

# Project Presentation

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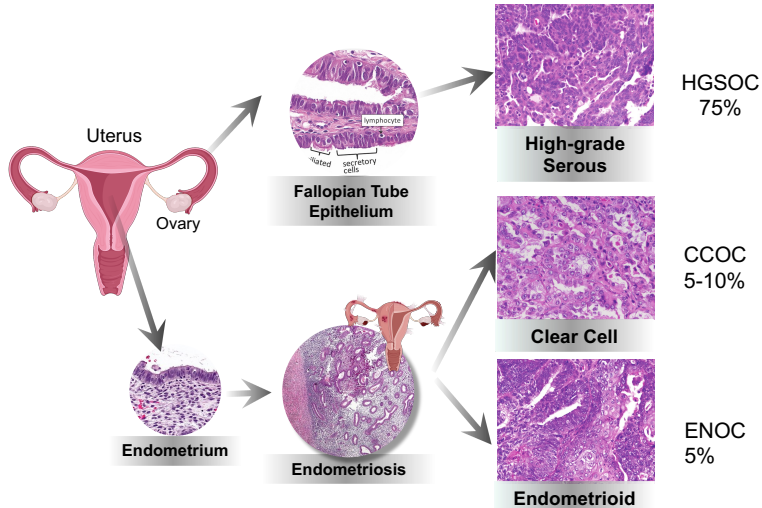
2 December 2025

# Background

# Introduction

- **Group Name:** Packing the Bits
- **Group Members:** Jacob Morrison
- **Project Title:** Investigating Tissue Differences in the Human Fallopian Tube across Ovarian Cancer Subtypes

# Ovarian Cancer Overview

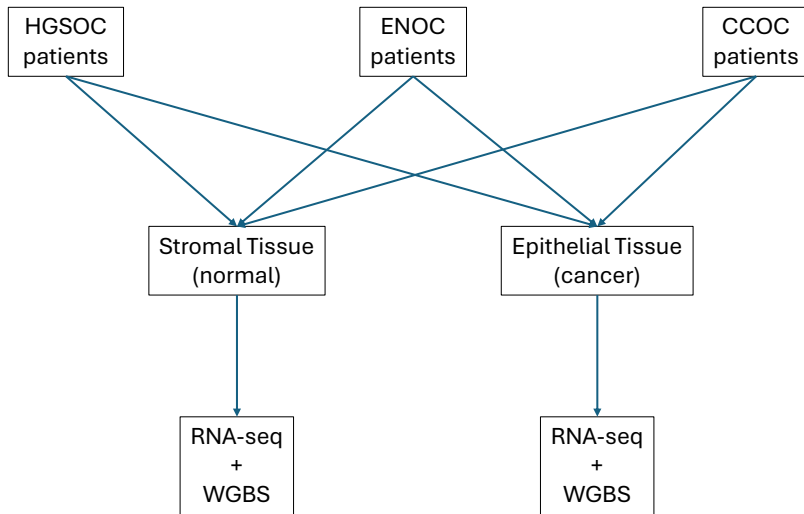


HGSOC=High grade serous ovarian carcinoma | ENOC=Endometrioid Ovarian Carcinoma | CCOC=Clear Cell Ovarian Carcinoma // Photo credit: Paige Matusiak

# Related Work

- Previous results in the lab showed that differences between ENOC and CCOC are driven by cell state rather than cell type
- But, this was done with bulk tumor samples
  - This makes it hard to distinguish between normal cells and cancer cells
- To better understand differences, new tumor samples were collected and laser capture microdissection was performed
  - Allows for precise separation of normal and tumor samples
- Initial analysis showed that differences in the normal tissue was not driven by the type of cancer, but rather by another set of factors

