

CANCER GENOMICS CONSORTIUM

Educating for Best Practices in Clinical Cancer Genomics



International Cancer Genome Consortium (ICGC) Data Portal

Data portal: http://dcc.icgc.org/

Project homepage: http://icgc.org/



ICGC Aim Statement

- ICGC Goal: To obtain a comprehensive description of genomic, transcriptomic and epigenomic changes in 50 different tumor types and/or subtypes which are of clinical and societal importance across the globe.
- 89 Cancer Genome Projects supported by ICGC
 - 76 projects on Data Portal
- Media link on Project homepage many PDFs related to this resource





ICGC Data Portal



Q e.g. BRAF, KRAS G12D, DO35100, MU7870, FI998, apoptosis, Cancer Gene Censu

Overview of Data Represented

About Us

The CICCC Data Portal provides tools for visualizing, querying and downloading the data released quarterly by the consortium's member projects.

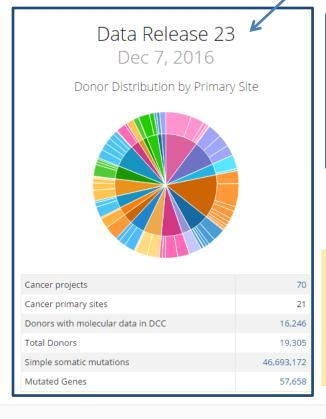
To access ICGC controlled tier data, please read these
instructions.

New features will be regularly added by the DCC development team.

■ Feedback is welcome.



The Pancancer Analysis of Whole Genomes (PCAWG) study is an international collaboration to identify common patterns of mutation in more than 2,800 cancer whole genomes from the International Cancer Genome Consortium.



Tutorial

EXAMPLE QUERIES

- 1. BRAF missense mutations in colorectal cancer
- Most frequently mutated genes by high impact mutations in stage III malignant lymphoma
- Brain cancer donors with frameshift mutations and having methylation data available

Can use as starting point if asking similar question





ICGC data is now available on commercial and academic compute cloud. Read more...

MAIN

Home Cancer Projects Advanced Search Compounds Data Analysis DCC Data Releases

Data Repositories

TOOLS

Genome Viewer ICGC GA4GH Beacor Software

DOCUMENTATION

Portal API
Submission

DCC

The Team Contact Us Twitter

ICGC

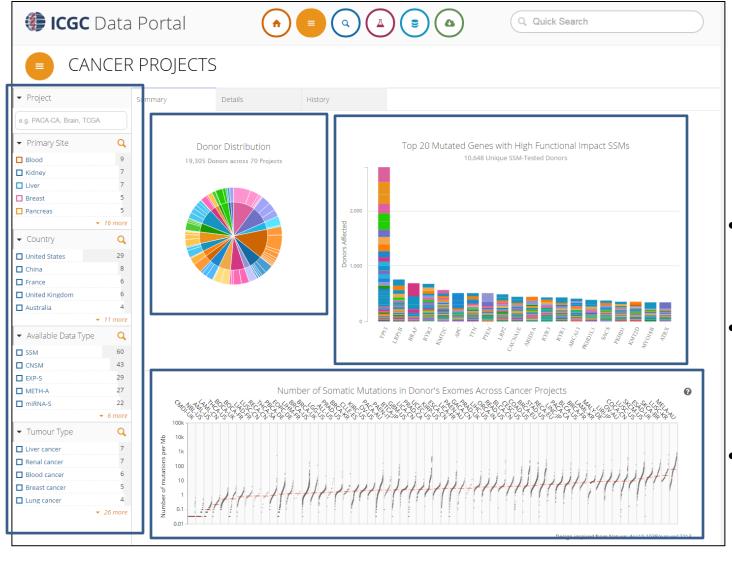
Data Access
Publication Policy
Privacy Policy

