

# Running Workflows in the Cloud

BFX Workshop  
Jason Walker, Chris Miller

# Getting data from dbGaP

## 1) Apply for access

- ERA Commons account
- Fill out a Data Access Request (DAR)
  - How are you going to use the data? Read the study information

### The Genetic Landscape of Metastasis and Recurrence in HNSCC (phs001007.v1.p1)

☐ Disease-Specific (Cancer, MDS)  
(phs001007.v1.p1.c1), [NCI DAC](#)

Use of the data must be related to Cancer. Use of the data includes methods development research (e.g., development of software or algorithms). .

Public Posting of [Genomic Summary Results](#) – **Allowed**.

Have data available in Amazon S3 Cloud Storage 

Have data available in Google Cloud Storage 

23

### Up for a Challenge: African American Breast Cancer Consortium (AABC) Study (phs000851.v1.p1)

☐ Up for a Challenge (Publication required)  
(phs000851.v1.p1.c1), [NCI DAC](#)

Use of this data is limited to research described for the National Cancer Institute (NCI) "Up for A Challenge" breast cancer genetic epidemiology competition. The goal of this challenge is to use innovative approaches to identify novel biology involved in breast cancer susceptibility including new genes, genetic variants, or sets of genomic features, leading to novel biological hypotheses. Individuals NOT participating in the challenge would NOT be granted access. Requestor agrees to make results of studies using the data available to the larger scientific community. .

Public Posting of [Genomic Summary Results](#) – **Undefined**.

4881

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- Wait 2-8 weeks for approval

https://gen3.biodatacatalyst.nhlbi.nih.gov/

NIH

National Heart, Lung, and Blood Institute

BioData

CATALYST

Powered by Gen3

Browse Data

Documentation

1CMILLER

Logout

Dictionary

Exploration

Discovery

Workspace

Profile

Data

File

Explorer Filters

Data Tools

Summary Statistics

Table of Records

Data Access

☒ Data with Access

☐ Data without Access

☐ All Data

Filters

Harmonized Variables

Project

Subject

Collapse all

Program

parent

180,102

topmed

59,680

tutorial

14,433

open\_access

3,202

Project Id

Export to Seven Bridges

Export All to Terra

Export to PFB

Export to Workspace

Subjects

257,417

Projects

34

Annotated Sex

female

162,730

(63.2%)

male

22,037

(8.6%)

no data

72,650

(28.2%)

Race

white

56.15%

black or african american

7.7%

other

2.38%

asian

2.08%

american indian or alaska native

0.26%

hispanic

0.16%

native hawaiian or other pacific islander

0.04%

multiple

0%

no data

31.22%

- 400,000 lines of code in the genome/analysis-workflows github repository
- Dozens of data types and approaches
  - exome/WGS/targeted (somatic/germline)
  - bisulfite
  - RNAseq
  - single-cell (TCR, 5'/3', ATAC)
  - RNAseq (expression, fusions, splicing)
  - ATAC/ChIPseq
  - etc

master analysis-workflows / definitions / pipelines /

Go to file Add file ...

johnegarza Merge pull request #988 from johnegarza/immuno\_vcf\_filter\_updates ✓ 77ec4f2 on Jan 21 History

..		
alignment_exome.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
alignment_exome_nonhuman.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
alignment_umi_duplex.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
alignment_umi_molecular.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
alignment_wgs.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
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aml_trio_cle.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
aml_trio_cle_gathered.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
bisulfite.cwl	Changed the documentation to a doc line	2 years ago
chipseq.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
chipseq_alignment_nonhuman.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
detect_variants.cwl	Generalized parameter names and improved documentation	last month
detect_variants_nonhuman.cwl	Add doc line about scatter_count to pipelines.	5 months ago
detect_variants_wgs.cwl	Generalized parameter names and improved documentation	last month
downsample_and_recall.cwl	Removed variant_index variables as not used in GATK4	7 months ago
gathered_downsample_and_recall.cwl	Gathered downsample_and_recall to store gVCFs in per-BAM directories.	14 months ago
germline_exome.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
germline_exome_gvcf.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
germline_exome_hla_typing.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
germline_wgs.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
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immuno.cwl	Merge pull request #988 from johnegarza/immuno_vcf_filter_updates	last month
maseq.cwl	update stringtie to accept indexed bam	last month
maseq_star_fusion.cwl	update stringtie to accept indexed bam	last month
maseq_star_fusion_with_xenosplit.cwl	made inputs for rna-xenosplit consistent with the other rna workflows	last month
somatic_exome.cwl	Merge pull request #988 from johnegarza/immuno_vcf_filter_updates	last month
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somatic_wgs.cwl	Merge pull request #988 from johnegarza/immuno_vcf_filter_updates	last month
tumor_only_detect_variants.cwl	adding docm documentation	5 months ago
tumor_only_exome.cwl	Propagate documentation of sequence inputs to other pipeline CWLs.	last month
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- *Should* be platform-independent
- Workflow systems are complicated
- Our cluster has some unique quirks
- Transition to a new workflow language that fits with our local/GCP model (WDL)
- **griffithlab/analysis-wdls**

master analysis-workflows / definitions / pipelines /

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# Workflows on Google Cloud

- Assumes that:
  - you have already set up a PO and billing and have IT support
  - you have access to the cloud console to grant permissions and such
- <https://github.com/griffithlab/cloud-workflows>