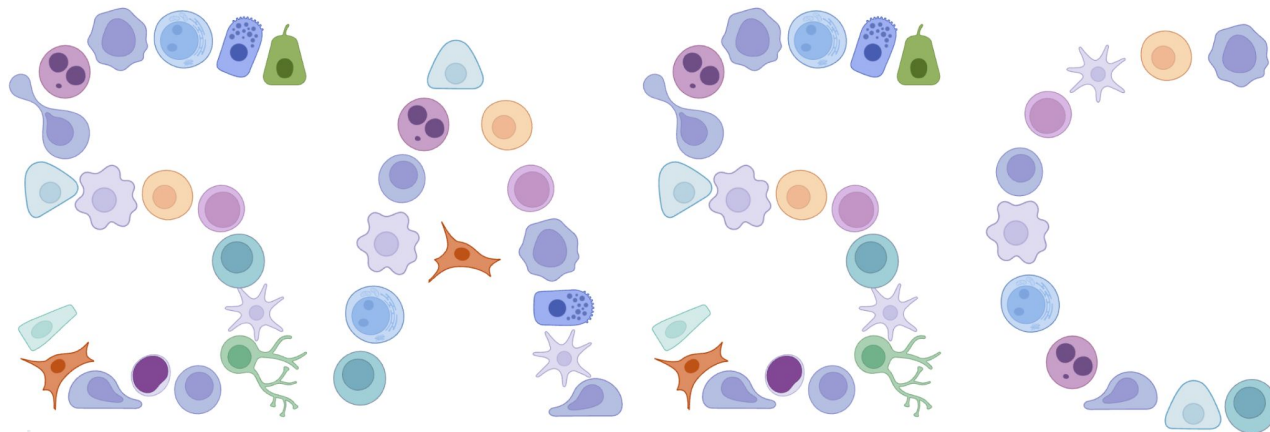


# Welcome to



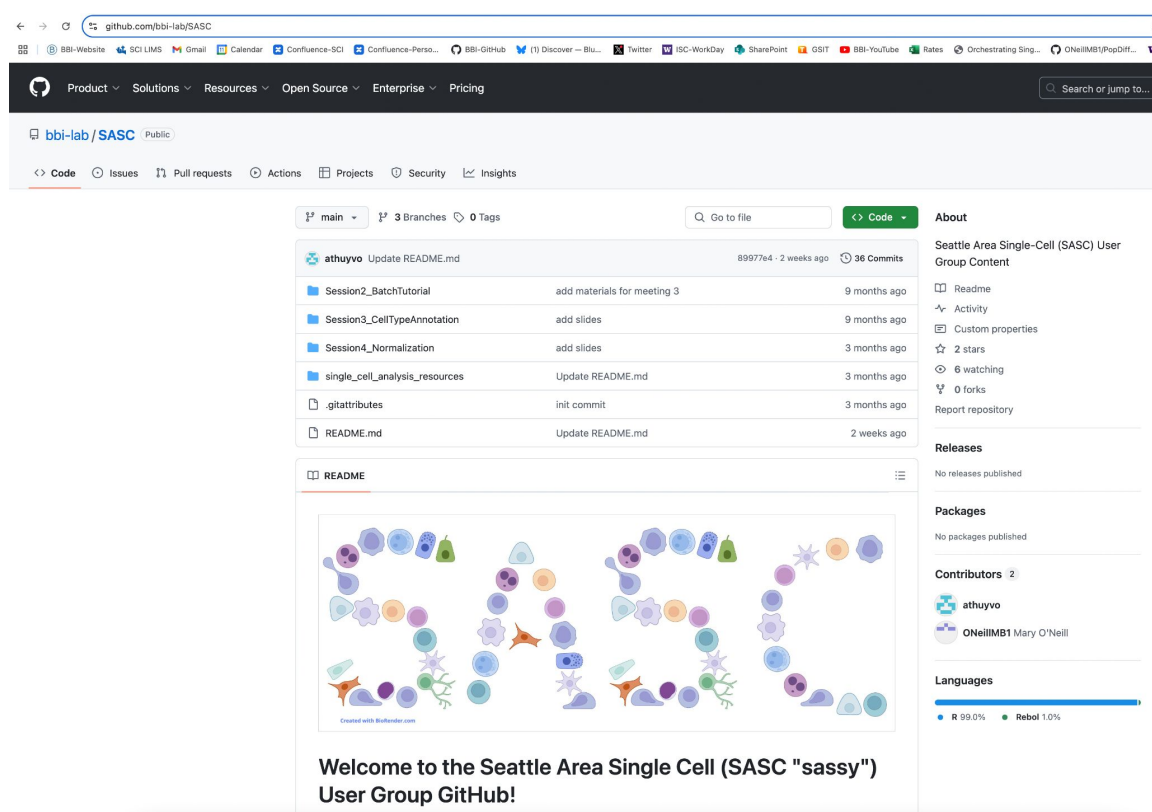
Created with BioRender.com

## Seattle Area Single Cell (SASC) User Group

June 17, 2025



# Github remains our 'website'



github.com/bbi-lab/SASC

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athuyvo Update README.md 89977e4 · 2 weeks ago 36 Commits

