

Cloud based access and analytics to advance precision medicine

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Thank you to our <u>participants</u> and community partners!



All of Us Community and Provider Partner Network (as of July 2020)





















































































Thank you to our consortium partners!

The Participant Center







· fitbit













WONDROS







COLLABORATIVE



PicnicHealth



Blood Bank



COLUMBIA UNIVERSITY

MEDICAL CENTER

NewYork-Presbyterian







HPO Network

(Health Care Provider Organizations)

RMCs

All of Us California

UC San Diego Health

UCI Health







Illinois Precision Medicine Consortium

Northwestern Medicine

THE UNIVERSITY OF CHICAGO

≠NorthShore

HEALTH

UI Health | @

ORUSH

THE UNIVERSITY OF ILLINOIS COLLEGE OF MEDICINE

All of Us New England











Trans America

Consortium







New York City

Consortium

HEALTH+

HOSPITALS





LL4B SCHOOL OF MEDICINE The University of Alabama at Birminghan



(SA HEALT

FQHCs (Federally Qualified Health Centers)







LIZE SCHOOL OF

LSU Health

Montgomery Regional Medical Campus

LAB MEDICINE

THE UNIVERSITY OF ALABAMA®

All of Us SouthEast **Enrollment Center**

> UNIVERSITY OF MIAMI MILLER SCHOOL











All of Us Wisconsin

Marshfield Clinic Health System

Participant Technology Systems

Center (PTSC)















Banner Health

University of Arizona and





























Commun*ty









vibrent 👭



Biobank

Center (DRC)

COLUMBIA UNIVERSITY

MEDICAL CENTER



















Thank you Data and Research Center (DRC) and NIH teams!











VANDERBILT UNIVERSITY







All of Us Research Program Objectives

Nurture relationships

with **one million or more** participant partners, from all walks of life, for decades



of researchers and funders hungry to use and support it



Our mission

To accelerate health research and medical breakthroughs, enabling individualized prevention, treatment, and care for all of us



Deliver the largest, richest biomedical dataset

that is easy, safe, and free to access



AllofUs Research Hub

Welcome, All of Us Researchers.

The All of Us Research Program, part of the National Institutes of Health, is building one of the largest biomedical data resources of its kind. The All of Us Research Hub will store health data from a diverse cohort of one million or more participants in the All of Us Research Program.

Here in the Research Hub, everyone can learn more about the types of data participants are providing and how approved researchers can use our data and tools to conduct studies that may speed up medical breakthroughs.

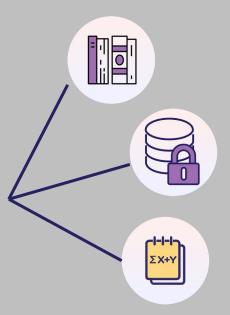
APPLY FOR ACCESS







Launched Beta in May 2020!



Outline for Today

1. Data Overview



2. Live demo of <u>researchallofus.org</u> & Researcher Workbench



Outline for Today

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Our Data are Growing. Here are the Current Data Types





Participant Surveys



Physical Measurements



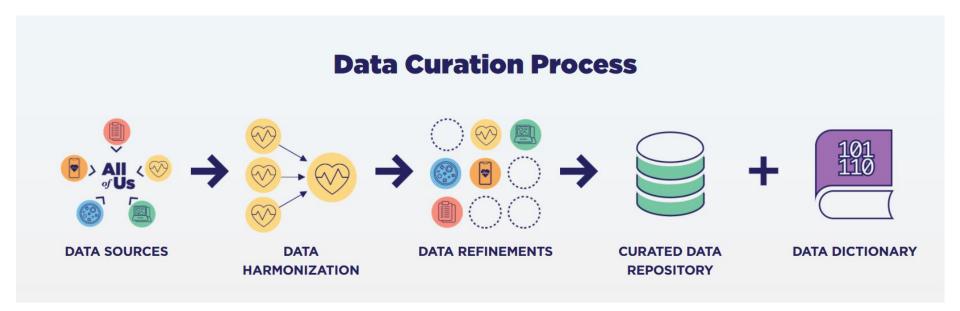
Biosamples



Available in the Current Research Dataset



Data are harmonized and standardized to a common data model



We use the OHDSI collaborative's OMOP Common Data Model (more info here)

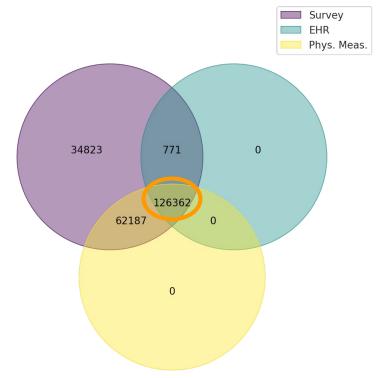


Research Data Available Now

Data Type	Participant Count
Survey	>224,000
Physical Measurement	>188,000
Electronic Health Record	>127,000

^{*}Counts reflect unique participants with ANY of data of the specified type.

