

A open-source, cloudnative platform for biomedical researchers

One place to

- Access data
 - Run analysis tools
- Collaborate

Designed with built-in security so you can focus on science

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Modular

Comprised of functional components with well-specified interface

COMMUNITY FOCUSED

Created by many groups to foster a diversity of ideas

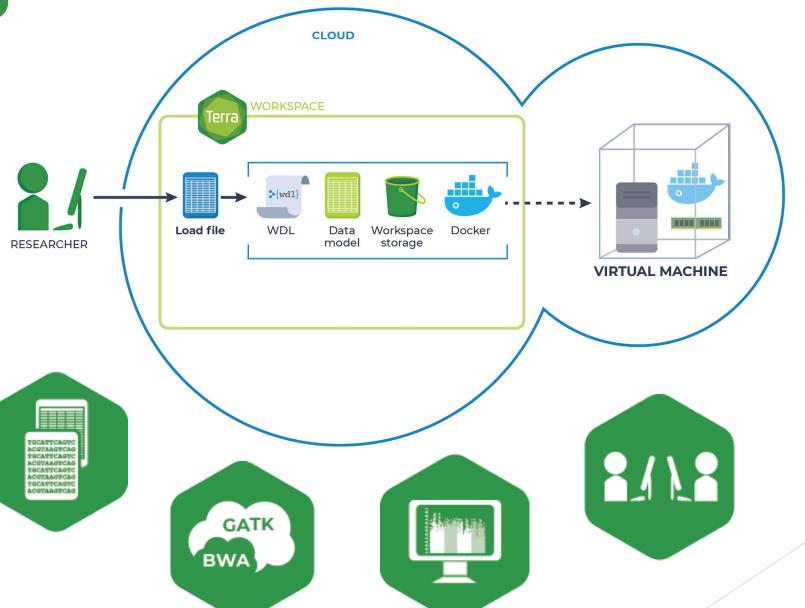
OPEN

Open-source licenses, software, arch to enable extensibility

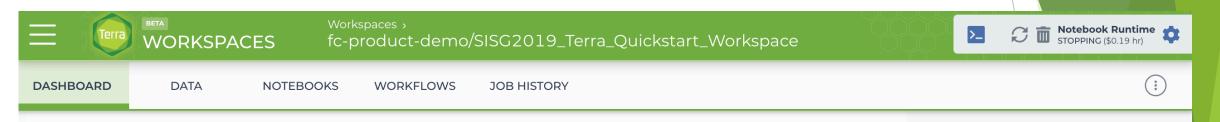
STANDARDS BASED Consistent with standards developed by coalitions such as GA4GH



Core Functionality and organization







ABOUT THE WORKSPACE 🧪

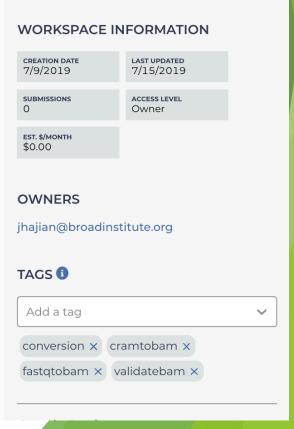
This hands-on practice workspace helps users get started processing and analyzing data on Terra. You'll get familiar with the Data, Notebooks, Workflows, and Job History tabs in your workspace. You'll import BigQuery data from the Data Library and run a quick analysis in a notebook, and run your first workflow. The workspace is intended to be useful for a broad audience, streamlined so you can see results quickly with hands-on practice that will give you the foundation to start your own work on Terra.

Part 1 focuses on using Terra's Data Explorer to create cohorts of data and access and analyze the BigQuery data in a Jupyter notebook. Part 2 focuses on running a simple workflow, including linking genomics data in a Google bucket to the workspace Data table for access by the workflow, and configuring the workflow to run on your data.

Scroll down to the end for links to additional resources. Note that you will need to clone your own copy of this workspace to run the notebooks and workflows.