Session2\_BatchTutorial add materials for meeting 3 9 months ago

Session3\_CellTypeAnnotation add slides 9 months ago

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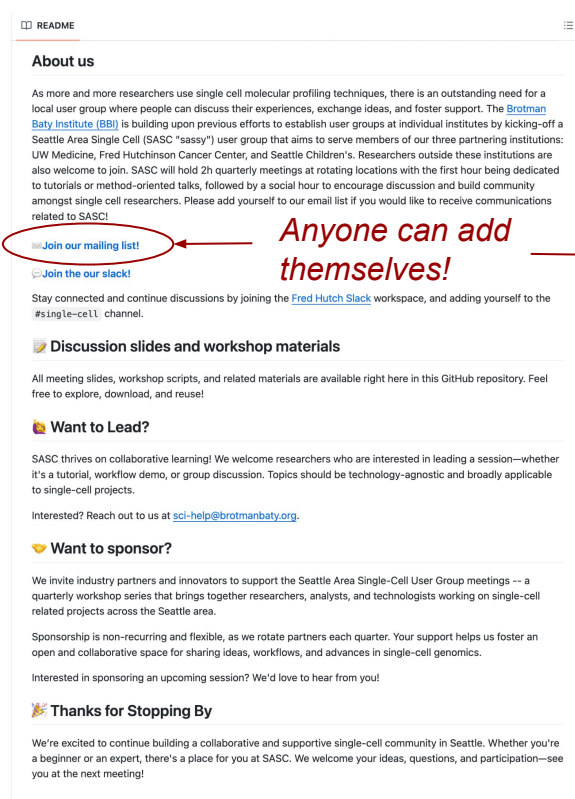
single\_cell\_analysis\_resources Update README.md 3 months ago

gitattributes init commit 3 months ago

README.md Update README.md 2 weeks ago

README

Welcome to the Seattle Area Single Cell (SASC "sassy") User Group GitHub!



README

About us

As more and more researchers use single cell molecular profiling techniques, there is an outstanding need for a local user group where people can discuss their experiences, exchange ideas, and foster support. The [Brotman Baty Institute \(BBI\)](#) is building upon previous efforts to establish user groups at individual institutes by kicking-off a Seattle Area Single Cell (SASC "sassy") user group that aims to serve members of our three partnering institutions: UW Medicine, Fred Hutchinson Cancer Center, and Seattle Children's. Researchers outside these institutions are also welcome to join. SASC will hold 2h quarterly meetings at rotating locations with the first hour being dedicated to tutorials or method-oriented talks, followed by a social hour to encourage discussion and build community amongst single cell researchers. Please add yourself to our email list if you would like to receive communications related to SASC!

[Join our mailing list!](#)

[Join the our slack!](#)

Stay connected and continue discussions by joining the [Fred Hutch Slack](#) workspace, and adding yourself to the #single-cell channel.

Discussion slides and workshop materials

All meeting slides, workshop scripts, and related materials are available right here in this GitHub repository. Feel free to explore, download, and reuse!

Want to Lead?

SASC thrives on collaborative learning! We welcome researchers who are interested in leading a session—whether it's a tutorial, workflow demo, or group discussion. Topics should be technology-agnostic and broadly applicable to single-cell projects.

Interested? Reach out to us at [sci-help@brotmanbaty.org](mailto:sci-help@brotmanbaty.org).

Want to sponsor?

We invite industry partners and innovators to support the Seattle Area Single-Cell User Group meetings -- a quarterly workshop series that brings together researchers, analysts, and technologists working on single-cell related projects across the Seattle area.

Sponsorship is non-recurring and flexible, as we rotate partners each quarter. Your support helps us foster an open and collaborative space for sharing ideas, workflows, and advances in single-cell genomics.

Interested in sponsoring an upcoming session? We'd love to hear from you!

Thanks for Stopping By

We're excited to continue building a collaborative and supportive single-cell community in Seattle. Whether you're a beginner or an expert, there's a place for you at SASC. We welcome your ideas, questions, and participation—see you at the next meeting!

