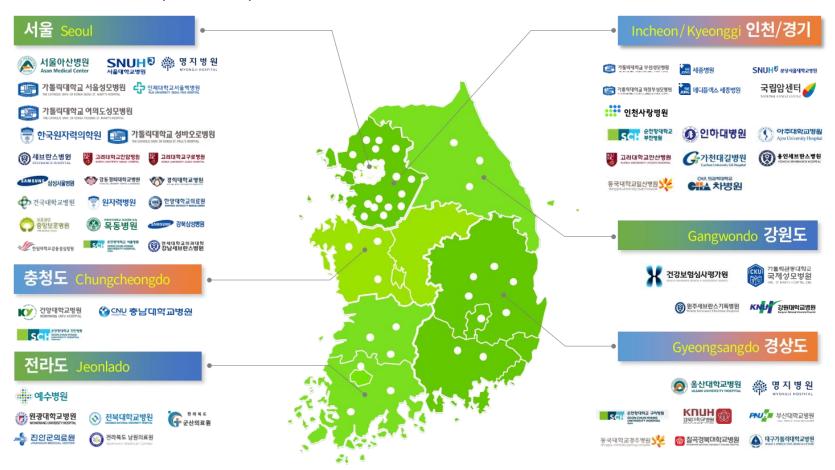
FeederNet Project in Korea



OMOP-CDM Data Network

Nation-wide CDM Data Network

- 53 hospitals (including 72% of all tertiary hospitals in Korea)
- 71,987,327 patients (including duplicates)
- X National claim data (2010-2020) of All Korean was also converted to OMOP-CDM



FeederNet Platform

FeederNet Platform

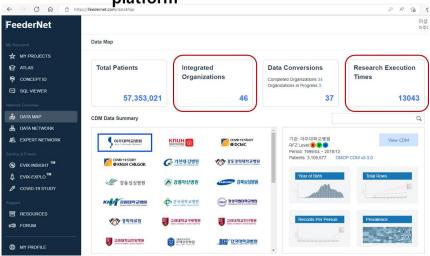
- 46 CDM DBs of hospitals have been integrated with FeederNet platform
- FeederNet Central + FeederNet Node
- Since May 2019, total 13,043 analyses have been executed

Research Free Zone

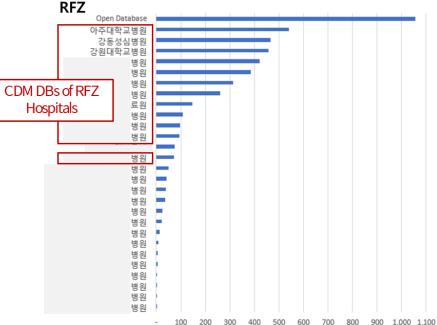
- Mutual Cooperative MOU for promoting joint research
- Clauses of RFZ
 - 1. Grant same authorities on CDM DB analysis to all researchers in RFZ hospitals
 - 2. Approve single IRB among RFZ hospitals

Currently, 18 Hospitals joined the RFZ

Main page of FeederNet platform



Comparison of CDM usage between RFZ and non-



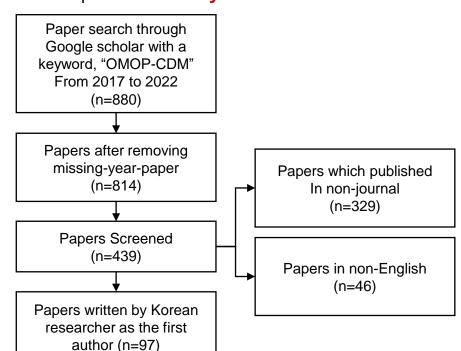
OMOP-CDM Research in Korea

Analysis on FeederNet

- Number of analyses continues to increase
- From June 2020, about 500 analyses have been being conducted every month

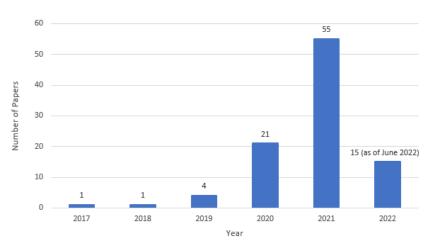
Paper Publications

 Total 97 papers of Korean researcher as a first author have been published since 2017 and 55 were published in only 2021



Cumulative number of analysis using FeederNet 14,000 10,000 8,000 4,000 2,000 10,000 2,000 10,

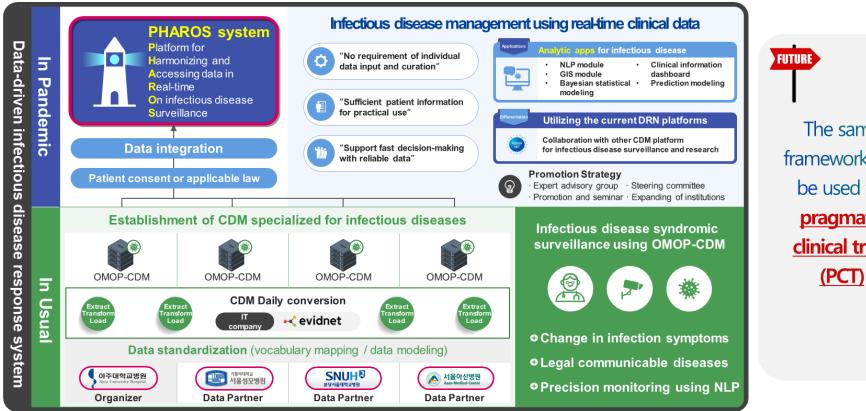
OMOP-CDM research status by Korean researchers



New CDM project on Infectious Disease Surveillance

'PHAROS' Project

- PHAROS (Platform for Harmonizing and Accessing data in Real-time On Infectious disease Surveillance)
 - **Purpose**: Establishment of infectious disease surveillance system using real-time CDM data of multi centers
 - Uniqueness
 - 1. Daily automatic CDM ETL
 - **2. Patient-centered Integration** of CDMs from multi-centers





Thank you



OMOP GENOMIC MAPPING CAPACITIES IN CONVERSION OF COMPREHENSIVE GENOMIC PROFILING RESULTS

Rogozhkina Maria, Odysseus

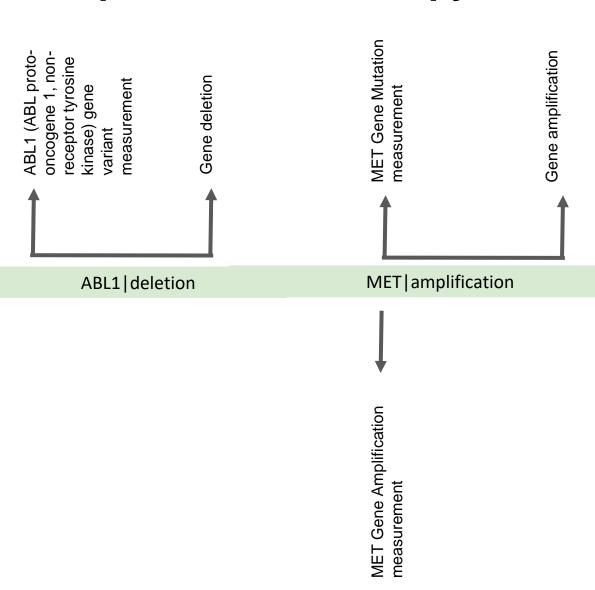
OMOP Genomic mapping capacities in conversion of comprehensive genomic profiling results