Count of participants with multiple data types



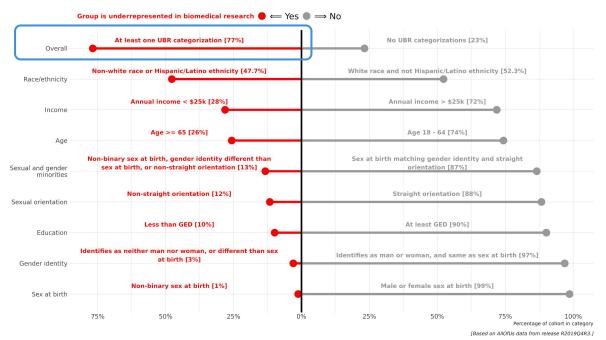


Participants included in this dataset are diverse.



Cohort diversity: underrepresented groups in biomedical research

Categorizations based on explicit responses to survey questions.



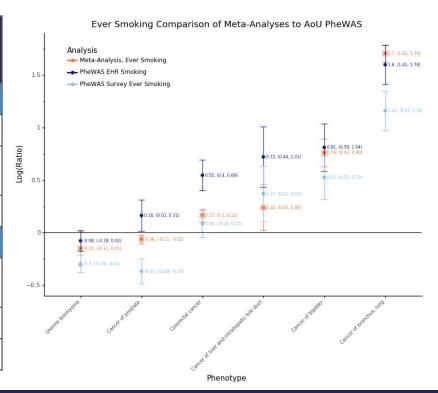


A. Ramirez, L. Suleiman, D. Schlueter, et al., *The All of Us Research Program: data quality, utility, and diversity*, medRxiv 2020.05.29.20116905; doi: https://doi.org/10.1101/2020.05.29.20116905

This dataset reproduces known associations.



Description	EHR Ever Smoking OR (95% CI)	Survey Ever Smoking OR (95% CI)
Top 3 Increased risk effects		
Cancer of the bronchus; lung	4.94 (4.11, 5.95)	3.19 (2.65, 3.84)
Cancer within the respiratory system	4.94 (4.12, 5.92)	3.15 (2.62, 3.78)
Malignant neoplasm of bladder	2.36 (1.87, 2.98)	1.76 (1.42, 2.18)
Top 3 Decreased risk effects		
Vascular hamartomas and non-neoplastic nevi	0.51 (0.42, 0.62)	0.55 (0.48, 0.64)
Nevus, non-neoplastic	0.52 (0.43, 0.64)	0.57 (0.49, 0.66)
Benign neoplasm of skin	0.53 (0.49, 0.58)	0.62 (0.58, 0.66)

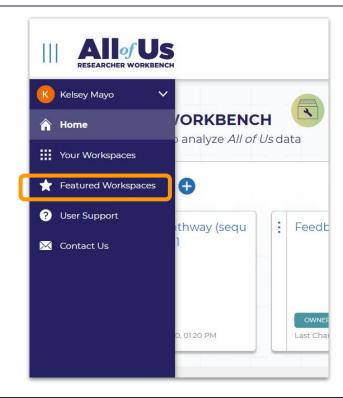


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Want to learn more?







A. Ramirez, L. Suleiman, D. Schlueter, et al., *The All of Us Research Program: data quality, utility, and diversity*, medRxiv 2020.05.29.20116905; doi: https://doi.org/10.1101/2020.05.29.20116905

Coming Soon (Late 2020)...



Electronic Health Records (EHR)



Participant Surveys



Physical Measurements



Experience
(COPE Survey)



Refreshed + New Data Types



Research Data Available Soon (Winter 2020)

Data Type	Participant Count*	
Survey	>315,000	~40% increase
COPE Survey	>63,000	New!
Physical Measurement	>263,000	~40% increase
Electronic Health Record	>203,000	~60% increase!
FitBit	>8,000	New!

^{*}Counts reflect unique participants with ANY of data of the specified Counts are approximate and subject to change upon final release of the dataset.