<https://mailchi.mp/668c21581425/seattle-area-single-cell-sasc>

# Special Thanks



*Dr Kevin Lin*

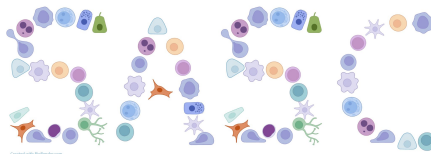


**Fred Hutch**  
Cancer Center

Data Science Lab (DaSL)  
*Sana Hirata*



**BROTMAN BATY**  
INSTITUTE  
Single Cell Genomics Group  
*Anh Vo*



*Everyone Here!*



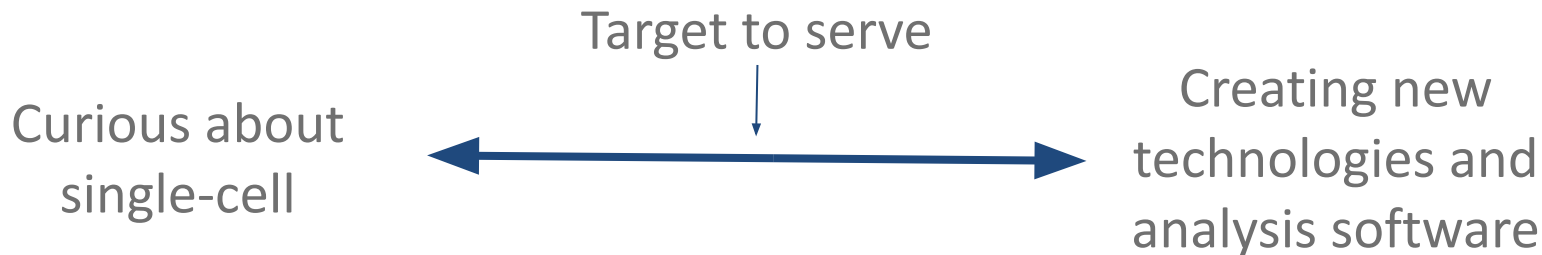
**Element**  
Biosciences

(Today's sponsor for refreshments)  
*Amy Klegarth*



# SASC Goals

- We are striving for this user group meeting be technology (and software) agnostic
  - **Focus on common challenges**





# Disclosures



I do not have any financial relationships to report.



# Agenda

- Brief introduction to multi-ome analysis
- Workshop/tutorial
- Networking (remaining time)
  - Refreshments sponsored by Element Biosciences - thank you!



# Multiome analysis

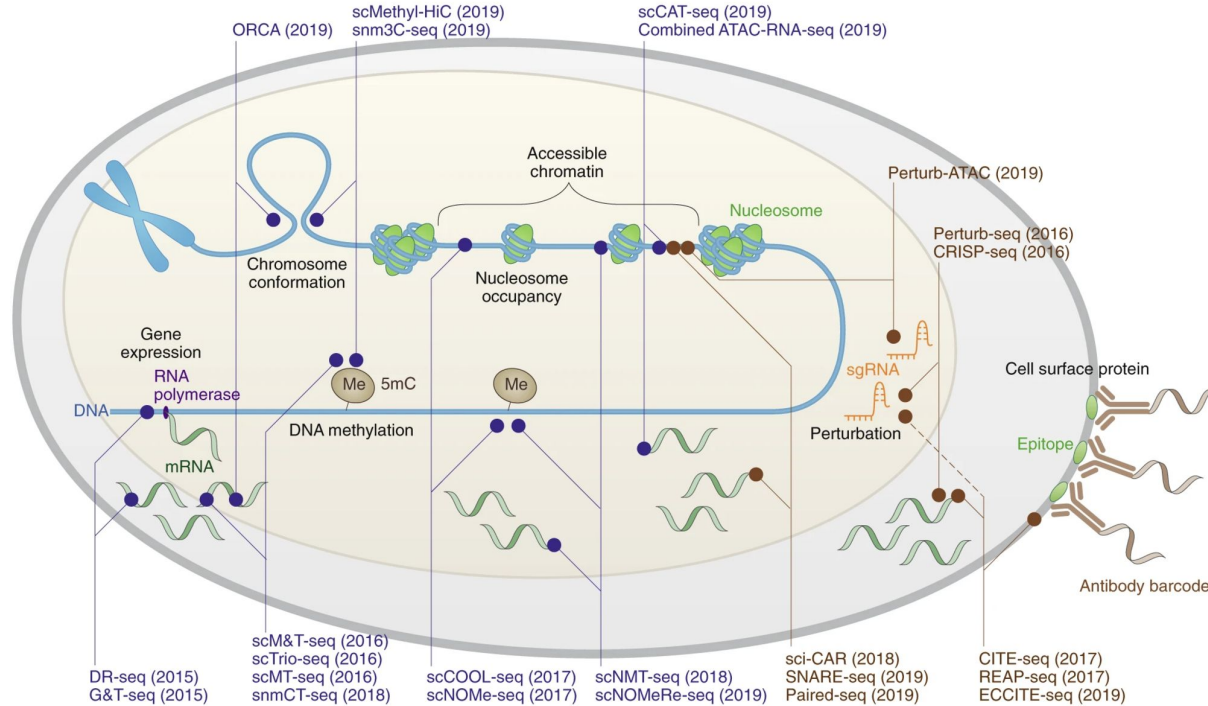


## Just a few words about myself before we get started:

- My lab at UW Biostatistics: developing new computational methods to study cellular mechanisms to advance our understanding of Alzheimer's disease?
  - Main theme of answering temporal questions: 1) Single-cell sequencing is destructive, and 2) brain tissue is mainly post-mortem
- Much of the content today came from the course I taught (BIOST 545/GENOME 545/PHG 545), which I have volunteers helping convert the course content into Quarto (Bookdown) website over the summer