Terms and Conditions

CLOUD

About Amazon Collaboratory User Guide

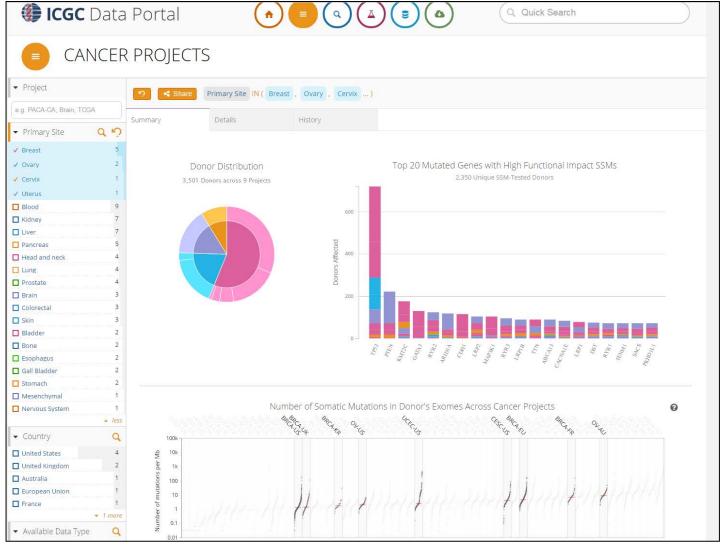




- Search by selecting project, primary site, country conducting study, data types (abbreviations defined by hovering over with mouse), or tumor types
- Search by selecting a primary site or study from pie chart
- See gene in formation by selecting part of bar graph
- See study

 information by
 selecting data set
 from # of somatic
 mutations graph



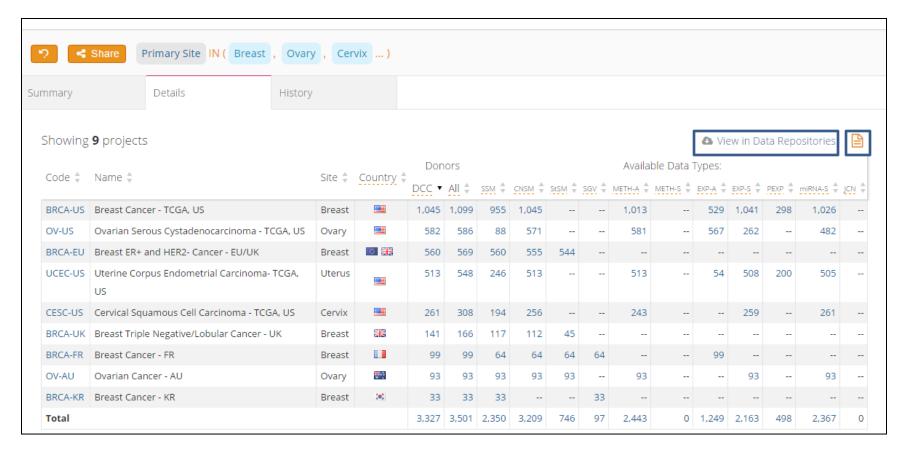


 Selection a subset of studies will update summary page to only include the desired information and help stratify data and identify genes that may be of importance in particular cancer types.



