




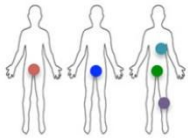
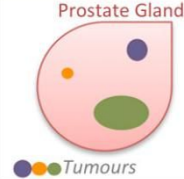
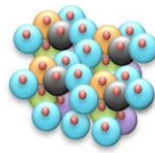
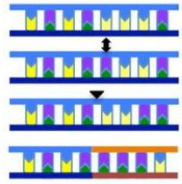
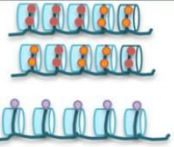
Data-driven precision medicine approaches to cancer

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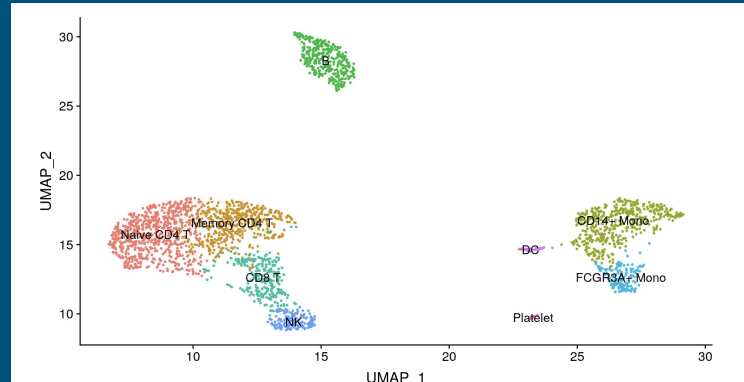
Cancer Background

- Mass of cells
- Heterogeneity

| Types of heterogeneity | | | | |
|--|---|--|--|---|
| Patient | Tumour | Cellular | Genomic | Epigenetic |
|  |  <p>Prostate Gland</p> <p>Tumours</p> |  |  |  <p>● Methylated ● Acetylation ● Unmethylated</p> |
| <p>Different primary tumours in different patients. Different metastatic tumours in different locations.</p> | <p>Different tumours in the same prostate of a single patient (multi-focal disease). Recurrent tumours that have been altered or selected for by treatment.</p> | <p>Different cell types within each tumour mass. These range from stem cells to terminally differentiated cells.</p> | <p>Different mutations (small nucleotide polymorphisms, insertions, deletions or genome rearrangements).</p> | <p>Different methylation and acetylation patterns between the same genes in the same cells, between normal and cancer cells and between different cancer cells.</p> |

Single Cell Sequences

- Seurat - R toolkit for single cell genomics
- Filter based on QC metrics (ex: total number of molecules detected in a cell)
- Identify highly variable features
- Determine dimensionality of dataset (look at p-values)
- Cluster Cells



Patient Data

- Look at DNA
- Compare treatment success
- Gene expression difference



Application

- Identify tumor/aggressiveness
- More effective treatment

Acknowledgements

Thank you

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