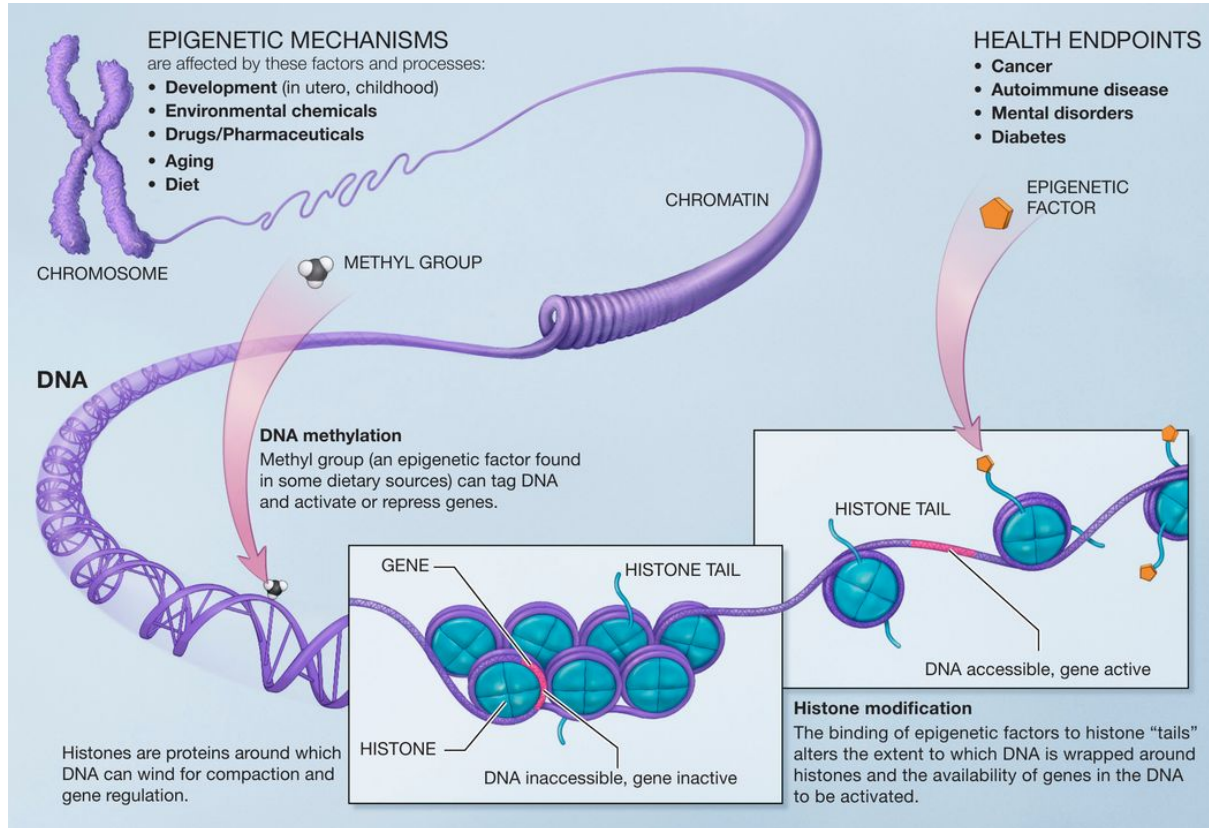


# The wacky world of cell biology



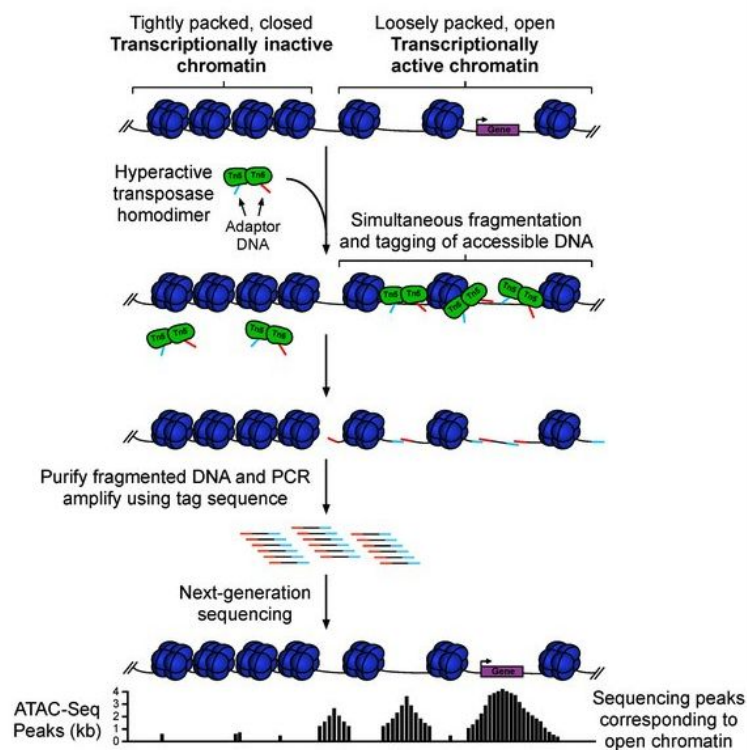
Teichmann and Efremova (2020)