Details Page



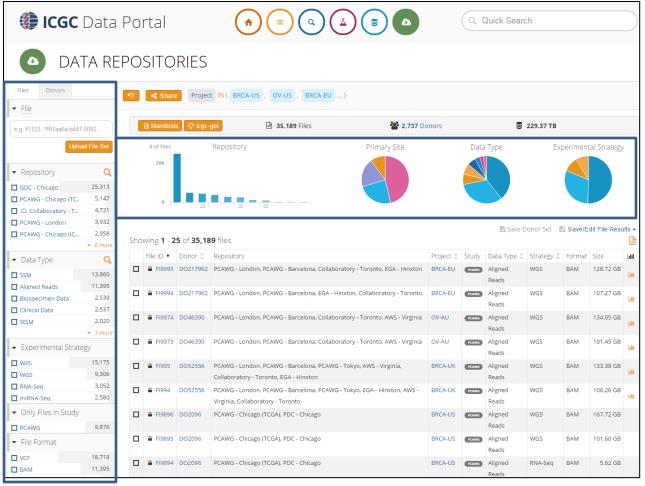


- Table is exportable as .TSV
- Can view data in Data Repositories





Data Repositories



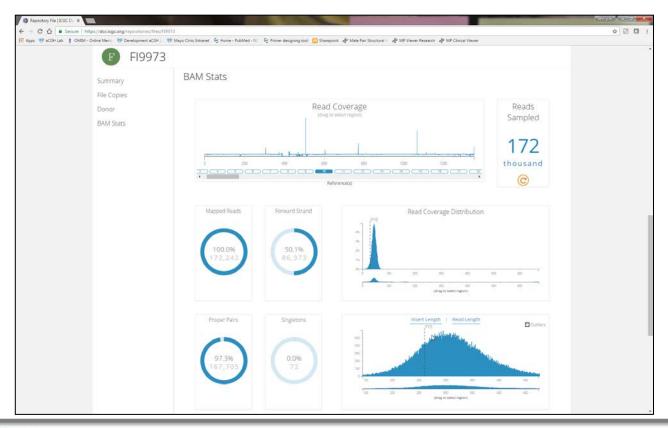
- Left panel: Can select by file, repository, data type, experimental strategy, file format, analysis software, and open/controlled access.
- Can also select by picking a segment of pie chart or bar graph.
- Repositories with Cloud icon have BAM metrics (or similar) available.



Inside Controlled Data



- Downloadable Reference Genome, BAM files
- Visible Quality metrics by chromosome
- Scroll down for % proper pairs, % singletons, % both pairs mapped, % duplicates, and mapping/base quality







Search for Gene or Mutation



Type gene name in search bar

- Start typing in gene name followed by mutation, then select appropriate mutation ID (MU###)
- Under 'Donors' Tab in main panel, find most common primary site and tumor stage for mutation/gene alteration



Search for Gene/Mutation

- On 'Genes' tab in main panel, mutations in table can be analyzed.
 - Oncogrid
 - Enrichment analysis
 - Genome Viewer
 - Save your gene set
 - Download as .TSV



Oncogrid







Can select to view enrichment in Go Molecular/Biological/Cellular or Reactome Pathways