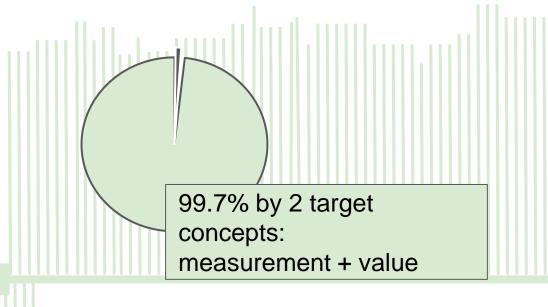


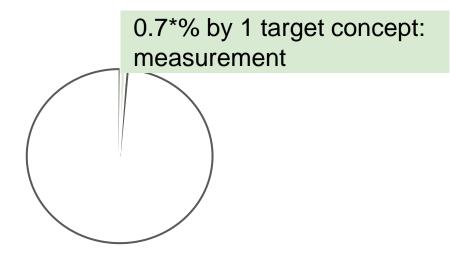


Representation of Copy Number aberrations





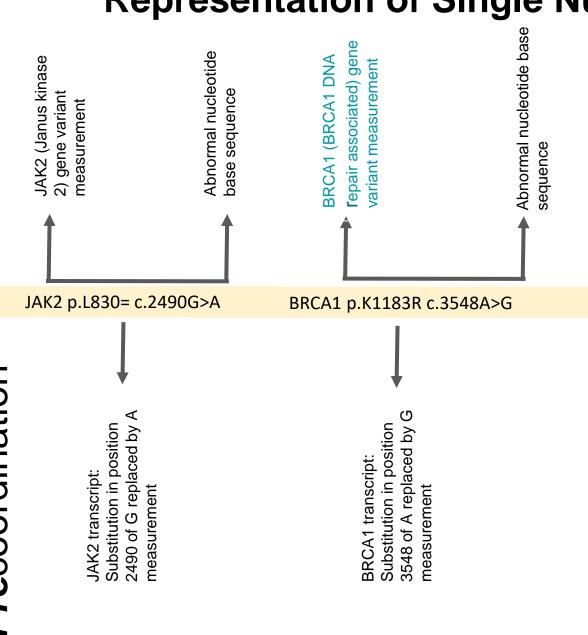


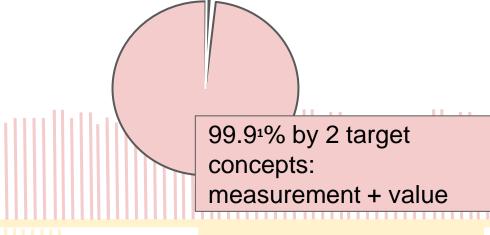


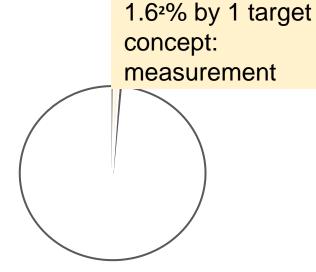
*All 5 codes for amplifications only

Representation of Single Nucleotide Variants





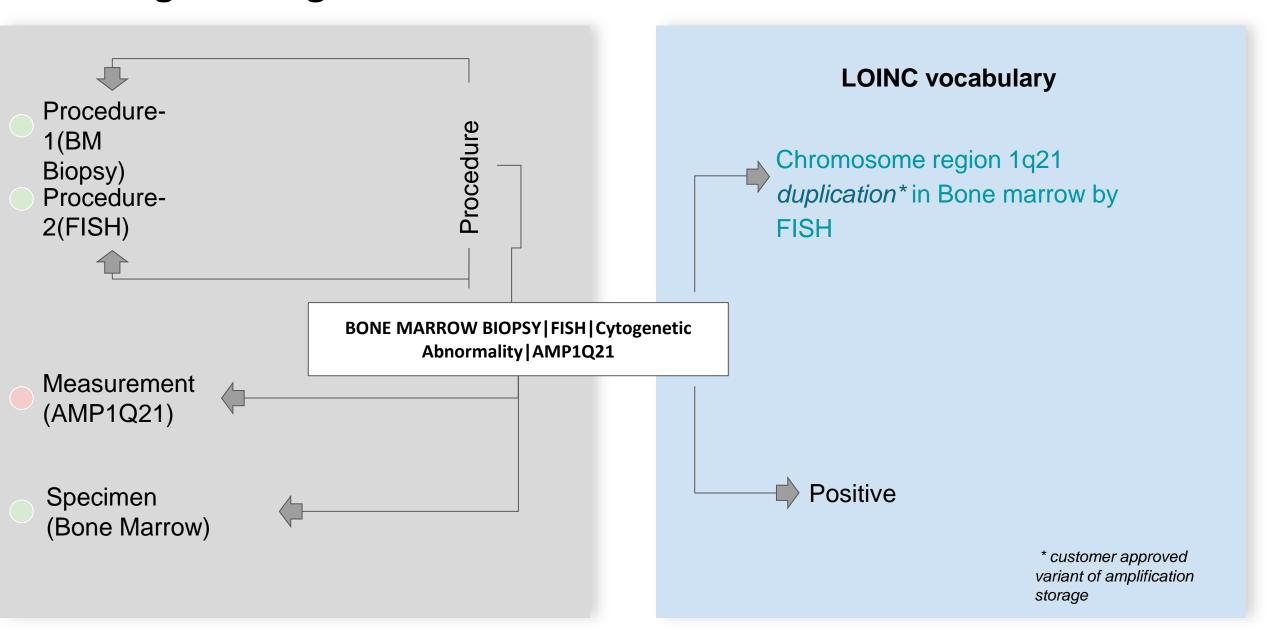




- 1- data about position is missing in all target concepts
- ² 0.3% was mapped to Protein level



Using existing vocabularies as a fallback



Conclusion



- We found <u>46</u> duplicate concepts for DNA, <u>1276</u> for RNA and <u>224</u> for Protein variants in OMOP Genomic. All of them to be deduplicated.
- A substantial number of codes are missing. **New codes** to be ingested in OMOP Genomics are essential to facilitate studies.
- Concepts from non-Genomic vocabularies may be a valid target, but lacks of consistency and only a fraction of required targets is available.
- Linking to other event tables such as Procedure and Specimen is required to properly
 represent source data. To efficiently run queries an easier way to link facts is required
 (direct fact modification)



OMOP MAPPING OF REAL-WORLD DATA FROM BRAZIL & PAKISTAN TOWARDS MANAGEMENT OF COVID-19 IN THE GLOBAL SOUTH

Sara Khalid, University of Oxford













OMOP Mapping of Real-World Data From Brazil & Pakistan Towards Management of COVID-19 In the Global South

Authors

Elzo Pereira Pinto Junior¹, Priscilla Normando¹, Renzo Flores-Ortiz¹, Muhammad Usman Afzal², Muhammad Asaad Jamil², Sergio Fernandez Bertolin³, Vinícius de Araújo Oliveira¹, Valentina Martufi¹, Edward Burn⁴, Maria Yury Ichihara¹, Maurício L. Barreto¹, Talita Duarte Salles³, Daniel Prieto-Alhambra⁴, Haroon Hafeez², Sara Khalid⁴

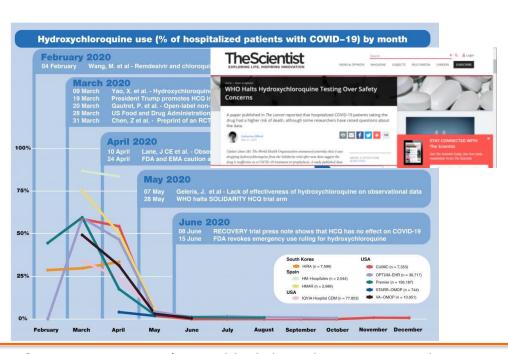
- ¹ The Center for Data and Knowledge Integration for Health (CIDACS FIOCRUZ), Brazil
- ² Shaukat Khanum Memorial Cancer Hospital and Research (SKMHR&C), Pakistan
- ³ The Information System for Research in Primary Care (SIDIAP), Spain
- ⁴ Centre for Statistics in Medicine, University of Oxford, UK.

Background





The COVID-19 pandemic highlighted need for rapid, reliable, **representative** evidence generation



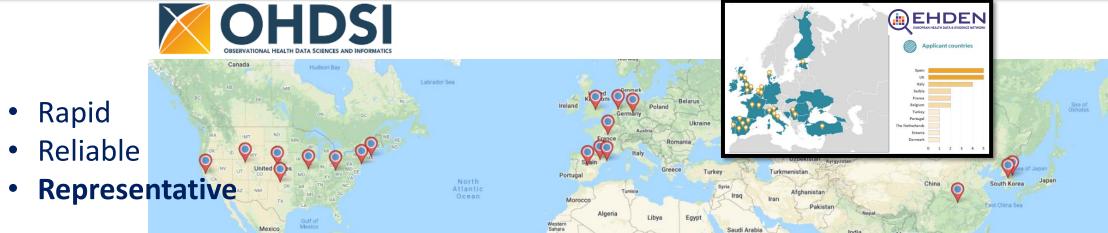


The OHDSI COVID-19 Data Network

Rapid







Cuba	Saudi Arabia	India Myanmar
USA (8)	EUROPE (7)	ASIA-PACIFIC (3)
Premier (National – Hospital Billing)	CPRD (UK – Electronic Health Records)	HIRA (South Korea – Administrative Claims)
HealthVerity (Claims linked to diagnostic testing)	SIDIAP (Spain – Electronic Health Records)	DCMC (South Korea – Electronic Health Records)
Optum EHR (National – Electronic Health Records)	SIDIAP-H (Spain – EHR hospital linkage	Nanfang Hospital (China – Electronic Medical Records)
IQVIA Open Claims (National – Administrative Claims)	HM Hospitales (Spain – Hospital Billing)	Together, OHDSI has studied (to date): • >4.5m patients tested for SAR-COV- 2 • >1.2m patients diagnosed or tested positive for COVID-19 • >250k hospitalized for COVID-19
Department of Veterans Affairs (National – Electronic Health Records)	ICPI (Netherlands – Electronic Health Records)	
Stanford University (CA – Electronic Health Records)	LPD France (France – Electronic Health Records)	
Tufts University (MA – Electronic Health Records)	Germany DA (Germany – Electronic Health Records)	
Columbia University (NY – Electronic Health Records)		·

The OHDSI COVID-19 Data Network









Data Partners – Pakistan





- Data source:
 - De-identified electronic health records
- Period:
 - 1994 2022 (ongoing)
- Unique records:
 - 8.3 million individuals
- Regional COVID-19 hub





Country Facts

Population: 220 million

5 Provinces

Clinical activity snapshot (2021)

















Data Partners – Brazil



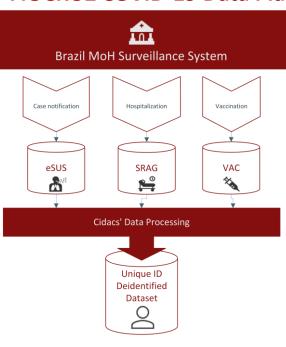


Data source/s:

- Brazil Ministry of Health Influenza Surveillance System (SIVEP-Gripe)
- Period
 - 2020 2022 (ongoing)
- COVID-19 records:
 - 2.6 million individuals
 - ~67,000 hospitalisations
 - ~27,000 deaths

