## Gene Sample Gene Interaction Network

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Read in a sample BRCA file to get an idea of what it looks like. May clean it a little.

STRING general network maps close to 19000 genes in the BRCA file. Save the list of mapped genes

```
string_db = STRINGdb$new(version="11", species=9606, score_threshold=200, input_directory="")
# 18812 distinct BRCA genes can be mapped
mappable_genes =
   string_db$map(sample_brca_data, "gene_name", removeUnmappedRows = T) %>%
   distinct(gene_name, .keep_all = TRUE) %>%
   distinct(STRING_id, .keep_all = TRUE)
```

```
## Warning: we couldn't map to STRING 68% of your identifiers
gene_list = mappable_genes$gene_name
write_lines(gene_list, file = "~/Desktop/Columbia/Spring_2022/P8139-Statistical_Genetic_Modeling/Project
```

Randomly sample 2000 mapped genes from the sample BRCA file, and save the indices of those sampled

```
# identify the 18812 distinct genes on the BRCA data
sample_brca_data =
    sample_brca_data %%
    mutate(gene_name = toupper(gene_name)) %>%
    filter(gene_name %in% gene_list) %>%
    distinct(gene_name, .keep_all = TRUE) %>%
    arrange(gene_name)

# sample 2000 from the 18812 distinct genes
set.seed(2022)
sample_ind = sample(1:nrow(sample_brca_data), 2000)
write_lines(sample_ind, file = "~/Desktop/Columbia/Spring_2022/P8139-Statistical_Genetic_Modeling/Proje
```

Attached the STRING ids onto the 2000 sampled BRCA genes

```
sample_brca_data =
  inner_join(sample_brca_data, mappable_genes) %>%
  arrange(gene name) %>%
  filter(row_number() %in% sample_ind)
## Joining, by = c("gene_name", "tpm_unstranded")
Find the interaction terms for the 2000 genes
gene_int = string_db$get_interactions(sample_brca_data$STRING_id) %>% distinct(.keep_all = TRUE)
Because this sample BRCA data contains all 2000 genes and their reference STRING ids, make use of it and
create a dictionary to translate STRING_ids to gene name
h = hash()
for(i in seq_len(nrow(sample_brca_data))){
  h[[sample_brca_data[i, 3]]] = sample_brca_data[i, 1]
Translate the interaction matrix and save
for(i in seq_len(nrow(gene_int))){
  gene_int[i, 1] = h[[gene_int[i, 1]]]
  gene_int[i, 2] = h[[gene_int[i, 2]]]
write_csv(gene_int, file = "~/Desktop/Columbia/Spring_2022/P8139-Statistical_Genetic_Modeling/Project/g
```

Build an empty sample gene correlation matrix

```
# create an empty colunn
S_0 = matrix(0, nrow=length(sample_brca_data$gene_name), ncol=length(sample_brca_data$gene_name)) %>% d
# name the columns and rows
col_names = sort(sample_brca_data$gene_name)
names(S_0) = col_names
rownames(S_0) = col_names
write_csv(S_0, file = "~/Desktop/Columbia/Spring_2022/P8139-Statistical_Genetic_Modeling/Project/s0_zer
```