

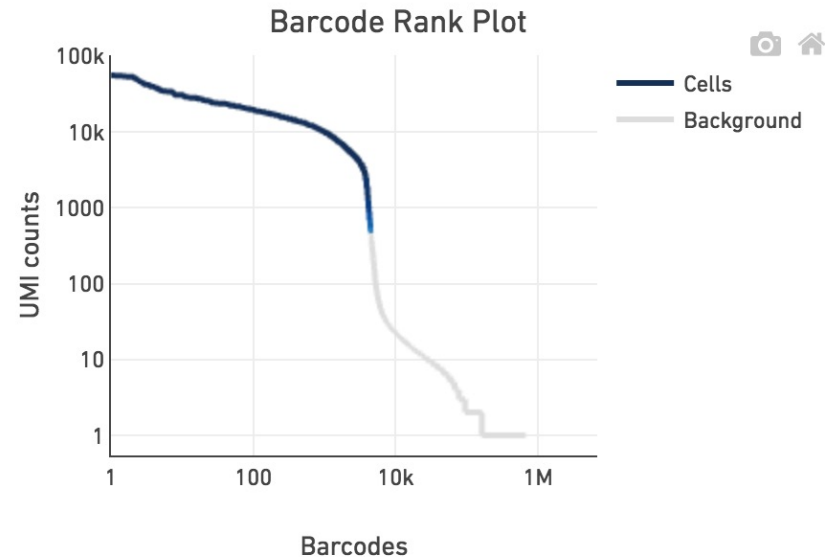
# Single cell transcriptomics

Quality control

# Cell calling

- Doublets: high #UMI/cell
- Dropouts: low #UMI/cell

## Cells ?



# Further cell filtering

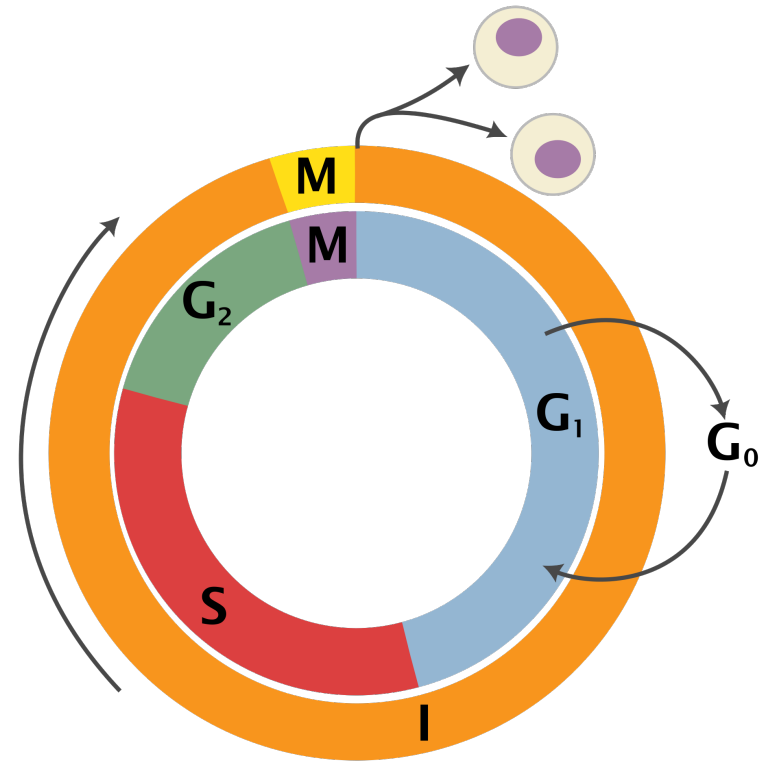
- % UMI in mitochondrial genes:
  - Points to dying/stressed cells
  - Filter by threshold (e.g. 20%)
- Number of detected genes
  - Can also point to dropouts/doublets

# Confounding variables

- % mitochondria
- % ribosomal genes
- % dissociation genes
- Library size
- Sex
- Age
- Cell cycle
- ..

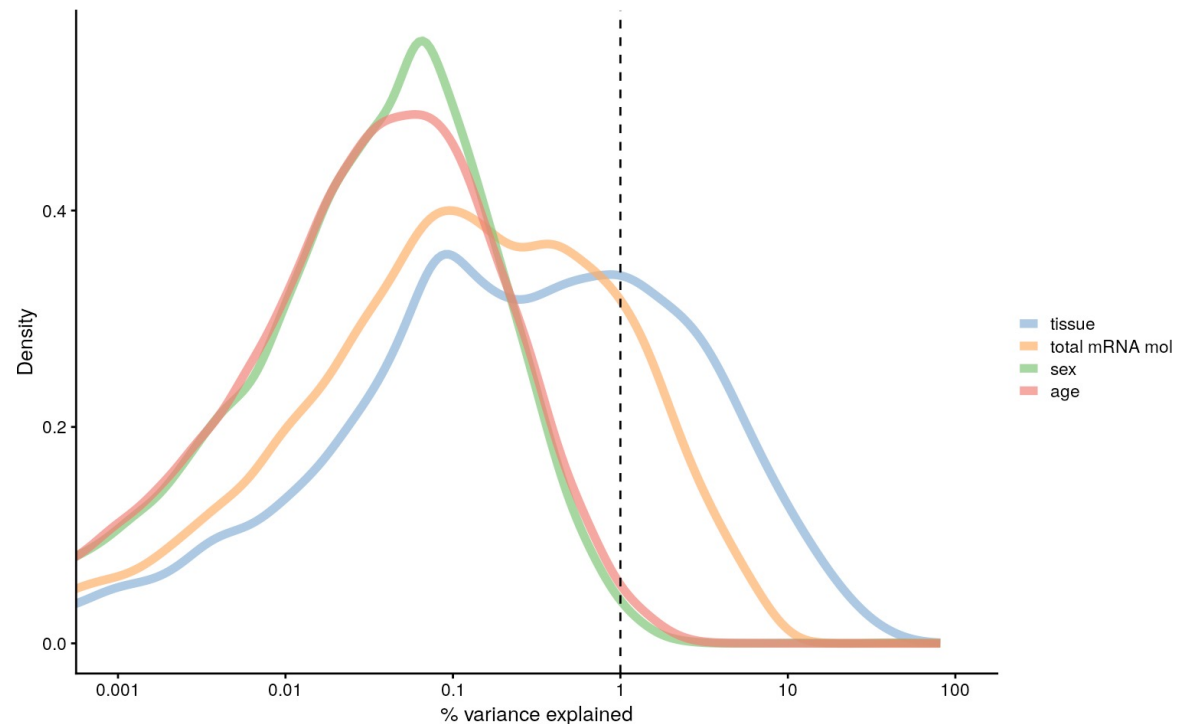
# Cell cycle

- Default clustering might be according to state
- Cycling state can be regressed out for clustering according to cell type



# Find explanatory variables

- With the package scater
- Plot variance explained per gene



# Regress out for clustering

- Regress out variables explaining a lot of variation
- At the scaling step -> affects PCA
- **OR**: integrate data (day 2)