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Date:	Scavenger Hunt		

#### TCGA (Home Page):

The Cancer Genome Atlas (TCGA), founded in December of 2005, is a cancer genomics program hosted by the <a href="National Cancer Institute">National Cancer Institute</a> and the National Human Genome Research Institute. The publicly available data from this project includes <a href="genomic">genomic</a>, epigenomic, <a href="transcriptomic">transcriptomic</a>, and proteomic data. This data was collected from 20,000 different samples that span 33 different cancer types, including breast cancer, which we will be focusing on this semester.

#### **Program History:**

Describe one outcome or impact of TCGA: Cancers of different tissues can share the same alterations and be biologically more similar to each other than to other tumors of		
the same tissue of origin		
Briefly skim the "Timeline & Milestones" page. When did TCGA publish their paper on breast cancer?		
Because TCGA is a public dataset, and one of the first of its kind, they faced some initial concerns		
regarding the ethics of releasing health data to the public. Choose one of the papers in the "Ethics &		
Policies" section to skim. What is one way that your paper addresses these privacy concerns?		
TCGA policies require that all PIs contributing annotated biospecimens provide documentation to the		
Biospecimen Core Resource and the Project Team that their IRBs have approved the use of data		
and specimens for TCGA studies		

#### **TCGA Cancers Selected for Study:**

List three criteria used to	select which cancers to study: poor prognosis, overall public health impact, Availability of samples meeting standards for patient consent
Open the breast ductal carcinoma page and read TCGA's provided background. List one interesting fact	
you found:M	en can also have breast cancer

#### **Publications by TCGA:**

TCGA published (at least) one paper on each of their studied cancer types. These papers, called marker	
papers, include an early analysis of the data, including any molecular characterizations that were	
performed. Read the abstract of the 2012 breast ductal carcinoma cancer paper. List any genes you	
come across (these may be good starting points for your future analyses of this cancer):	
TP53, PIK3CA and GATA3	

#### **Using TCGA:**

Name: _	
Date:	

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sing TCGA (Continued)		
Under the "Program" tab, select just TCGA studies. According to the graph at the top of the page, <u>TP53</u> is the most mutated gene in TCGA projects, affecting approximately <u>35</u> % of cases.		
Return to the GDC Portal home page. Now click the breast image in the diagram to the right of the page. This directs you to the "Exploration" tab and automatically selects all primary sites associated with breast cancers. Now select TCGA as the program, and TCGA-BRCA as the as the project. This is the data we will be focusing on this semester.		
The table on this page shows each patient along with their data. Feel free to explore the data files by clicking on any of the links provided.		
Now explore the Cases, Genes, Mutations, and OncoGrid tabs above the pie charts. What is one takeaway from the plots provided here:ductal and lobular neoplasms is the most common disease type;		
As you can see, the GDC portal provides an overwhelming amount of information. Feel free to continue		

to explore it on your own time!

### **Discussion:**

Think through the following questions, and record your answers below:  1. What is the goal of TCGA?
To provide a publicly available dataset for researchers to have multi-omic data analysis on various cancers
What are some ways that we use TCGA's data for our own cancer research? (Think about the types of data available and brainstorm some research questions that can be proposed given that data.)      Predict or model the mutation rate of TP35/other genes, and the frequency of different kinds of mutation      Does a particular gene affect a particular kind of breast cancer subtype more than the others?
3. What are the benefits and drawbacks of TCGA or other large publicly available datasets?  TCGA has a clear classification of the statistics of different cancer subtypes, and provide us with data from different -omics aspects. But TCGA lacks the normal sequencing data that can be compared with the tumor data. We need to search for other datasets such as GTEx for more data.