Part 1: How to use Jupyter Notebooks to access and analyze data, including data in Terra's data library

Notebooks overview





Integrated Data Library, associated Google bucket















Project Baseline by verily

AMP Parkinson's Disease

The Accelerating Medicines Partnership (AMP) is a public-private partnership between the National Institutes of Health (NIH), multiple biopharmaceutical and life sciences companies, and non-profit organizations to identify... READ

Participants: > 4,700

BROWSE DATA



Baseline Health Study

Baseline Health Study is a longitudinal study that will collect broad phenotypic health data from approximately 10,000 participants, who will each be followed over the course of at least four years. The study is part of a broader effort designed to develop a well-defined reference, or "baseline," of

Participants: > 1,500

BROWSE DATA



ENCODE Project

The Encyclopedia Of DNA Elements (ENCODE) project aims to delineate all functional elements encoded in the human genome. To this end, ENCODE has systematically mapped regions of transcription, transcription factor association, chromatin structure and histone modification

Donors: > 650; Files: > 158,000

BROWSE DATA



FireCloud Dataset Library

Search for datasets sequenced at the Broad Institute, or public datasets hosted at the Broad. Datasets are pre-loaded as workspaces. You can clone these, or copy data into the workspace of your choice.

Samples: > 158,629

BROWSE DATASETS



GTEx presented by NIH Commons

The Genotype-Tissue Expression (GTEx) Program established a data resource and tissue bank to study the relationship between genetic variation.



HUMAN CELL

Human Cell Atlas

The Human Cell Atlas (HCA) is made up of comprehensive reference maps of all human cells — the fundamental units of life — as a basis for



Nurses' Health Study

The Nurses' Health Study and Nurses' Health Study II are among the largest investigations into the risk factors for major chronic diseases in



TopMed presented by NIH Commons

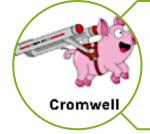
Trans-Omics for Precision Medicine (TOPMed),



Analysis Tools including both interactive and batch processing



Interactive development environment for working with notebooks, code and data. Supports 100 programming languages with 1.7M public notebooks available



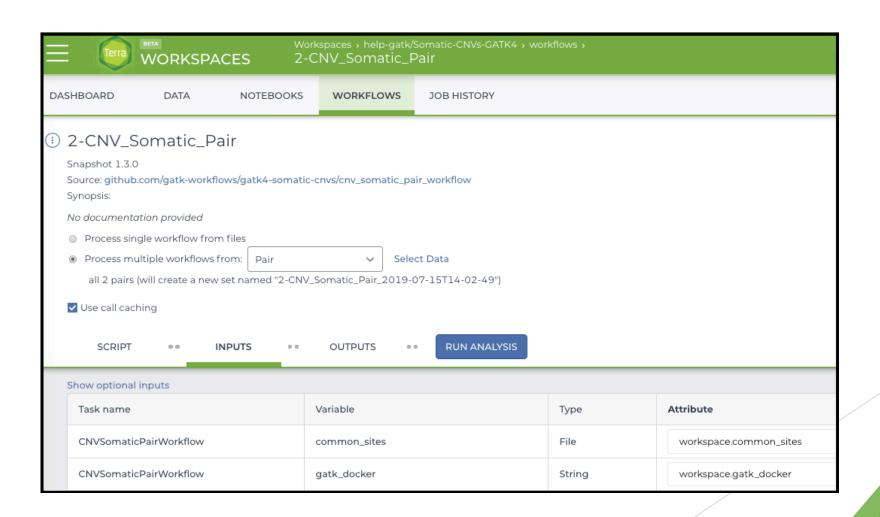
Scientific workflow engine designed for simplicity & scalability. Supports WDL and CWL, the two workflow languages adopted by GA4GH



Open-source, scalable framework for exploring and analyzing genomic data. Hail can generate variant/sample annotations, and perform variant, gene-burden, eQTL association analyses



Workflows to align, QC, call short variants per sample, joint-call across populations, and filter





Jupyter Notebooks for interactive visualization and analysis

"An open-source web application that allows users to create and share documents containing live code, equations, visuals, and narrative text"

Interactive Analysis

- Visualize and iterate in real time
- Couple with BigQuery to analyze data of any size or format in real time

Sharing

- Code and visualization tools, data, and narrative text in one package
- Lets your collaborators easily access and understand your results
- Open and interoperable standards



Our Ecosystem Partners

Terra is product of $\bigotimes \underset{\text{INSTITUTE}}{\text{BROAD}}$ and \bigvee

We are privileged to be part of the



Our goal is to build momentum for an *open, compatible, and* secure approach to data within the larger research community.

Acknowledgements

We are grateful to the how team for their support.



This walkthrough features a reproduction of the subset of analysis in:

Haas, ME et al. (2018) Genetic Association of Albuminuria with Cardiometabolic Disease and Blood Pressure. AJHG volume 103, issue 4, p461-473. doi:10.1016/j.ajhg.2018.08.004

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