

Kaiser Permanente Research Bank

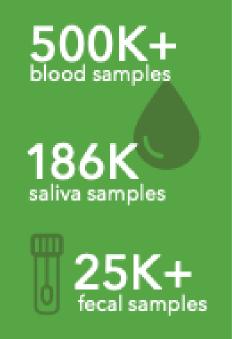


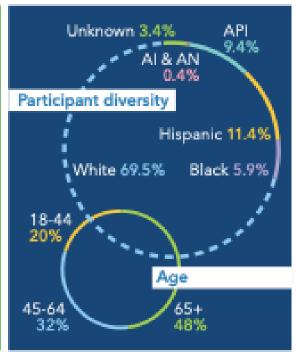
CELEBRATING 10 YEARS 185+ Research Studies



400K+ health surveys Covid-19 surveys

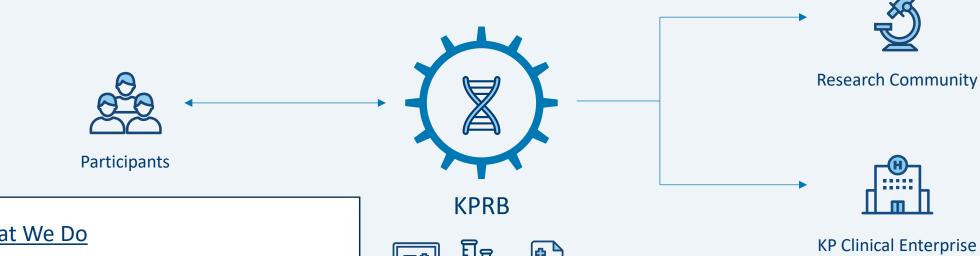
Metabolomic Genomic 400K Data 55K participants participants







KPRB Overview



What We Do

- Support research projects (internal and external)
- **Support new cohort development**
- **Develop new data resources**
- **Leverage our data** to support research and clinical programs

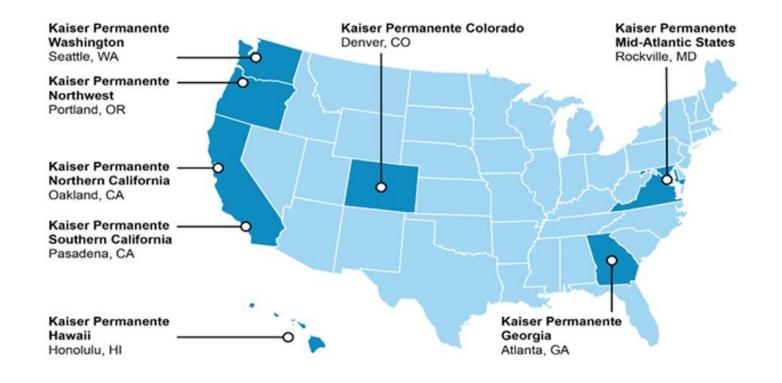






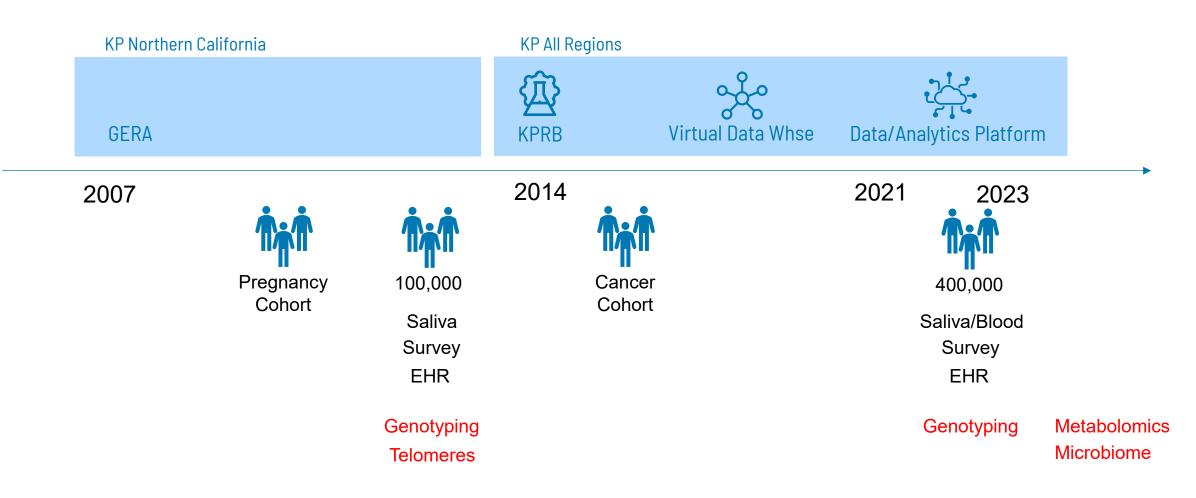
Medical Records Biospecimens Surveys **Genomic Data** Other Omics

Kaiser Permanente: Organized Around Eight Regions





KPRB History





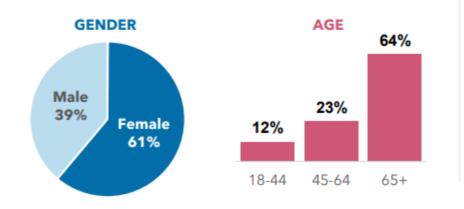
KPRB Demographics

POPULATION & DEMOGRAPHICS*:

Among the total KPRB population there were special recruitment efforts focused on enrolling pregnant women and members with a cancer diagnosis.

