

# Double Trouble: Understanding Sex Differences in Synthetic Lethal interactions in Human Cancers

## Project Updates

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# Recall: The Combined Inactivation of Two Genes Can Lead to Synthetic Lethal Interactions

- ▶ Synthetic lethal interactions describe the relationship between two genes whose coupled inactivation, but not their individual inactivation, causes cell death or reduces cell viability

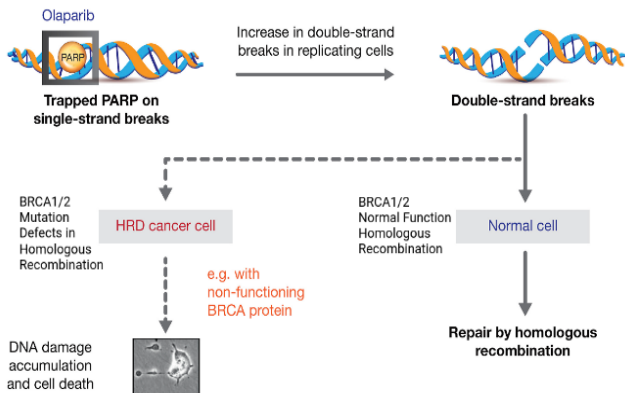


- ▶ Inactivation: Preventing or disabling normal function of a gene (e.g. mutation)

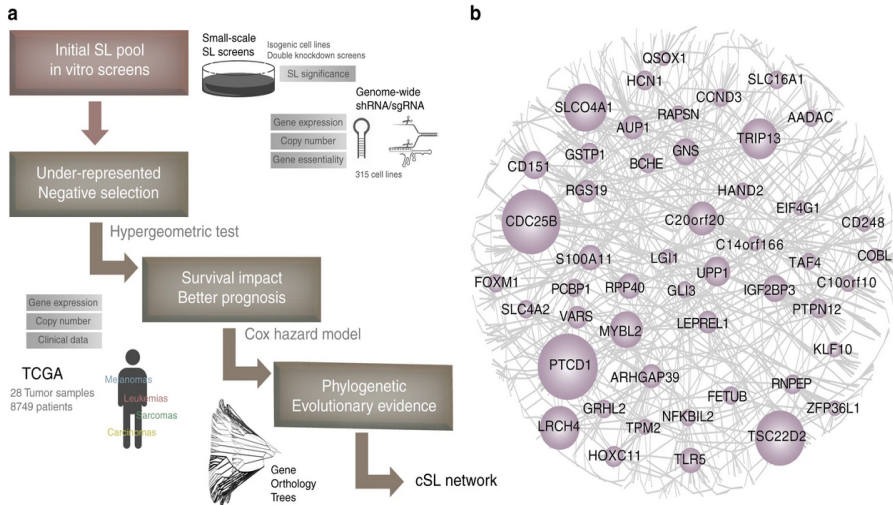
**Synthetic = Combining**

# Recall: Synthetic Lethal Interactions are Harnessed for Precision Oncology

- ▶ Four FDA approved anti-cancer drugs are Poly [ADP-ribose] polymerase 1/2 (PARP1/2) inhibitors that work via a synthetic lethal mechanism



# Recall: Building Pan-Cancer Synthetic Lethality Networks



# Recall: Sex Differences Add an Additional Layer of Complexity

Human sex differences are mainly caused by;

- 1 Gonadal hormone secretions
- 2 Genes located on the sex chromosomes (X and Y)

This leads to differences in the frequency of certain cancer types and the efficacy of treatments in males and females

# The Objective

Can we build sex-specific synthetic lethality networks for various cancer types?

More specifically, we are trying to elucidate the differences in synthetic lethal interactions between males and females using a network based approach.

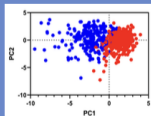
# Overall Project Workflow

## DATA COLLECTION

	Sample 1	Sample 2
Gene 1	6	3
Gene 2	1438	739
Gene 3	2361	1852
Gene 4	400	951
Gene 5	299	142

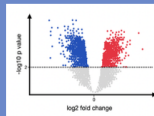
Obtain RNA-seq data from TCGA in the form of raw counts. (Genes are features, samples are data points/variables)

## DATA PROCESSING



Normalize and transform raw RNA-seq data, identify sources of variation, batch effects. Do samples cluster according to biological conditions?

## DIFFERENTIAL EXPRESSION



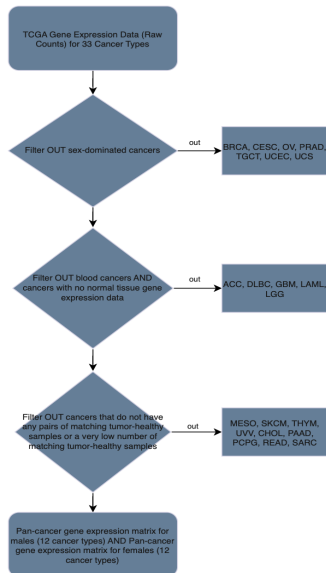
Perform differential gene expression analysis to identify genes that are differentially expressed in tumor tissue.

## SYNTHETIC LETHALITY



Find potential candidate SL pairs for differentially expressed genes using CRISPR gene essentiality data from DepMap.

# Selection of TCGA Cancer Types



- ▶ There is a lack of healthy tissue RNA-seq samples in the TCGA database (12 cancers with more than 10 pairs in M and F)



# Creating Pan-Cancer Expression Matrices

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# Normalizing Pan-Cancer Expression Matrices

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