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2. Live demo of <u>researchallofus.org</u> & Researcher Workbench



ABOUT

DATA

TOOLS D

DISCOVER

FAQ

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APPLY

Research Hub

Data Browser

Apply Page

Researcher Workbench

Support Hub

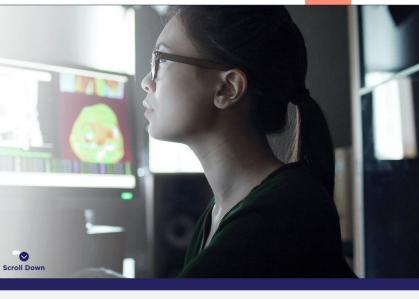
Ex. Workspace

Welcome, *All of Us* Researchers.

The *All of Us* Research Program, part of the National Institutes of Health, is building one of the largest biomedical data resources of its kind. The *All of Us* Research Hub will store health data from a diverse cohort of one million or more participants in the *All of Us* Research Program.

Here in the Research Hub, everyone can learn more about the types of data participants are providing and how approved researchers can use our data and tools to conduct studies that may speed up medical breakthroughs.

APPLY FOR ACCESS



Data Snapshots

Our Data Snapshots are the fastest way to view the diversity of the All of Us Research Program participant data.

These snapshots provide visualizations of participant demographics, geographic distribution, and more.







Data includes 225,140 participants and is current as of 2/11/2020. **Data Browser** Researcher Workbench Support Hub Ex. Workspace View Complete Survey 57

Research Hub

Apply Page

Search Across Data Types • Q Keyword Search





Videos



FAQs

EHR Domains: 0

Conditions () 20,776

medical concepts

113,200 participants in this domain

View Top Conditions

Drug Exposures 6

20,951

104,500 participants in this domain

View Top Drug Exposures

Labs & Measurements 6

10.049

medical concepts

109,100 participants in this domain

View Top Labs & Measurements

Procedures 6

20,546

medical concepts

102,140 participants in this domain

View Top Procedures

Survey Questions:

The Basics

16

questions available

225,140 participants in this domain

This survey includes participant demographic information.

Overall Health 6

questions available

219.800 participants in this domain

Survey includes information about how participants report levels of individual health.

View Complete Survey

Family Medical History (1)

Lifestyle 6

26

questions available

218,500 participants in this domain

Survey includes information on participant smoking, alcohol, and recreational drug use.

View Complete Survey

Personal Medical History

465

questions available

39,320 participants in this domain

This survey includes information about past medical history, including medical conditions and approximate age of diagnosis.

View Complete Survey

Health Care Access & Utilization (1)

questions available 45.820 participants in this domain

Survey includes information about a participant's access to and use of health

questions available 41,600 participants in this domain

Survey includes information about the medical history of a participant's

Research Hub

Data Browser

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Researcher Workbench

Support Hub

Ex. Workspace

Apply to be an All of Us Researcher

The Researcher Workbench is open to researchers whose institutions have signed a Data Use and Registration Agreement with the All of Us Research Program. Researchers at this time must also have an eRA Commons account and complete the All of Us Research Program data access process before they can access the Researcher Workbench and Registered Tier data. For more information, please visit the Data Use Policies page. If you are a researcher who does not have an eRA Commons account, please work with your institution to create one. Once your account is established, you may apply for access to the Researcher Workbench. For more information, visit the eRA Commons website.

About the Beta Researcher Workbench

Currently, an institutional agreement & eRA Commons account must be in place.

Feedback is welcome.

The tools will continue to evolve.

The program cohort is actively growing, and so is our data.

The goal is to be a true game changer for understanding health.

How the Application Process Works









Research Hub

Data Browser

Apply Page

Researcher Workbench

Support Hub

Ex. Workspace

Welcome to the

RESEARCHER WORKBENCH

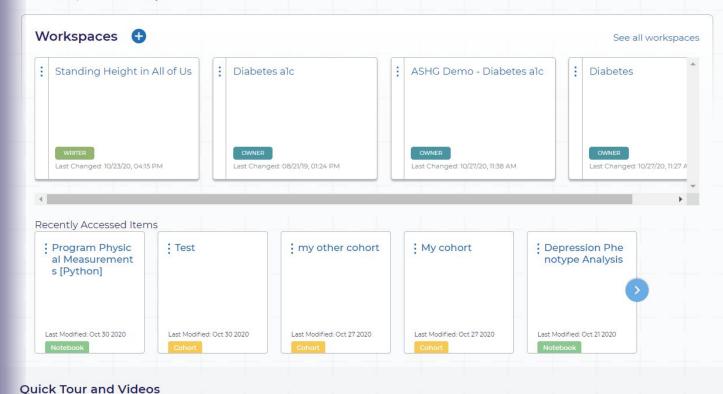






The secure platform to analyze All of Us data

orkspac



User Support Hub

Search our knowledge base for answers to common questions

Q Enter a question, topic, or a keyword...