Total Population General + Cancer + Pregnancy Cohorts	430,967
General Cohort Any KP member age 18 or older	316,252
Cancer Cohort KP members with a known history of cancer	89,739
Pregnancy Cohort KP members over age 18 in KP Northern California enrolled during pregnancy	24,976

^{*}Data is among members who have provided a sample to KPRB (430,967); enrolling through 11/2023



KAISER PERMANENTE REGIONS



RACE/ETHNICITY

American Indian & Alaska Native	1,053
Asian American	35,663
Hispanic	46,687
Native Hawaiian/Other Pacific Islander	1,543
Non-Hispanic Black	23,604
Non-Hispanic White	309,670
Other	6,754
Unknown	5,816



KPRB EHR Data

Medical Records History and Follow-Up

(All participants, 1996-2022)

Median 75th Pctl 25th Pctl Mean Max 19.35 Total Years of KP Membership 13.24 20.91 27.33 28.36 Total Years of Follow-Up 3.65 7.36 11.73 15.85 6.84

Medical Records History and Follow-Up

(Pregnancy Cohort)

	25th Pctl	Median	Mean	75th Pctl	Max
Total Years of KP Membership	8.06	14.47	15.02	22.13	36.07
Total Years of Follow-Up	5.58	8.19	7.56	9.98	14.61

Medical Records History and Follow-Up

(Cancer Cohort)

	25th Pctl	Median	Mean	75th Pctl	Max
Total Years of KP Membership	13.61	19.94	20.70	28.87	42.10
Total Years of Follow-Up	1.98	5.60	6.34	11.00	14.77