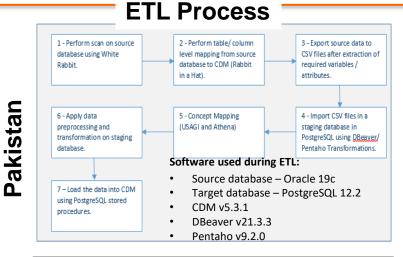


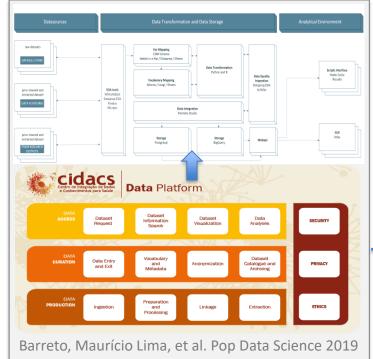
CIDACS-FIOCRUZ COVID-19 Data Platform



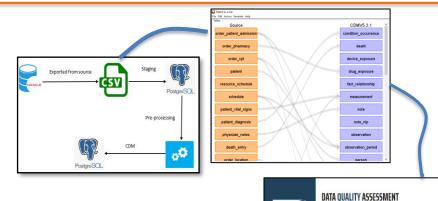
Harmonisation to OMOP





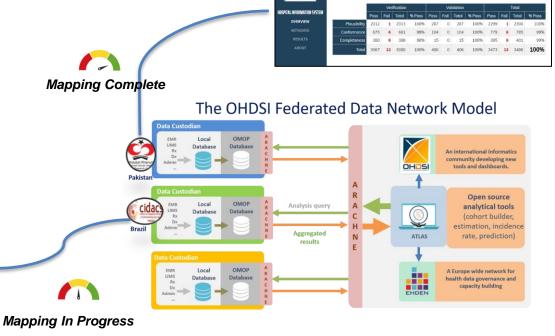


Brazil



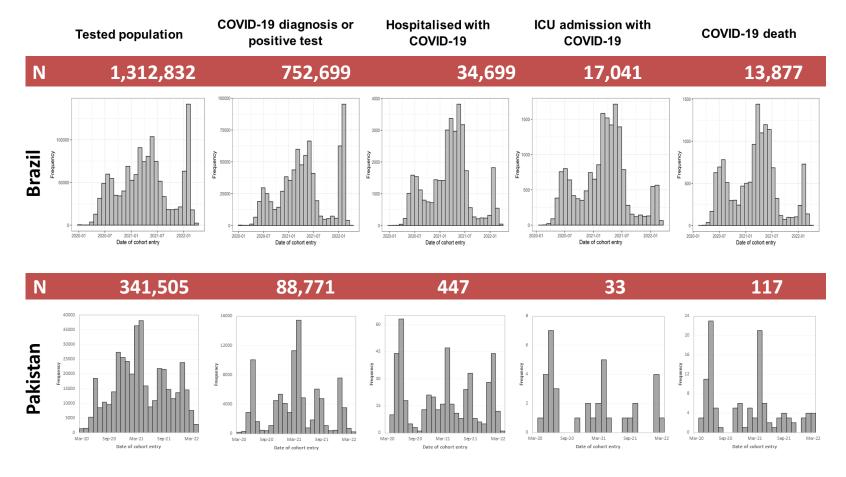
Mapping in Numbers

- >100K (source) to 108K (CDM) concepts
- >33M measurements
- >2M procedures
- >600K observations
- <1% missing matching concepts



COVID-19 cohorts – cases over time



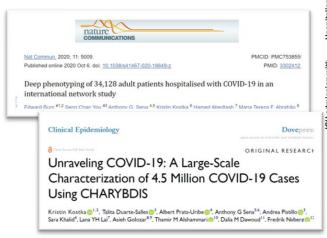


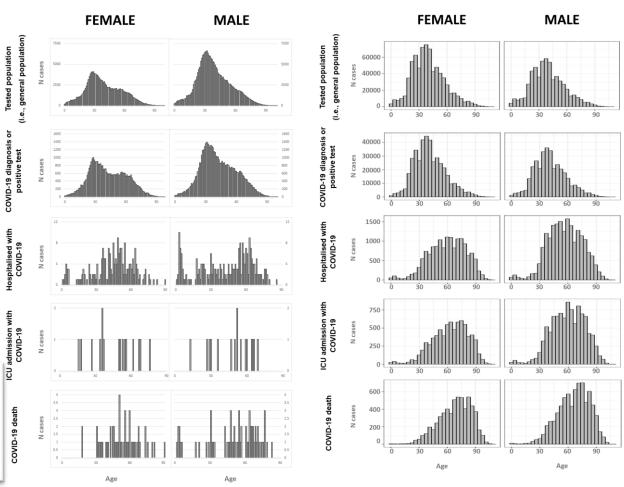
Distribution of cases over time (Jan/March 2020 – April 2022)

COVID-19 cohorts – baseline characteristics



COVID-19
 outcomes were
 more severe in
 men, elderly,
 and those with
 co-morbidities

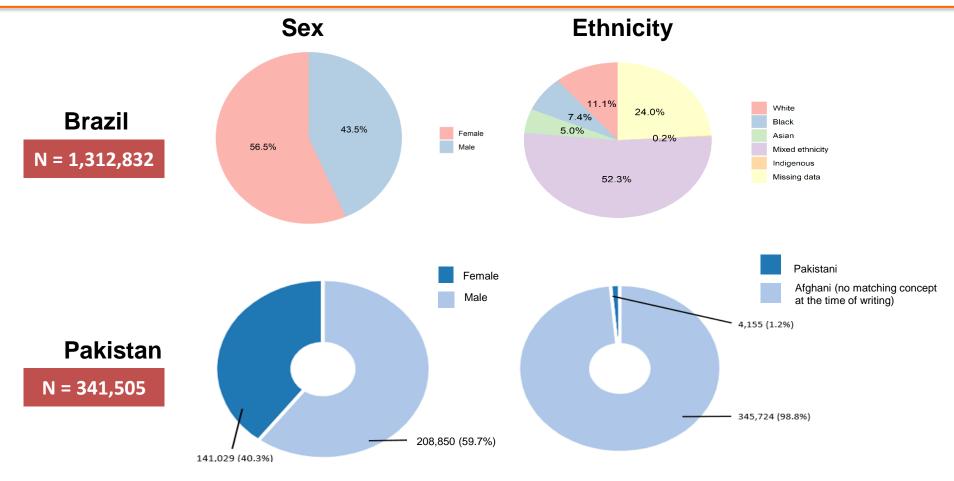




Distribution of cases by age and sex in each cohort

COVID-19 cohorts – baseline characteristics





General population tested for COVID-19:

- Age: median (IQR) was 36 (25 -75) and 38 (27 50) for Pakistan and Brazil
- Sex: 45.5% and 55% were female in Pakistan and Brazil
- Ethnicity/race: 1.2% Pakistan individuals had "Afghan" ethnicity. In Brazil, 52.3% had "Mixed" ethnicity.

Summary



- Two health databases covering 8.3 million people from Pakistan and 2.6 million people from Bahia, Brazil were analysed.
- 109,504 (Pakistan) and 921 (Brazil) medical concepts were harmonised to OMOP CDM.
- 341,505 (4.1%) people in the Pakistan dataset and 1,312,832 (49.2%) people in the Brazilian dataset tested for COVID-19 between 1st Jan 2020 and 30 April 2022.
- In agreement with international findings, COVID-19 outcomes were more severe in men, elderly, and those with underlying health conditions.
- This proof-of-concept study demonstrates potential for OMOP-harmonised data from underrepresented regions for global knowledge mobilisation and clinical translation for timely response to healthcare needs in pandemics and beyond.

What's next





- COVID-19 variant and vaccine surveillance study
- Communicable, NCDs
 - Cancer (OHDSI Oncology WG)
- Environment, equity, and artificial intelligence
 - Social deprivation dashboard (OHDSI GIS WG, OHDSI Equity WG)
- Data science ecosystem
 - Capacity building
 - Data re-use projects
 - Data governance















Acknowledgements

This work was supported by funding from the Bill & Melinda Gates Foundation















































































OMOP Mapping of Real-World Data From Brazil & Pakistan Towards Management of COVID-19 In the Global South

Authors

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- ⁴ Centre for Statistics in Medicine, University of Oxford, UK.



IMPACT OF RANDOM OVERSAMPLING AND RANDOM UNDERSAMPLING ON THE DEVELOPMENT AND VALIDATION OF PREDICTION MODELS USING OBSERVATIONAL HEALTH DATA

Cynthia Yang, Erasmus MC

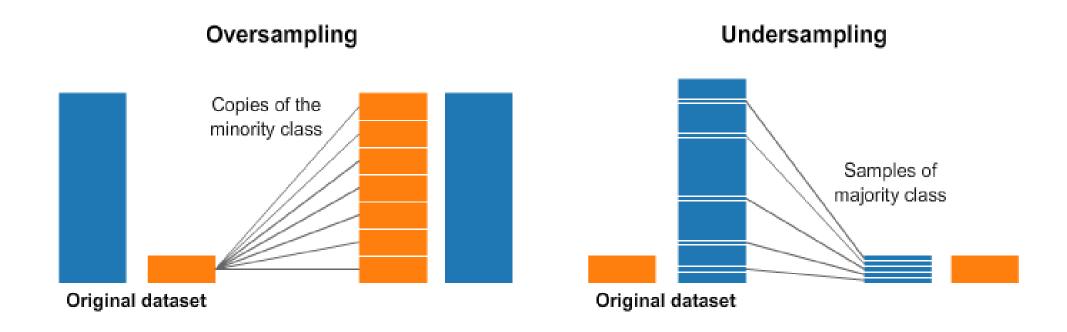
Impact of random oversampling and random undersampling on the development and validation of prediction models using observational health data

Cynthia YangPhD Student at Erasmus MC



The class imbalance problem

Class imbalance: a small proportion of patients in a study population (minority class)
experiences a certain outcome of interest.