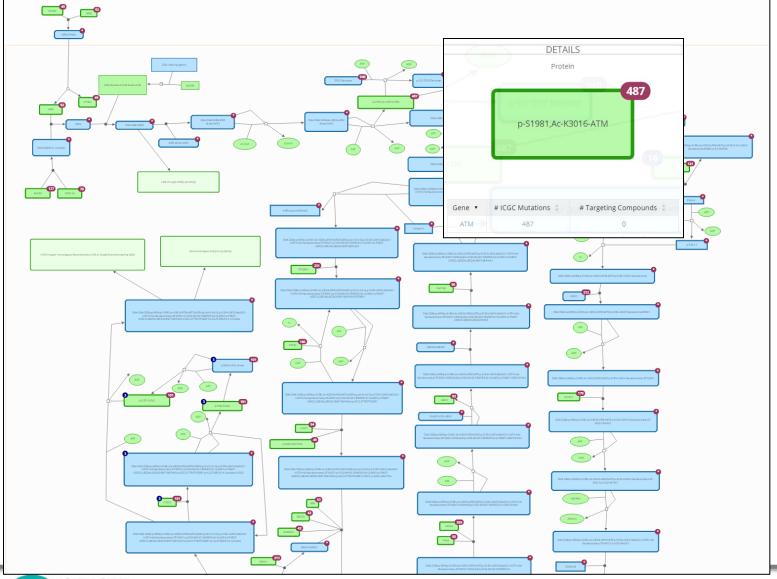
Enrichm	ent Analysis							
Please see the 🗗 En	richment Analysis opcumentation resource for a g	eneral overvi	ew of this feature.					
Overview								
Universe # Gene sets in U		rse # Gene sets in overlap		p # Genes in Universe		# Genes in overlap		# Input genes
Reactome Pathy	vays 2,0	065	9	99 9,775			2	2
							< ∜ Open	In Pathway Viewer
	s out of 58 enriched gene sets with FI							In Pathway Viewer
Top 50 gene se	s out of 58 enriched gene sets with FI		# Genes in overlap	# Donors affected	# Mutations	Expected		
			# Genes in overlap	# Donors affected 220	# Mutations	Expected 2.05e-4		
ID	Name Abnormal conversion of 2-oxoglutarate		# Genes in overlap 1		# Mutations 1		P-Value •	Adjusted P-Value \$
ID R-HSA-2978092	Name Abnormal conversion of 2-oxoglutarate to 2-hydroxyglutarate		# Genes in overlap 1 1	220	1	2.05e-4	P-Value ▲ 2.05e-4	Adjusted P-Value \$ 1.01e-2
ID R-HSA-2978092 R-HSA-389542	Name Abnormal conversion of 2-oxoglutarate to 2-hydroxyglutarate NADPH regeneration	# Genes 1	# Genes in overlap 1 1 1	220 220	1	2.05e-4 2.05e-4	P-Value A 2.05e-4 2.05e-4	Adjusted P-Value \$\displaystar 1.01e-2





Pathway Viewer









Genome Viewer







Scenario #1

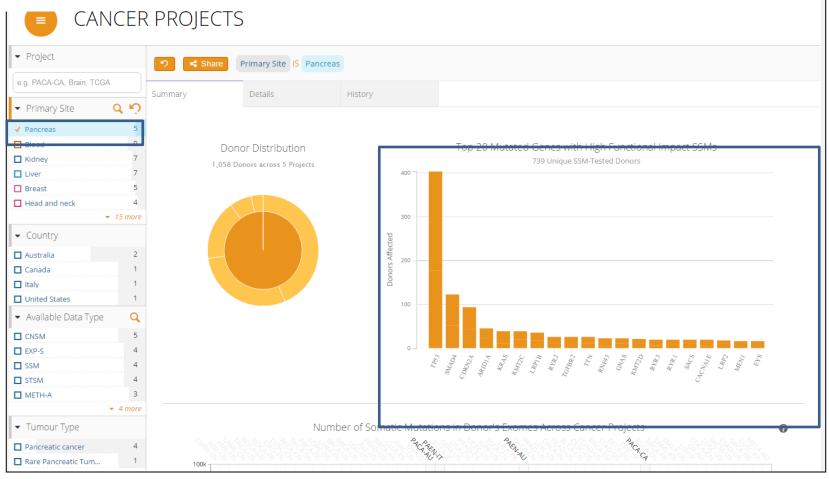
- You are a laboratory professional looking to add new genes to your pancreatic cancer NGS panel. How can you find candidate genes?
 - Find Top Mutated genes
 - Pathway Analysis





Find Top Mutated Genes





• Select Pancreas from Primary Site menu on left panel. Top 20 mutated genes for that cancer type will appear on bar graph.