Research Hub

Data Browser

Apply Page

Researcher Workbench

User Support Hub

Ex. Workspace

Explore All of Us Researcher Workbench Resources



New to the Researcher Workbench? Looking for some quick help for working with our tools or data? Start here.



Find helpful resources to better understand the All of Us dataset and how to analyze it.



Post your questions and comments for other users here.



Frequently Asked Questions

Look for answers on workspaces, concepts, and everything in between.



Announcements

Get the latest All of Us news and announcements.





Workspaces >

Demonstrating All of Us Anthropomorphic Data Qualit

Cohorts 0 Datasets 0 A cohort is a group of participants based on specific A dataset is a table containing data about a cohort that can be exported for analysis. criteria. Participant ID 1 Participant ID 1 ID 2 Participant ID 2 \rightarrow Participant ID 2 Labs Participant ID 3 Participant ID 3 **Your Cohort Data About** Your Dataset All of Us **Your Cohort Your Cohort Participants** Show: Show All Cohorts Cohort Reviews Concept Sets Datasets

Research Hub

Data Browser

Apply Page

Researcher Workbench

Support Hub

Ex. Workspace

Physical Measur ements Concep t Set 2

Physical Measur ements Concep t Set 2 : Height : PPI he

: PPI height

Live Demo on **Synthetic Data***

"Demonstrating All of Us Anthropomorphic Data Quality"

Primary Investigator

Kelsey Mayo, Ph.D.



Co-Investigator

Adrienne Roman, Ph.D.



Data Scientist

Francis Ratsimbazafy, Ph.D.



Graduate Student

Michael Holmes, B.S.





Outline for Today

1. Data Overview



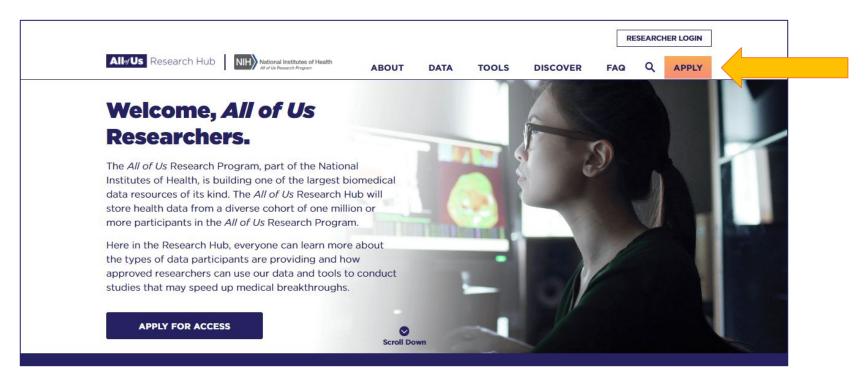
2. Live demo of researchallofus.org & Researcher Workbench

Thank you for listening!



Interested in becoming a Beta Researcher?

Visit researchallofus.org to apply!





Acknowledgements

The All of Us Research Program is supported by the National Institutes of Health, Office of the Director: Regional Medical Centers: 1 OT2 OD026549; 1 OT2 OD026554; 1 OT2 OD026557; 1 OT2 OD026556; 1 OT2 OD026550; 1 OT2 OD 026552: 1 OT2 OD026553: 1 OT2 OD026548: 1 OT2 OD026551: 1 OT2 OD026555; IAA #: AOD 16037; Federally Qualified Health Centers: HHSN 263201600085U; Data and Research Center: 5 U2C OD023196; Biobank: 1 U24 OD023121; The Participant Center: U24 OD023176; Participant Technology Systems Center: 1 U24 OD023163; Communications and Engagement: 3 OT2 OD023205; 3 OT2 OD023206; and Community Partners: 1 OT2 OD025277; 3 OT2 OD025315; 1 OT2 OD025337; 1 OT2 OD025276.

In addition, the *All of Us* Research Program would not be possible without the partnership of its participants.