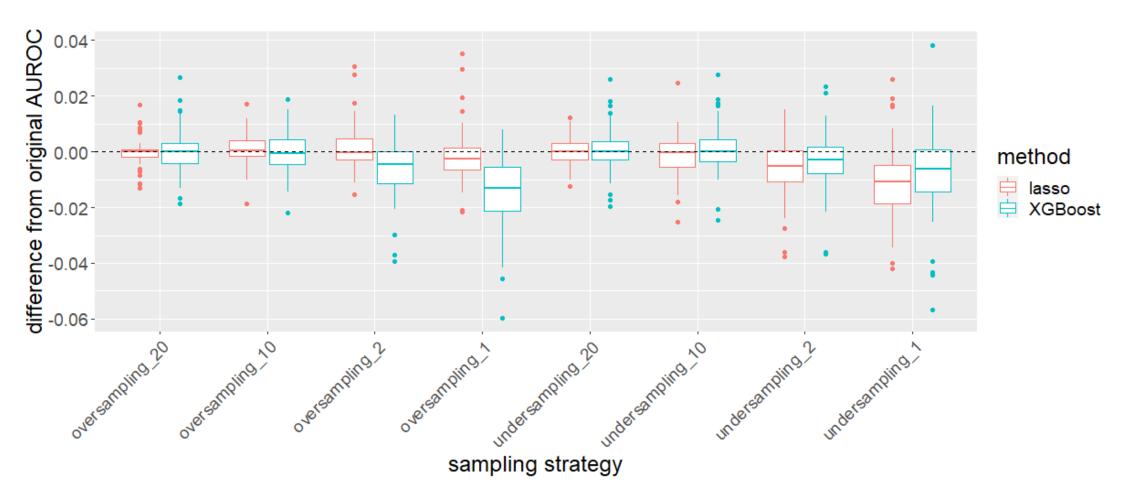


Methods

- Random oversampling and random undersampling
- Imbalance ratio (IR) = the total number of patients without an outcome event (majority class) / the total number of patients with an outcome event (minority class)
- We vary $IR = min(IR_{original}, x)$ with $x \in \{20, 10, 2, 1\}$
- 21 outcomes in depression (PLP framework paper)
- CCAE, MDCD, MDCR, IQVIA Germany
- XGBoost, lasso logistic regression
- Area under the receiver operating characteristic curve (AUROC)



Results





Conclusions

- The impact of random sampling on the AUROC is limited.
- Random sampling strategies on average do not improve the AUROC.
- Next steps: model calibration, random forest, external validation.
- For more information or questions please visit me at my poster #57.







A SUMMARY OF 'LIVE' REQUESTS FOR RWE STUDIES PUBLISHED BY A EUROPEAN HEALTH TECHNOLOGY ASSESSMENT (HTA) AGENCY

Jamie Elvidge, National Institute for Health and
Care Excellence
(NICE)

Real-world evidence is in demand!

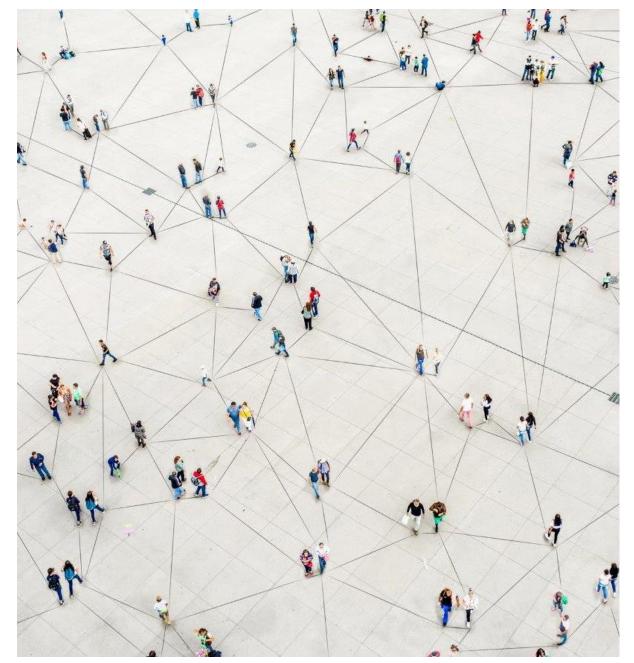
Live requests for real-world evidence (RWE) studies from a health technology assessment (HTA) agency

Jamie Elvidge

jamie.elvidge@nice.org.uk

Ravinder Claire, Shane Collins, Dalia Dawoud

NICE National Institute for Health and Care Excellence



About NICE

- The HTA agency for England
- Provides guidance to the healthcare system
- Key principles:
 - evidence-based medicine
 - opportunity cost





Actively exploring ways to use RWE for decision making

NICE real-world evidence framework

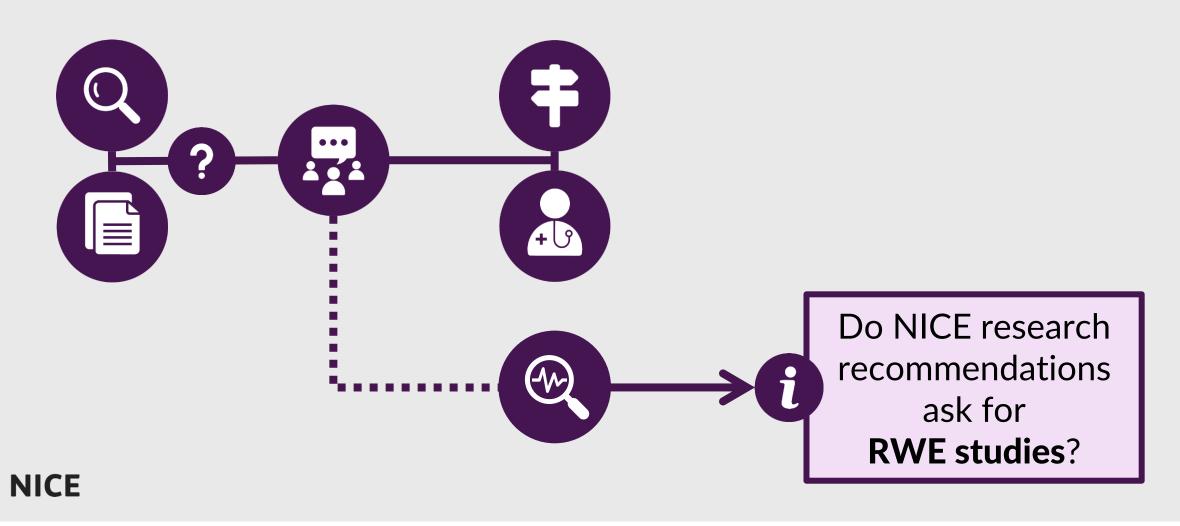
Overview

Key messages

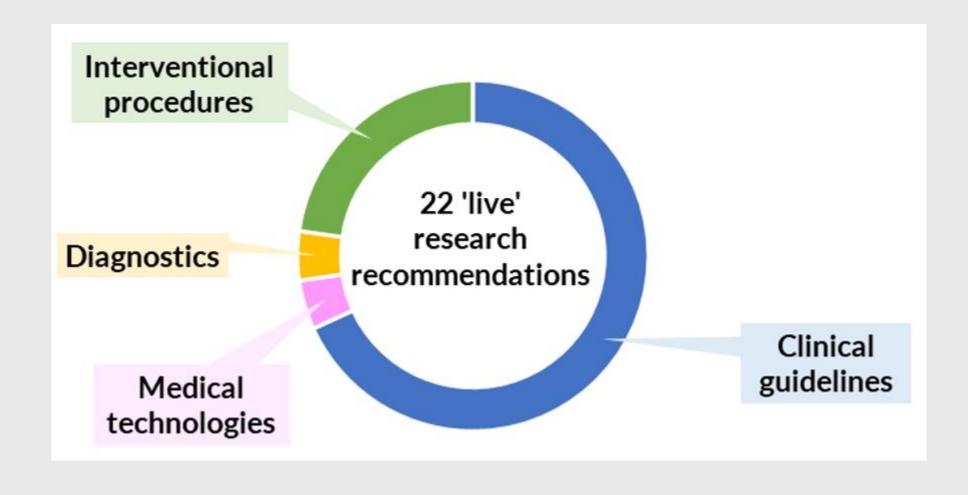
- The <u>NICE Strategy 2021 to 2026</u> states our ambition to use real-world data to resolve gaps in knowledge and drive forward access to innovations for patients. Real-world data is essential to enabling rapid, robust, and responsive technology evaluations and dynamic, living guidelines.
- We developed the Real-World Evidence Framework to help deliver on this ambition. It does this by:
 - Identifying when real-world data can be used to reduce uncertainties and improve

NICE

NICE recommends research to fill evidence gaps identified during guidance development

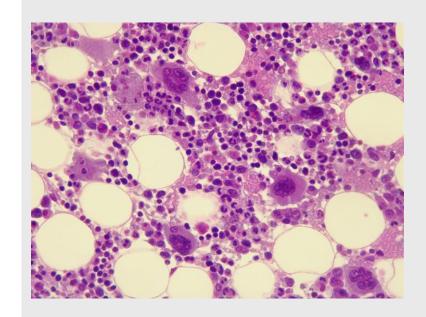


There are calls for RWE from across NICE guidance



NICE





Ophthalmology

Routine treatment can be effective, but also unpleasant, costly and risky.

→ Observational studies to characterise treatment benefit and predict when the benefits cease.





Delirium

Observational studies to characterise prevalence of delirium in long-term care settings, and explore whether delirium is predictive of adverse outcomes and death.

Lung cancer

Large retrospective analyses to predict determinants of lung cancer, characterise current practice, and predict likelihood of unnecessary surgery.





WHY PREDICTING RISK CAN'T IDENTIFY 'RISK FACTORS' EMPIRICAL ASSESSMENT OF MODEL STABILITY IN MACHINE LEARNING ACROSS OBSERVATIONAL HEALTH DATABASES

Aniek Markus, Erasmus MC

WHY PREDICTING RISK CAN'T IDENTIFY 'RISK FACTORS'

EMPIRICAL ASSESSMENT OF MODEL STABILITY IN MACHINE LEARNING ACROSS OBSERVATIONAL HEALTH DATABASES

CO-AUTHORS: PETER R. RIJNBEEK, JENNA M. REPS

Aniek Markus

PhD Student
Department of Medical Informatics, Erasmus MC
a.markus@erasmusmc.nl







Motivation

Some researchers incorrectly interpret prediction models:

1. Prediction models do not assess causality

2. Also problematic to use models for 'risk factor' detection

Why?