# Sample Overview



# Methods

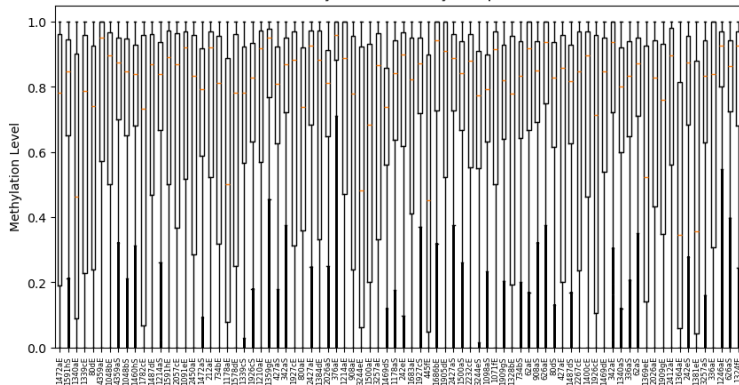
- Sample filtering
  - Started with 92 samples
  - Removed 10 samples due to contamination
- File processing
  - Data read in with `pandas`
  - Raw data points retained and logit-transformed values calculated
  - Data points removed that had coverage  $< 10$  or were not on the canonical chromosomes (chr. 1-22, X, Y, or mitochondria)
  - Only positions found in all samples were retained to eliminate missing data
- Analysis notes
  - Dissimilarity metric used:  $1 - \frac{\vec{x} \cdot \vec{y}}{||\vec{x}|| \times ||\vec{y}||}$ , used the 10,000 most variable positions
  - PCA used the logit-transformed values and retained 2 dimensions ( $\sim 30\%$  of variance explained)



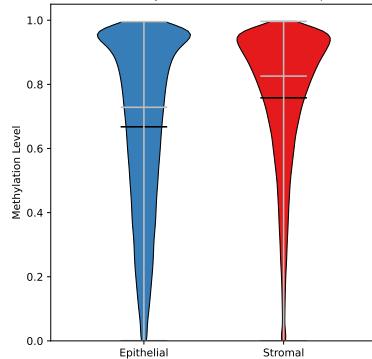
# Results

# Descriptive Statistics

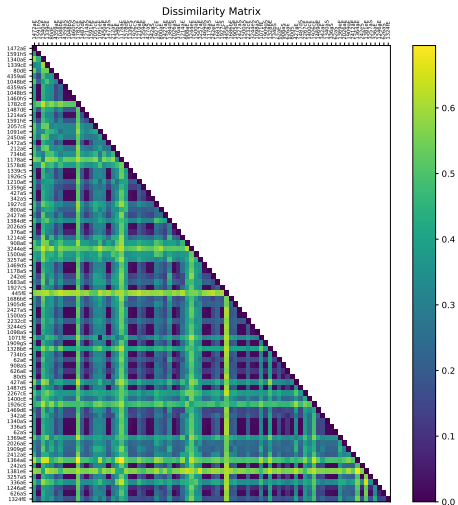
Methylation Levels By Sample



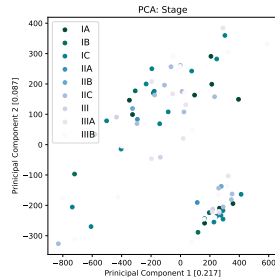
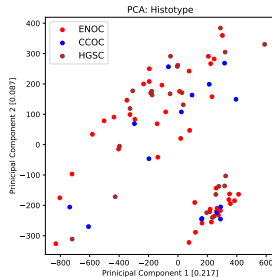
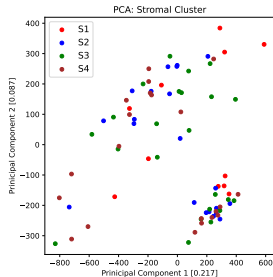
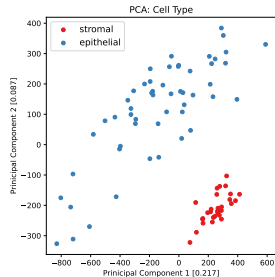
Mean Methylation Levels Across All CpGs



# Dissimilarity



# PCA



# Conclusion

# Summary and Future Work

- Limitations and Future Work
  - Need to overcome differences of cancer versus normal samples → separate stromal and epithelial samples within PCA
  - All data compared across entire genome → look at biologically relevant regions of interest
- Summary
  - Broadly confirmatory of data quality rather than noteworthy results
  - Tumor samples differ from normal samples
  - Some samples do stand out as interesting to explore in more detail