KPRB Resources

405,232

KPRB Biomolecular Data

90.23% Percentage of KPRB participants with at least one biomolecular data type

415K+ Total number of assays

View Biomolecular Data Types

449,056

At least one address matched to a U.S. Census Geocode

99.99% Percentage of KPRB Cohort with at least one data point

2M+ Total number of data points

448,925

Diagnoses

99.96% Percentage of KPRB Cohort with at least one data point

333M+ Total number of data points

View Key Diagnoses

433,444

KPRB Biosamples

96.51% Percentage of KPRB participants with at least one biospecimen

737K+ Total number of biosamples

View Biosample Types

448,957

Clinical Encounters

99.96% Percentage of KPRB Cohort with at least one data point

171M+ Total number of data points

447.747

Lab Results

99.69% Percentage of KPRB Cohort with at least one data point

266M+ Total number of data points

View Key Laboratory Results

382,305

KPRB Surveys

85% Percentage of KPRB participants with at least one intake survey

432K+ Total number of intake surveys

View Survey Types

389,688

Clinically-Administered Medications

86.77% Percentage of KPRB Cohort with at least one data point

60M+ Total number of data points

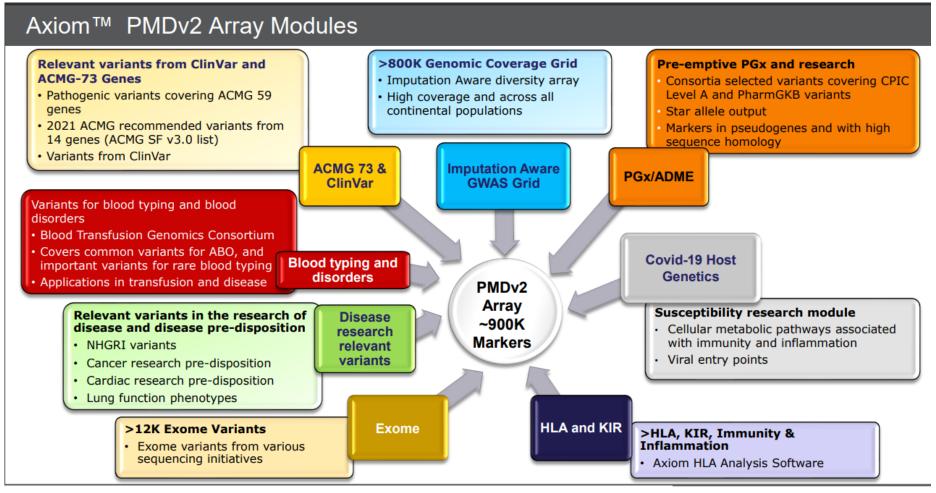
447,545

Outpatient Medications

99.65% Percentage of KPRB Cohort with at least one data point

132M+ Total number of data points

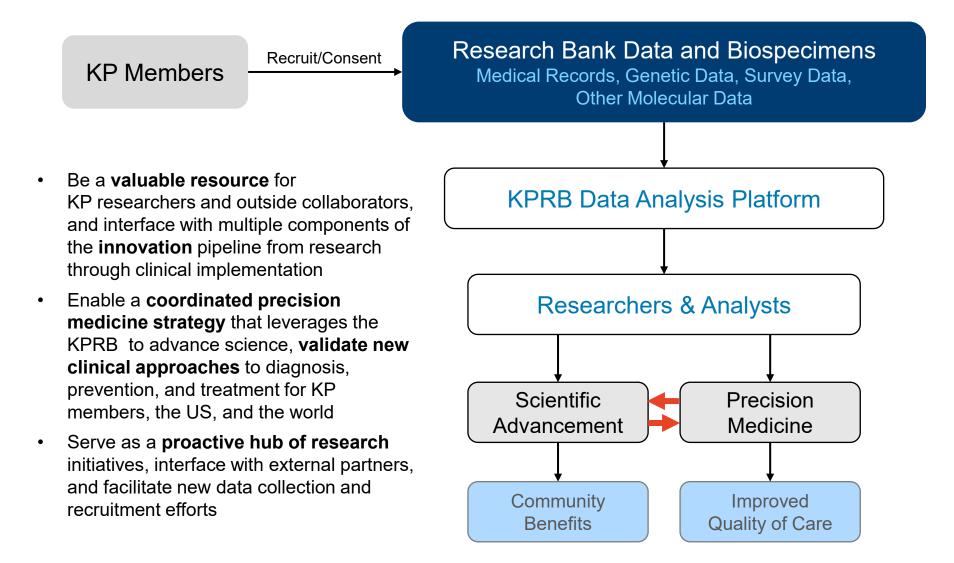
KPRB Genomic Data





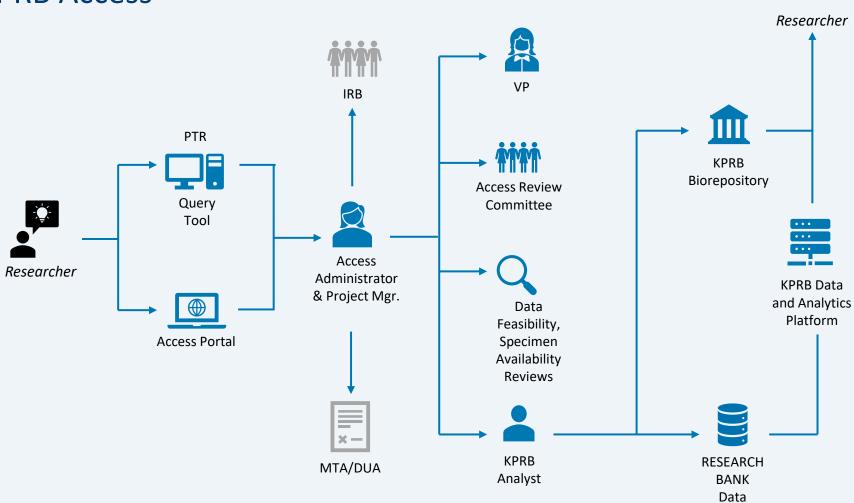


KP Research Bank Strategy and Mission





KPRB Access



KPRB Recent and Current Projects

- Genotyping of 400,000 participants on modern array with clinically relevant content
- Imputation complete (TopMED)
- Genetic ancestry, PCs, local ancestry estimates
- PGS Calc & PGS Catalog
- Partnership with Genomics, Inc (UK) to develop polygenic risk models for 8-10 diseases
- NMR-based metabolomics on 55K participants with longest follow-up time
- FIT-kit based fecal samples from colorectal cancer screening from microbiome work (40k by 2025)
- Longitudinal blood sample collection from 5+ year participants (60k by 2025)
- Return of results survey and pilot projects



KPRB Partnerships

Access Initiative

- Exploring partnerships with key institutions that would give their researchers easier access
 - HCSRN partner institutions use a common data model (VDW)
 - Opportunity for easier data integration and validation
 - Partnership would include common data use and research agreements to speed application process
 - Partner institutions could reduce the need for KP co-Investigator
 - Other process improvements
 - Access to Trusted Research Environment
 - Shared best practices, disease/outcomes definitions



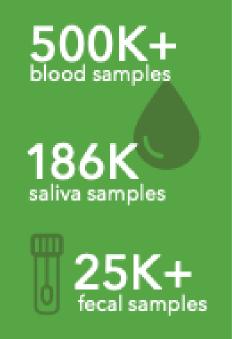


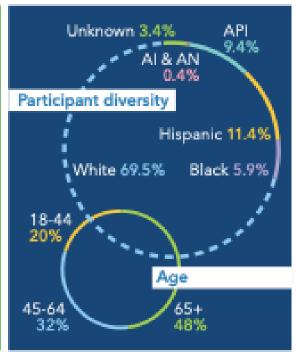
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Questions