Procedures, medical conditions, drugs often co-occur

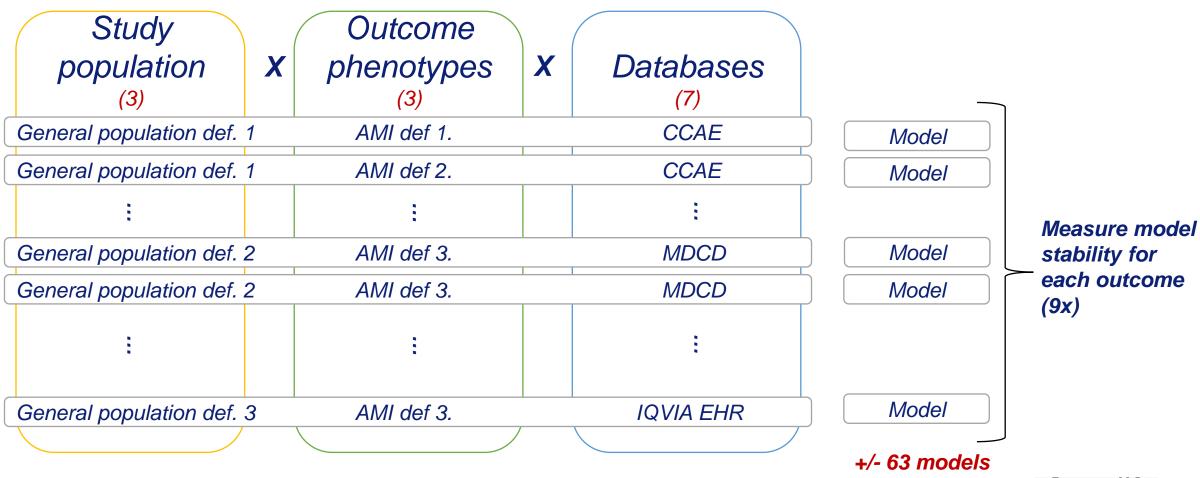
LASSO logistic regression might ignore some of them

'Variable
associated to
the outcome'





Study design



Erasmus MC z afuns

Measuring model stability

Example model:

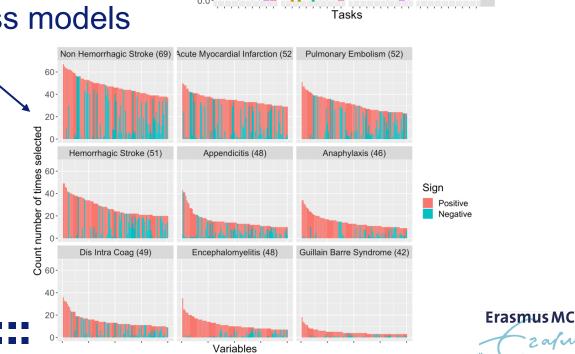
```
probability(acute myocardial infarction within 1 year) = \sigma(+1.33 * age - 0.2 female + 0.43 obesity)
```

- 1. How many variables are selected across models?
- 2. Are the same or different variables included across models?
- 3. Is the direction of the effect (+/-) of variables the same across models?



Findings

- Substantial variation in the selected variables
 - Different databases -> different 'risk factors'
- Sign of 'risk factors' can differ across models
- For 'risk factor' detection:
 - Investigate model robustness
 - Use other techniques (e.g. univariate analysis)



10.0-

10 variables

Overlap in top

Same O

⇔ Non Hemorrhagic Stroke
 ⇔ Acute Myocardial Infarction
 ⇔ Pulmonary Embolism
 ⇔ Hemorrhagic Stroke

Guillain Barre Syndrome

Appendicitis
Anaphylaxis
Dis Intra Coag
Encephalomyelitis

Take home message:

Be careful interpreting prediction models as the identified 'risk factors' appear to depend on study design choices.

'Risk factor' = 'variable associated to the outcome'





TRAJECTORYVIZ: INTERACTIVE VISUALIZATION OF TREATMENT TRAJECTORIES

Maarja Pajusalu, Institute of Computer Science, University of Tartu



TrajectoryViz: Interactive visualization of treatment trajectories

♣ PRESENTER: Maarja Pajusalu maarja.pajusalu@ut.ee

Co-authors: Marek Oja, Sirli Tamm, Markus Haug, Raivo Kolde Institute of Computer Science University of Tartu, Estonia





Create discrete sequences with R package

Cohort2Trajectory



Create interactive visualisations with R package TrajectoryViz

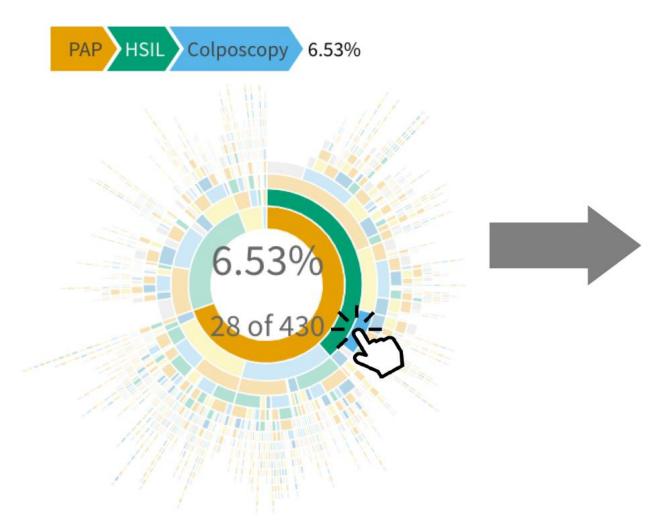


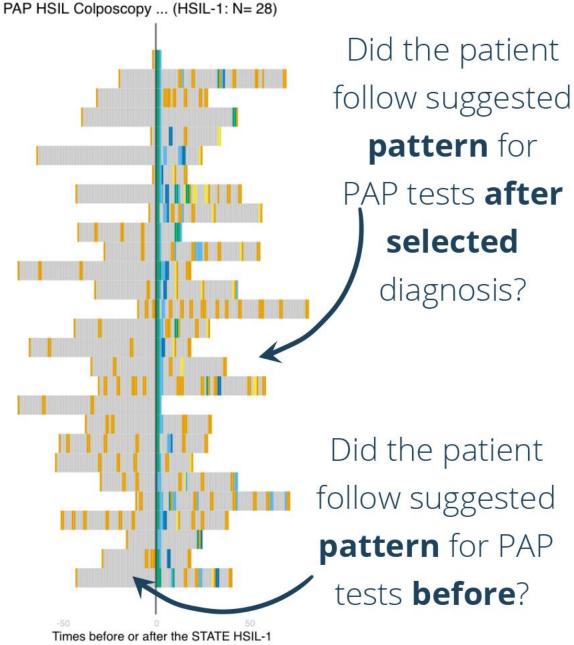


Use Case 1:

Cervical Cancer patients

Sequences on a Sunburst Chart

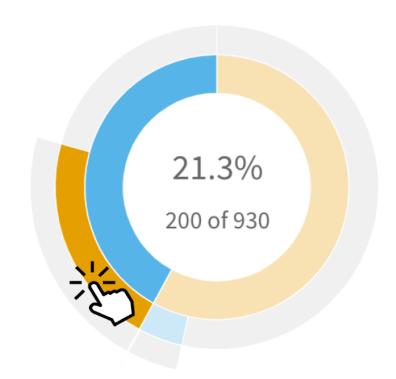


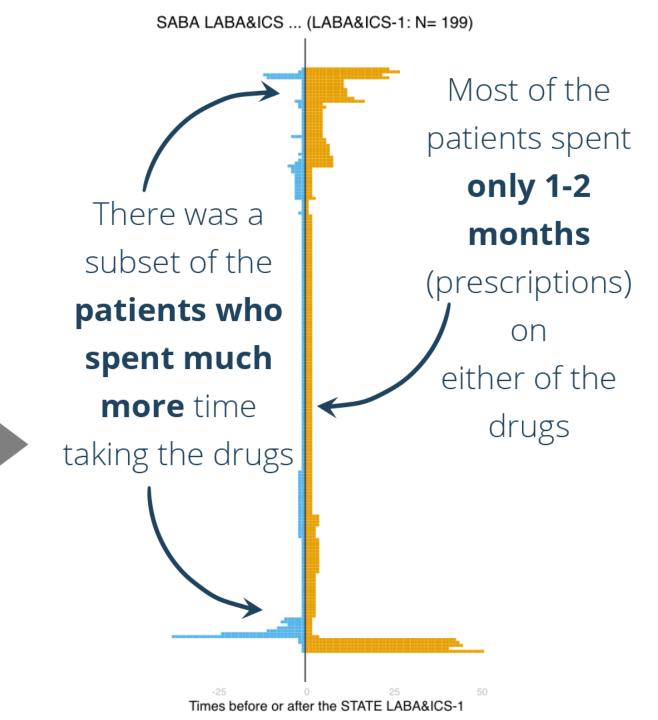


Use Case 2: Asthma patients

Sequences on a Sunburst Chart

SABA LABA&ICS 21.3%





Thank You!