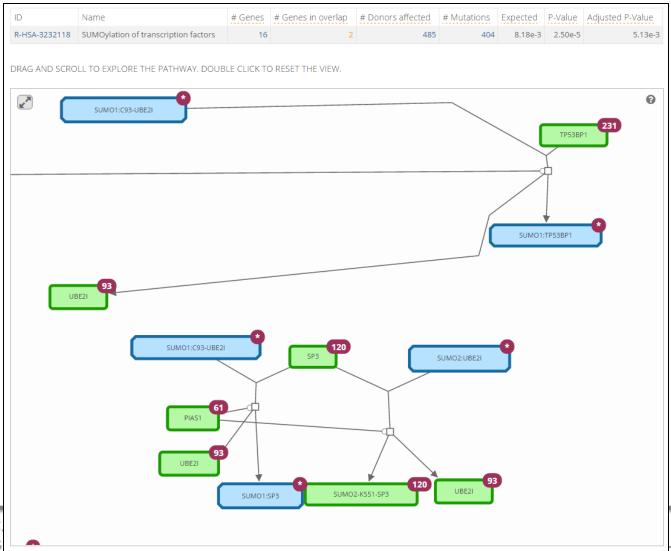
Pathway Analysis

- Input
 - Top 5 mutated Pancreatic Cancer Genes into Advanced Search window under 'Gene' tab on left panel
 - Select 'Pancreas' from primary site on 'Donor' tab on left panel
 - Select "Enrichment Analysis" from above data table on 'Gene' tab on main panel
 - Select Reactome and Open in Pathway Viewer





Pathway Analysis





enomics



Scenario #2

- You would like to know survival statistics and prognostic data for a mutation observed in your laboratory.
 - TP53 MU397
 - chr17:g.7577114C>A





Survival Analysis

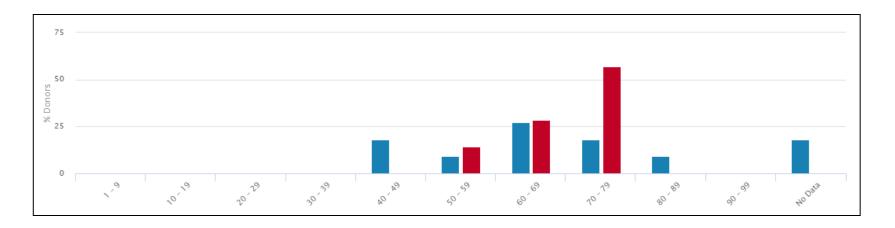






Demographics Available

- Age at Diagnosis
- Vital Status Alive or Deceased
- Gender







Scenario #3

 You are looking to combine your lab's research computing effort with ICGC to better compare data with TCGA data.





Cloud Computing and Collaborative Effort

ICCG Compute Cloud Partners



Amazon Web Services is a well established commercial cloud providing a highly reliable, scalable, low-cost infrastructure platform in the cloud in 190 countries around the world. ICGC datasets are currently hosted at the US East (Northern Virginia) EC2 facility. Read more...



The Cancer Genome Collaboratory is an academic compute cloud resource built by the Ontario Institute for Cancer Research and hosted at the Compute Canada facilities. This infrastructure is still under intensive development and is currently storing only a small subset of the ICGC data for beta testing. Read more...

https://dcc.icgc.org/icgc-in-the-cloud

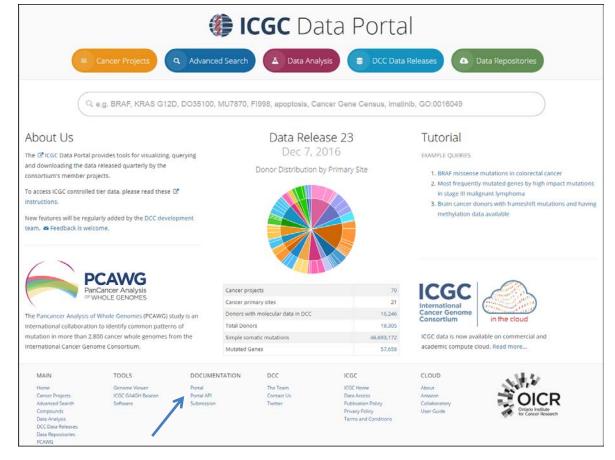


Portal API



API Endpoints are available for trial

and use.







Online Tutorial

https://vimeo.com/75522669





Contact

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