# The wacky world of cell biology



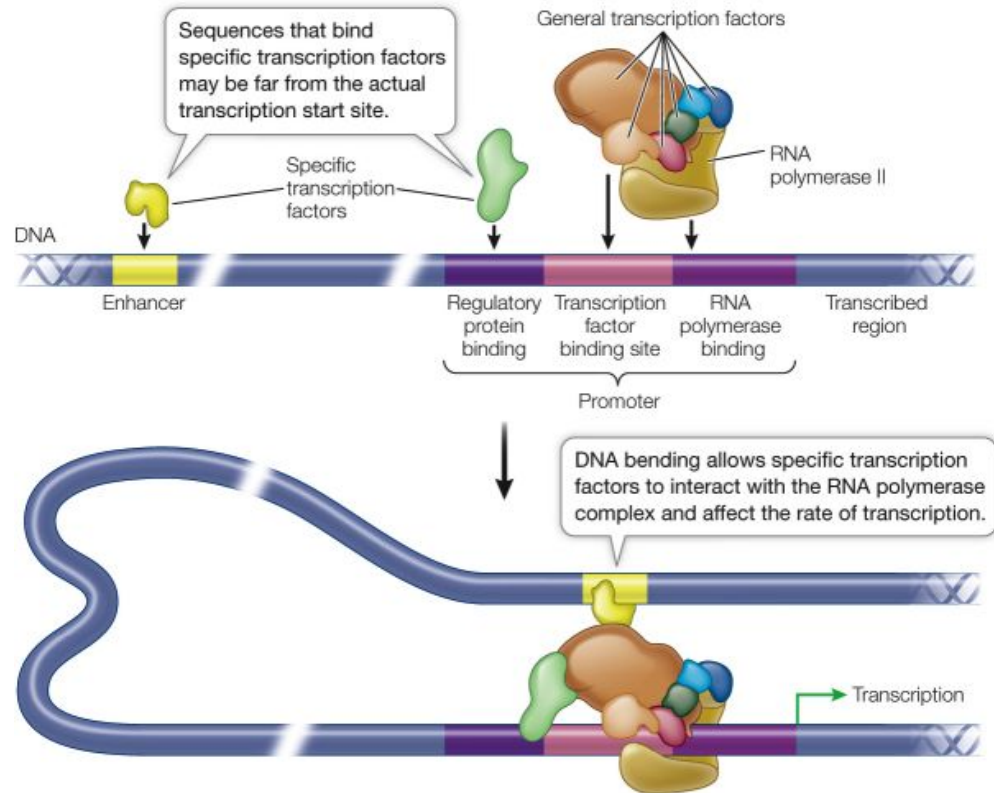
From: <https://en.wikipedia.org/wiki/Epigenetics>

# Accessible DNA



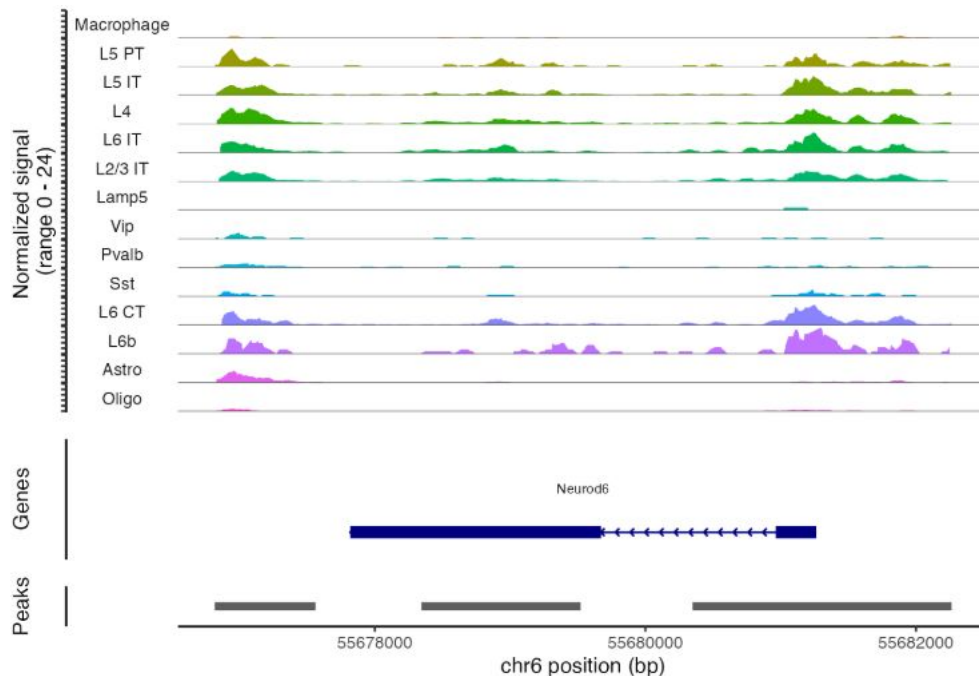
From: <https://www.quickbiology.com/ngs-services/atac-seq-service>