ASSESSING TREATMENT EFFECT HETEROGENEITY USING THE RISKSTRATIFIEDESTIMATION R-PACKAGE

Alexandros Rekkas, Erasmus MC

Assessing treatment effect heterogeneity using the RiskStratifiedEstimation R-package

Alexandros Rekkas



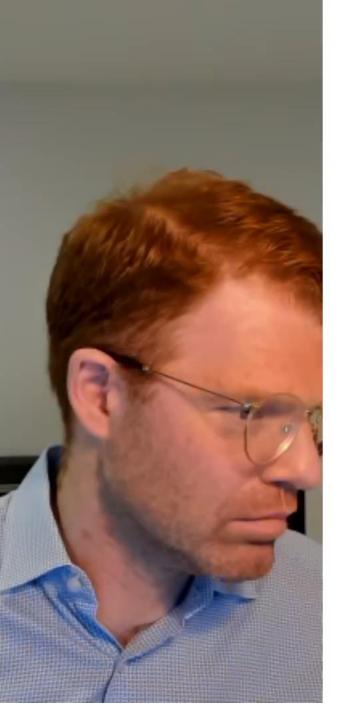






DEFINING THE VALID ANALYTIC SPACE FOR QUANTITATIVE BIAS ANALYSIS IN PHARMACOEPIDEMIOLOGY

James Weaver, Janssen R&D





Defining the analytic space for valid QBA

OHDSI Europe Symposium 24 June 2022

James Weaver [jweave17@its.jnj.com] Global Epidemiology, Janssen R&D NDORMS, University of Oxford







Background

- Bias from outcome misclassification often ignored in casual estimation
- QBA: a proposed solution

Objective

 Evaluate QBA performance across large set of plausible scenarios

Methods

 Applied QBA across incidence x effect size x measurement error analytic space

Results

- Small specificity change has large impact on effect estimate
- Limited impact of sensitivity at low incidence

Discussion

QBA produces implausible or invalid estimates in many common comparative effect estimation scenarios



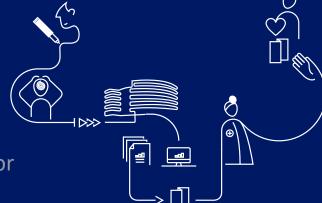
A PILOT STUDY TO EVALUATE THE FEASIBILITY
OF USING OBSERVATIONAL HEALTH
DATA SCIENCES AND INFORMATICS
ANALYTICS TOOLS FOR SUPPORTING THE
VALIDATION OF SAFETY SIGNALS

Ceyda Pekmez Kristiansen, Novo Nordisk



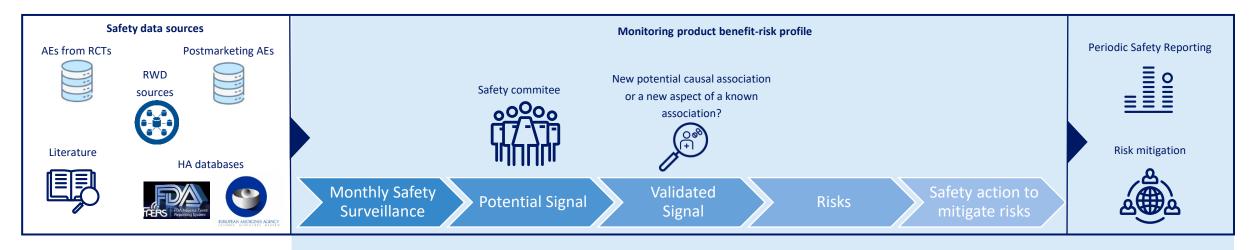
A pilot study to evaluate the feasibility of using OHDSI analytical tools for supporting safety surveillance

Ceyda Tugba Pekmez Kristiansen¹; Lasse Christensen¹; Michael Stellfeld¹; Atheline-Major Pedersen¹; Ditte Mølgaard-Nielsen¹; Mark White¹; Peter Jelnes¹



cypk@novonordisk.com

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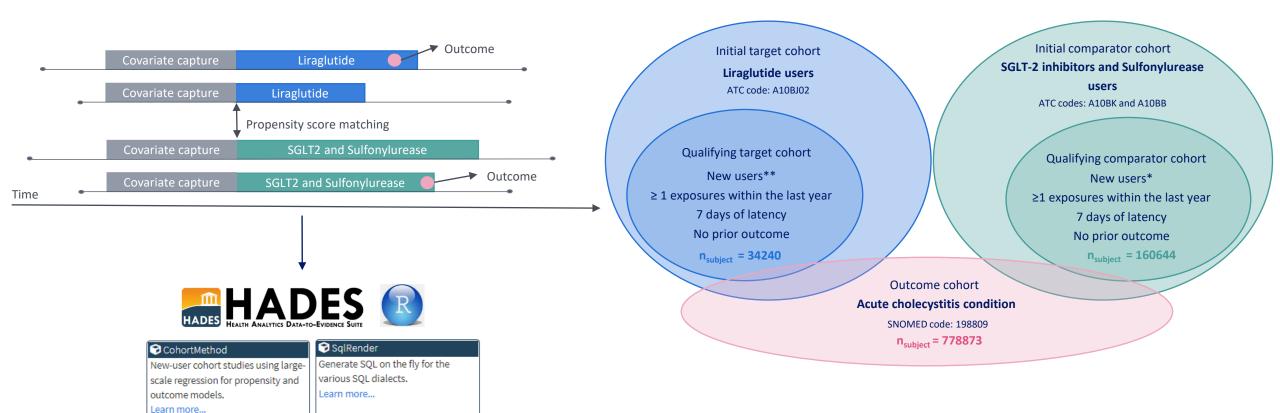


Real world data (RWD) for supporting safety surveillance

- RWD supports the safety signal validation process, especially when the evidence from traditional safety data sources is scarce
- Gallbladder- or biliary tract-related events (including acute cholecystitis) are known risks for Victoza® (liraglutide) and Saxenda® (liraglutide)
- A known risk for liraglutide was chosen for the pilot study to evaluate the feasibility of implementing population level effect estimation into the safety surveillance process

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Cohort definition – Truven Marketscan CCAE* 2020



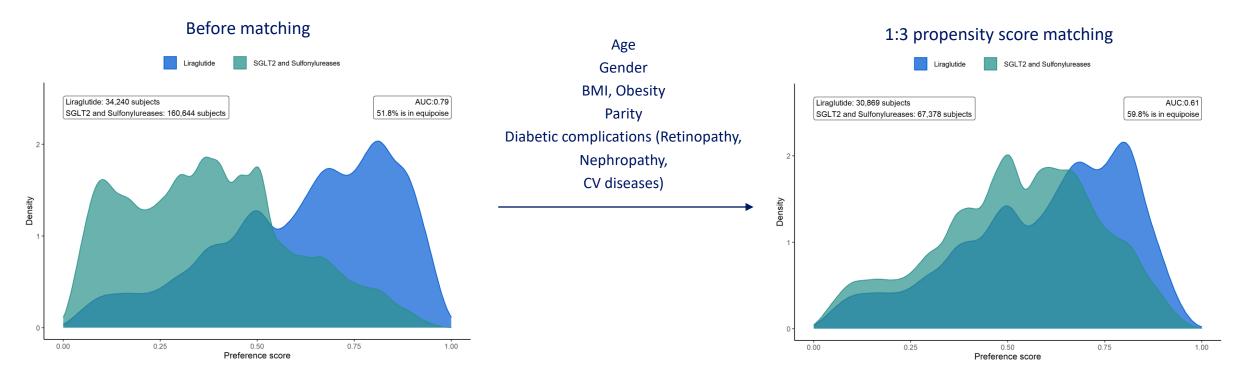
IBM® Watson Health™. Commercial Claims and Encounters Database and Medicare Supplemental and Coordination of Benefits Database IBM Marketscan Research Databases User Guide. Certain data used in this study were supplied by International Business Machines

^{*}Commercial Claims and Encounters

^{**}First time use of drug within last 1 years of observation period

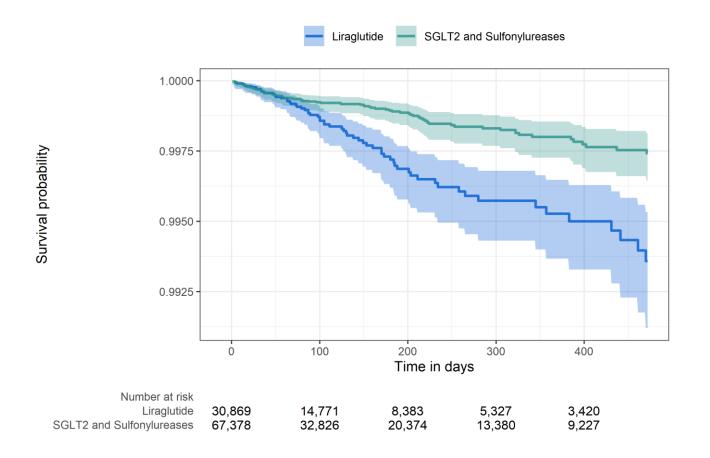
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Propensity score matching



	Cases (n)	Prevalence (per 1000)
Target	71	2.30
Comparator	83	1.23
Total	154	1.57

Population level risk estimation



Cox-proportional Hazard ratio: 2.26 CI: [1.70 – 3.03]

Minimum detectable relative risk: 1.62 ± 0.17 (SE)

Conclusion – Next steps

- OHDSI analytics tools have promising potential for utilising RWD sources to support the validation of safety signals
- The result supports a known risk of acute cholecystitis for liraglutide on a RWD source
- A new test case for another therapeutic area
 - Negative outcome controls
 - Data driven selection of covariates





Rapid Fire Presentations of Collaborators

THANK YOU!





- Findable, standardised data at scale through the EHDEN Database Catalogue -

24 June 2022

Julia Kurps, The Hyve









EUROPE: AN OCEAN OF DATA & A DESERT FOR ANALYSIS







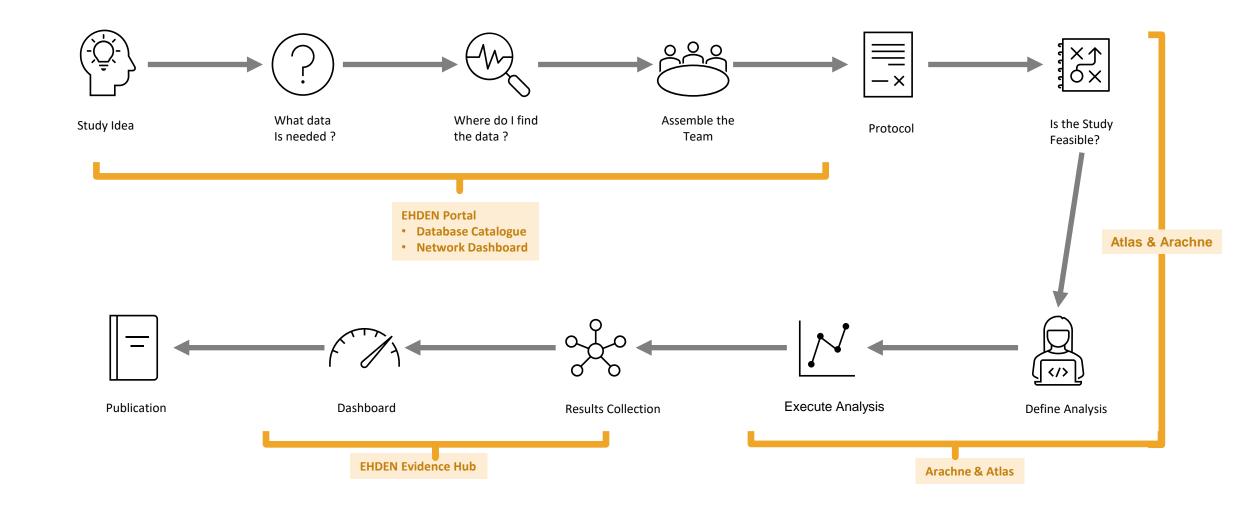


OPEN UP — KEY RESEARCH TOOL IN EHDEN





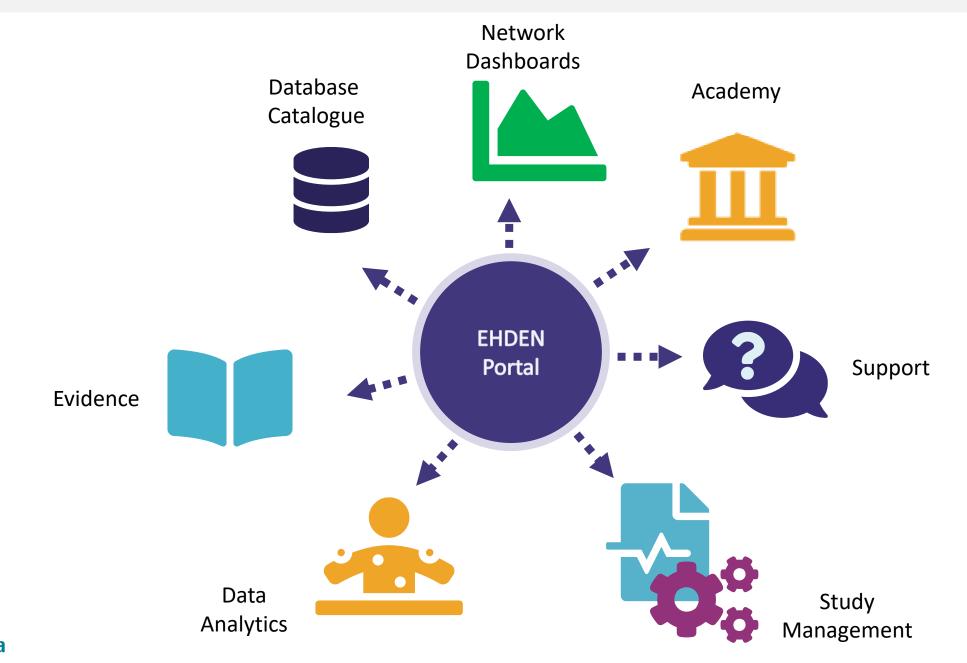
ROLE OF THE EHDEN PORTAL IN THE STUDY WORKFLOW







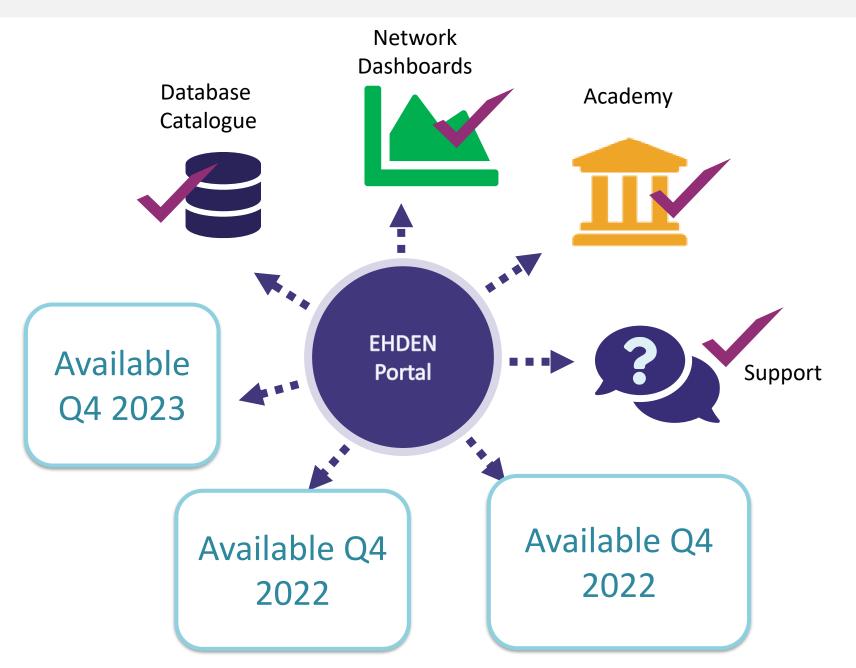
THE VISION OF THE EHDEN PORTAL — ONE-STOP-SHOP







EHDEN PORTAL RELEASE TODAY







DATABASE CATALOGUE







GEOGRAPHICAL SPREAD DATA PARTNERS

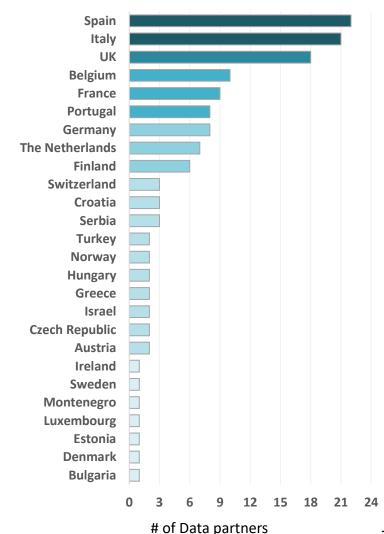


Geographic spread of data partners. The shade of blue indicates the # of data partners in that country (darker = more)

Applications (n=283)

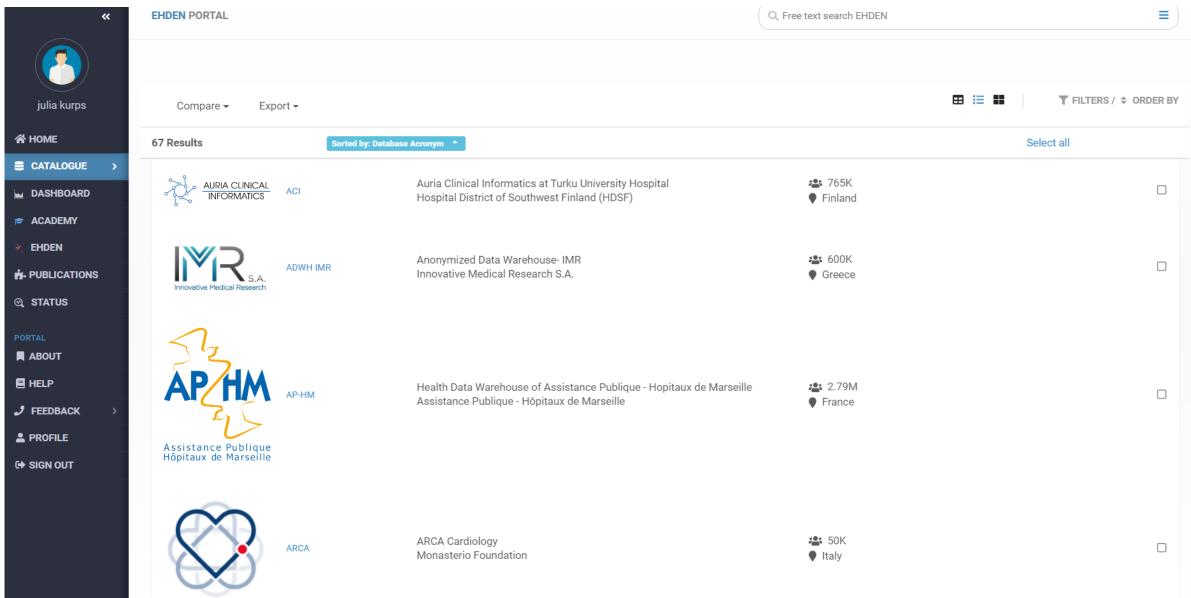


Awarded applications (n=140)





Database Catalogue – 67 Data Partners...and Counting











COMPONENTS OF THE DATABASE CATALOGUE

Database Fingerprint

- meta data -

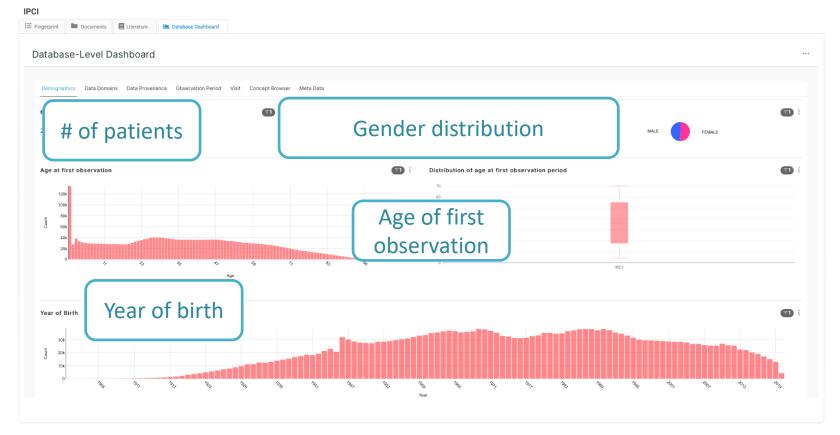


Database Dashboards

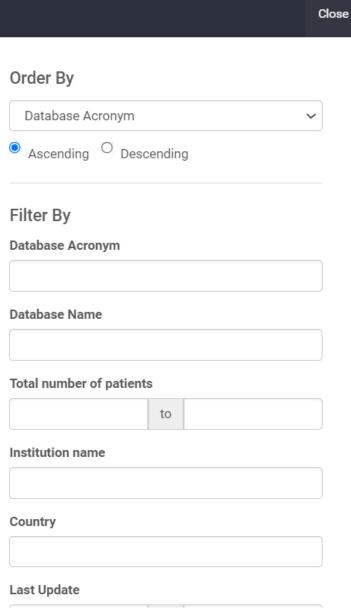
- based on CDM data -



1. Database Description (20/20)	100%
2. Contact Details (10/10)	100%
3. Technical Details CDM (5/5)	100%
4. Data Governance and Ethics (9	100%
5. Publications (2/2)	100%







All databases in France?