# Transcription factors



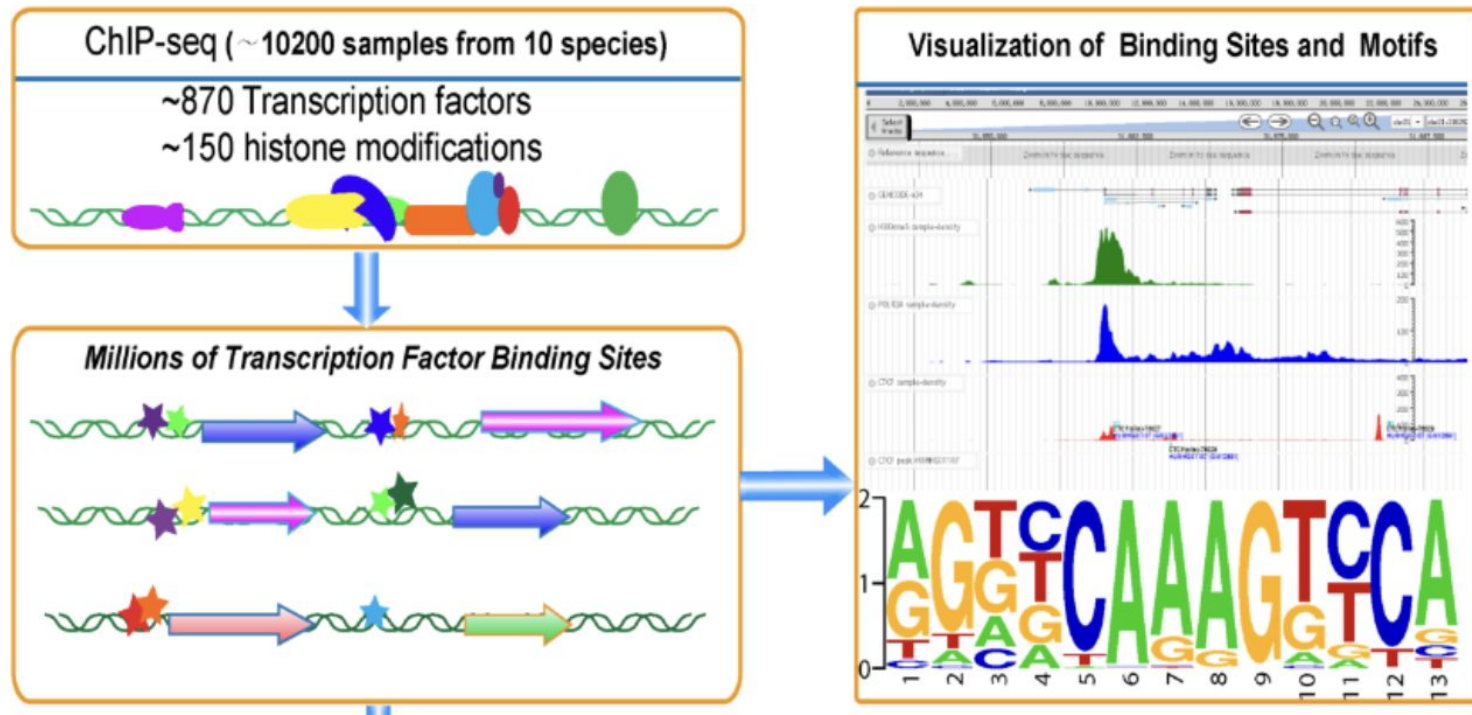
From: [https://digfir-published.macmillanusa.com/life11e/life11e\\_ch16\\_11.html](https://digfir-published.macmillanusa.com/life11e/life11e_ch16_11.html)

# What we're going to cover using ATAC-seq data (Coverage track)



From: [https://stuartlab.org/signac/articles/mouse\\_brain\\_vignette](https://stuartlab.org/signac/articles/mouse_brain_vignette)