Which databases have occurrences of myocardial infarction?

Which databases have **GP data** in combination with **hospital data**?

In which countries do we have up to date data on COVID-19 vaccinations?

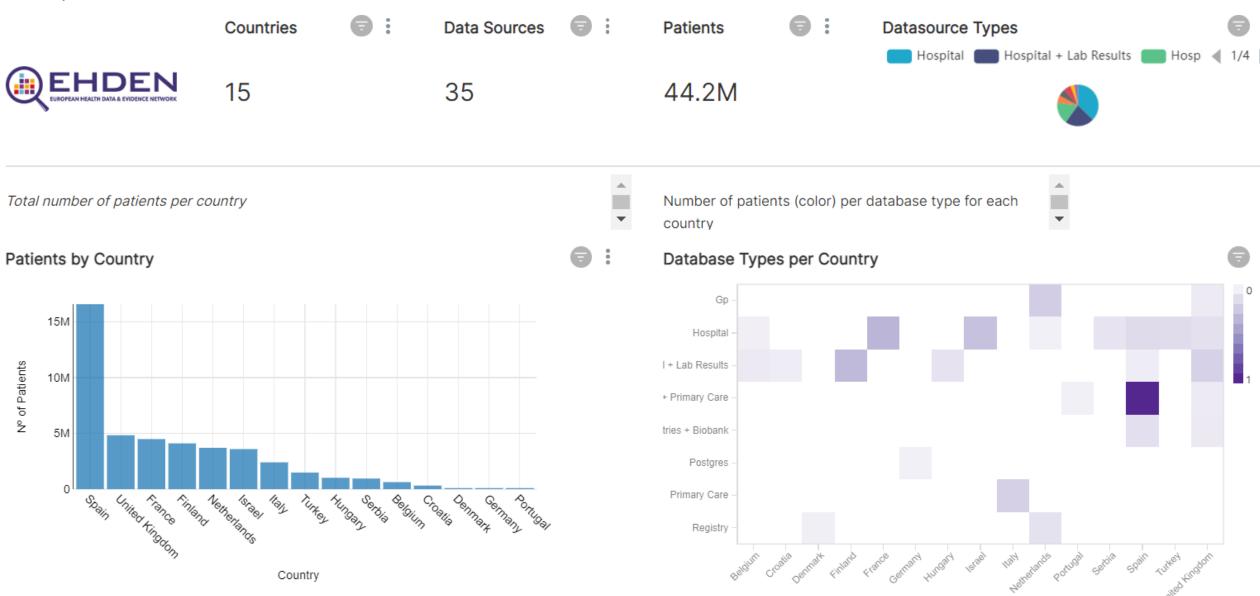








NETWORK DASHBOARDS - OVERVIEW







FILTER TO FIND RELEVANT DATA



Use case examples

- Geographical spread
 Identify all EHDEN Data Partners from France
- Range of Source data systems
 Identify Data Partners that can provide GP data in combination with hospital data





Network Dashboards - Browse - Explore - Discover

Overview Demographics **Data Domains** Data Provenance **Observation Period** Concept Browser Visit About Distribution of age at first observation period 100 80 60 40





GETTING STARTED

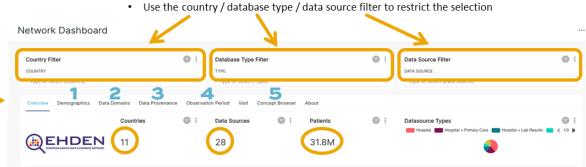


EHDEN Network Dashboard at a Glance

Getting Access to the Network Dashboard

- Enter credentials on the EHDEN portal (https://portal.ehden.eu)
- Select the Network Dashboard icon on the left panel





Total Number of datasources and patients in the network

Understanding gender or year of birth or age coverage

- Go to the tab 'Demographics'
- Gender distribution is shown in the top pie chart
- Age at first observation is presented in tabular format, bar chart and boxplot
- Year of birth is presented as bar chart.

Understanding what data are available for one or more data sources

- Go to the tab 'Data Domains'
- The 'Average number of records per person' indicates how many records per entity type are available. Use in conjunction with the above filters to select on country/database type or data source
- The pie chart gives the total number of records per entity type

What is the source of the data?

- Go to the tab 'Data Provenance'
- of the respective concept types e.g. EHR problem list entry, primary or secondary condition, hospitalization cost record etc.

What's the available observation period?

- Go to the tab 'Observation Period'
- Check the second graph cumulative observation time
- Click in the legend on a single item to select / deselect
- Double click to select a single data source

Which data sources do contain a particular concept?

Go to the tab 'Concept Browser'

- In the concept type, start typing part of the text string
- Select the concept that you want to search for
- Hit APPLY

COUNTRY France ×

• In the pivot table you can see approx. total number of records and descendant number of records. Further down, you see a graphical representation

Nice-to-know

- 1. Knowing the last update per datasource
- · On the landing page of the network dashboard
- Scroll to the bottom meta data view
- Cutoff date from the source data: source release date
- · CDM date: cdm release date



2. Filtering results and other tricks

- Option A: top filters for selection on country / database type or data source
- Option B: When a legend is displayed, individual items can be selected / unselected or double click select a single item from another list
- 3. Maxime your graph
- · When a graph is not readable: click on the icon and select 'maximise graph' from the pop-up



• The pivot table 'Visit Type Pivot' gives a count per entity type





AND THAT'S JUST THE BEGINNING...



More Data Partners will be added

Submission deadline DP Call #6 just closed

Integration with Data Analytics and Study Management capabilities







Implementation of Production Instance of service desk for study requests

WELCOME – WE ARE OPEN



Register your account today at

Portal.EHDEN.eu

After approval
you can start exploring
high quality
real world data from
> 44 M patients
from 15 European countries



Welcome to the EHDEN Portal

The European Health Data & Evidence Network (EHDEN) project aspires to be the trusted observational research ecosystem to enable better health decisions, outcomes and care. Its mission is to provide a new paradigm for the discovery and analysis of health data in Europe, by building a large-scale, federated network of data sources standardised to the OMOP common data model.

The EHDEN Portal provides an entry point to the growing list of tools in our ecosystem. Of these tools, Database Catalogue (including Dashboards) and EHDEN Academy are available for general use, while Arachne, Atlas and the Service Desk are still in development and only available to EHDEN Consortium members. Click on the icons below for more information. Your feedback is highly appreciated.



Database Catalogue

Provides metadata on the databases in the EHDEN data network



Network Dashboards

Allows to analyse and compare aggregated data from the OMOP CDM databases in the



EHDEN Academy

Our free, online and publicly available learning platform



Publications

An overview of all deliverables and publications of EHDEN

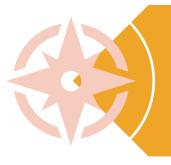






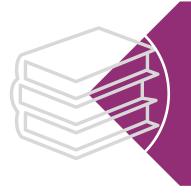


WP4 – BUILDING THE INFRASTRUCTURE



Mapping

- White Rabbit / Rabbit in a Hat
 - Usagi
 - Data Quality Dashboard



Study Management and Execution

- EHDEN Portal
- Database Catalogue
 - Arachne
 - Atlas
- EHDEN Evidence Hub

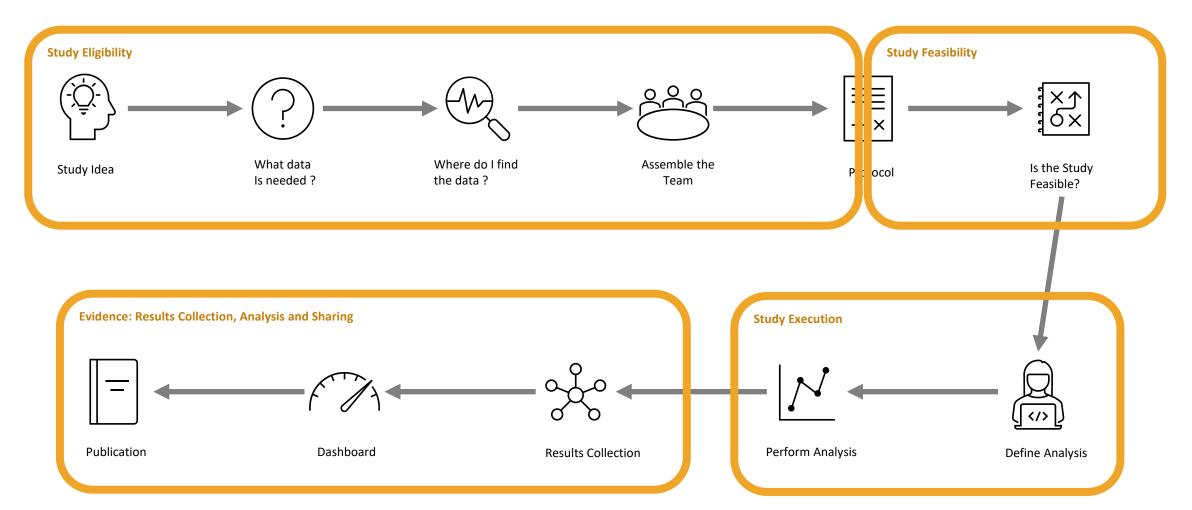


Auxilliary

- EHDEN Academy
- Virtual Training Environment
 - Website
 - Forum



STUDY WORKFLOW IN EHDEN — TECHNICAL PERSPECTIVE







Lunch, Collaborator Showcase, and Early Investigator meetings



The lunch is made possible with the help of Synapse



The collaborator showcase is made possible with the help of Promptly



Early Investigators mentor meetings in the Queen's Lounge