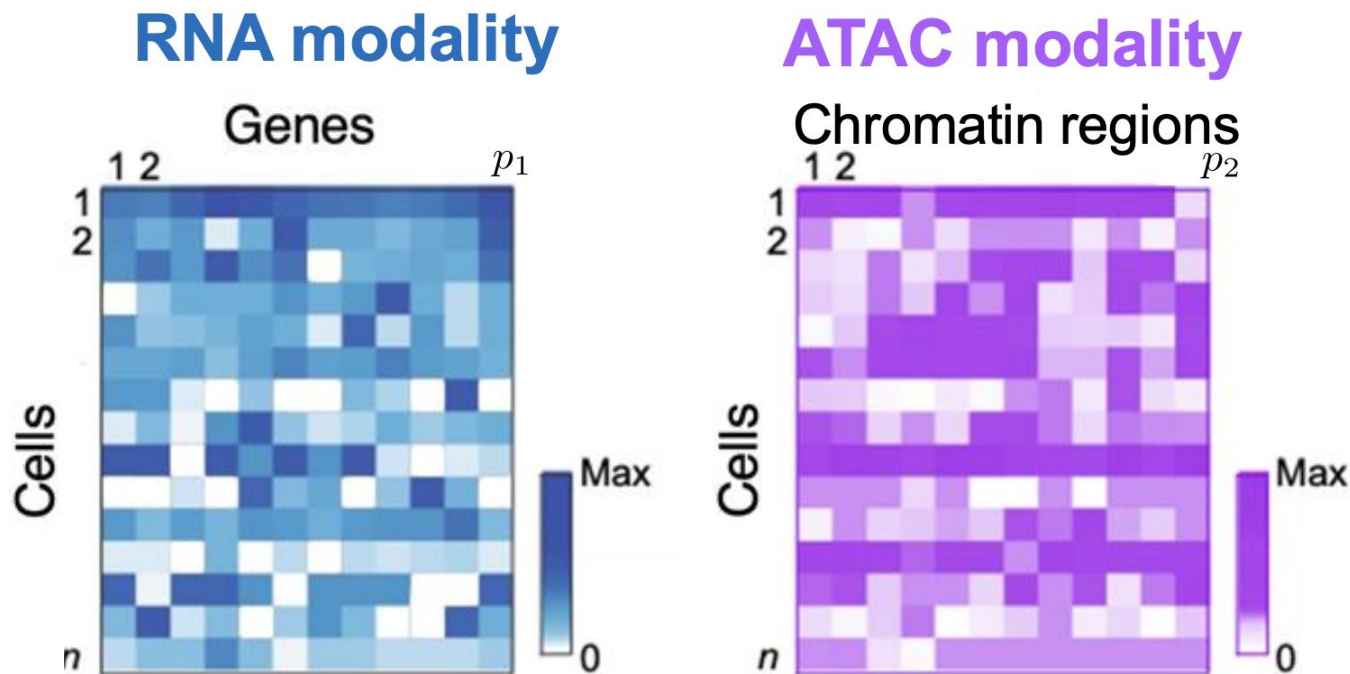
# What we're going to cover using ATAC-seq data (TF motif)



From:

<https://www.seqanswers.com/forum/bioinformatics/bioinformatics-aa/59455-transcription-factor-binding-sites-motifs-and-profiles-from-chip-seq-data>

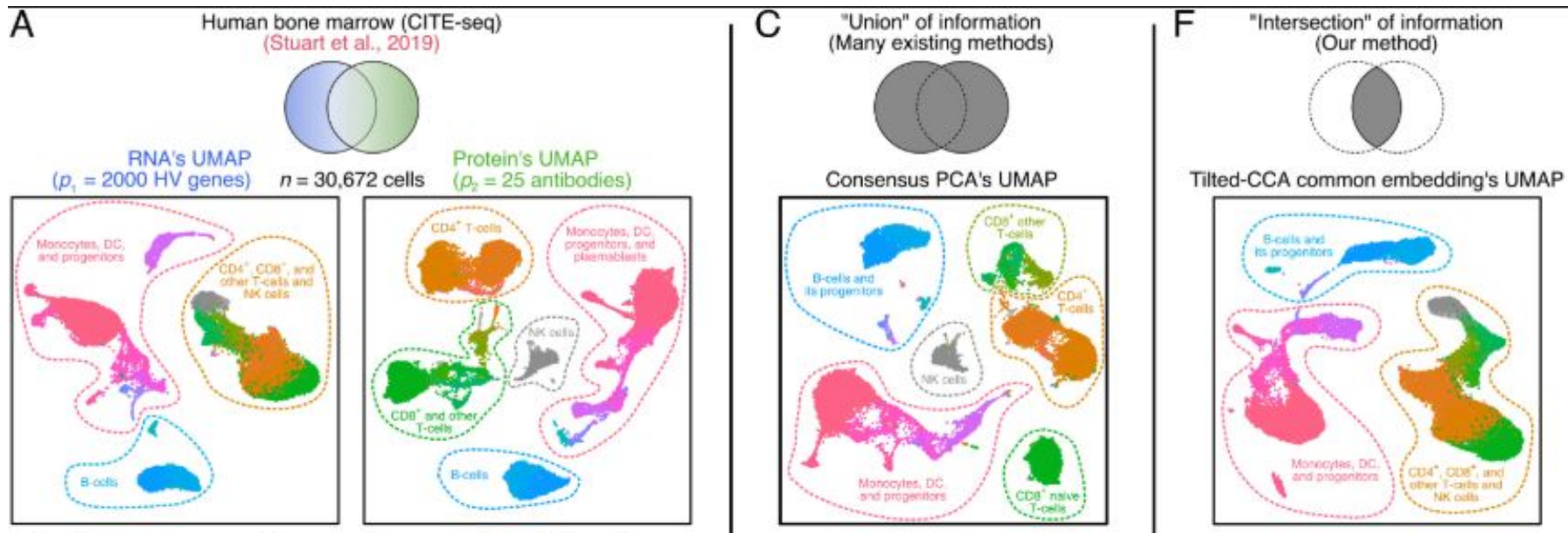
Then, what we'll do with multiome data



Same  $n$  cells, where we have sequence both gene expression (“what programs the cells are using”) and DNA accessibility (“what are the switches that turn on/off programs with a time lag”)



# Then, what we'll do with multiome data (Integration)



**Union:** Give me a “bird’s eye” view of both modalities. (We’ll discuss WNN for this.)

**Intersection:** Tell me what cellular programs are reflected in both modalities, because the complement will tell us what programs in one modality is “invisible” to the other. (We’ll discuss Tilted-CCA for this.)



# Tutorial



- Scripts: [https://github.com/bbi-lab/SASC/tree/main/Session5\\_ATAC-RNA](https://github.com/bbi-lab/SASC/tree/main/Session5_ATAC-RNA)

Additional datasets:

- [http://cf.10xgenomics.com/samples/cell-atac/1.1.0/atac\\_v1\\_adult\\_brain\\_fresh\\_5k/atac\\_v1\\_adult\\_brain\\_fresh\\_5k\\_filtered\\_peak\\_bc\\_matrix.h5](http://cf.10xgenomics.com/samples/cell-atac/1.1.0/atac_v1_adult_brain_fresh_5k/atac_v1_adult_brain_fresh_5k_filtered_peak_bc_matrix.h5)  
(<https://tinyurl.com/huznmdnh>)
- [http://cf.10xgenomics.com/samples/cell-atac/1.1.0/atac\\_v1\\_adult\\_brain\\_fresh\\_5k/atac\\_v1\\_adult\\_brain\\_fresh\\_5k\\_fragments.tsv.gz](http://cf.10xgenomics.com/samples/cell-atac/1.1.0/atac_v1_adult_brain_fresh_5k/atac_v1_adult_brain_fresh_5k_fragments.tsv.gz)  
(<https://tinyurl.com/58yshx2j>)
- [http://cf.10xgenomics.com/samples/cell-atac/1.1.0/atac\\_v1\\_adult\\_brain\\_fresh\\_5k/atac\\_v1\\_adult\\_brain\\_fresh\\_5k\\_fragments.tsv.gz.tbi](http://cf.10xgenomics.com/samples/cell-atac/1.1.0/atac_v1_adult_brain_fresh_5k/atac_v1_adult_brain_fresh_5k_fragments.tsv.gz.tbi)  
(<https://tinyurl.com/yu4aw96h>)



Element  
Biosciences

- *Thank you for sponsoring refreshments this meeting!*

**Teton Atlas OPS 'Grant' due June 20<sup>th</sup>** – just a 250 word blurb to apply – aimed at optical pooled screening, guide RNA sequencing on our low-output kit for in-situ sequencing on adherent cells or cell suspensions.

<https://www.elementbiosciences.com/apply-for-the-2025-teton-atlas-global-grant>

Coming Next on AVITI24 Blog

<https://www.elementbiosciences.com/blog/whats-next-for-aviti24-continued-innovation-for-your-research>

Direct In-Situ Sequencing Webinar On-Demand

<https://www.elementbiosciences.com/resources/webinar-on-demand-from-sample-to-insight-accelerating-discovery-with-direct-in-sample-sequencing-in-cells>

And two pre-prints, one about the technology, the other demonstrating the application in drug testing:

AVITI24 Pre-Print

<https://www.biorxiv.org/content/10.1101/2025.05.03.651997v1>

TKI Resistance Experiment Pre-Print

<https://www.biorxiv.org/content/10.1101/2025.05.06.652479v1>