The Complex Genome: From Structure to Function

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## Overview

In eukaryotes, the genome is extensively compacted inside the nucleus, hierarchically organized at a multiscale of structural units. One of the most critical roles of these delicately regulated structures is the control of transcriptional dynamics. Transcriptional regulations have been recognized as the primary regulatory mechanisms coordinating cellular activities, responses, cell type and fate determinations. No doubt, regulations on transcription are hugely achieved by changes in regulatory factors and structural configuration. Yet, with more than two hundred years of study on the genome, we are just starting to realize the complexity lies within …

The course is designed for students to learn:

1. Basic knowledge of the genomic structure and its role in transcriptional regulation.
2. Cutting-edge techniques and bioinformatics used in the field.
3. Practical experience in bioinformatic analysis.
4. Frontier research principles and methodology.
5. Mysteries and dilemmas remained unsolved.

## Schedule

Class meets online using Zoom/Lark on Tuesday and Saturday at 10:00 am – 12:00 pm (Beijing Time). Class on Saturday, Week 5 (Week 5 – 2\*) is subject to change depending on the actual pacing.

Each class comprises 1 hour of lecture and 1 ~ 1.5 hour of discussions or hands-on sessions, subject to change based on the progress of study.

Literature discussion is scheduled for each week. A research article and a related synopsis will be posted one or half week prior to the discussion. Students should read the article and complete the synopsis before the discussion. For each discussion, 1 ~ 2 students will be selected as discussion leader to guide the discussion.

Week 2, 4 will have hands-on sessions, which requires Win/Mac setup, specific details to be determined.

Week 5 is scheduled for research proposal project. Recommended topics will be provided on week 3. However, students can pick other topics related to the course and discuss with the instructor no later than week 9.

## Syllabus

### Week 1 – 1

**DNA, Chromatin, and Transcription**

**Introduction to the structure basis of DNA, nucleosomes, chromatins, and nucleus; Histone modifications and DNA methylations; Elements of genes and the transcription program; Cis- and trans-regulation.**

* Additional readings: Perspectives on transcription regulation

Lis, J. T. A 50 year history of technologies that drove discovery in eukaryotic transcription regulation. Nat. Struct. Mol. Biol. **26**, 777-782 (2019).

### Week 1 – 2

**Methods to Discovery**

**Quantification of gene expression (from RNA to proteins); Functional analysis of genes: fluorescent reporter assay, in situ hybridization, genetic perturbations (RNAi, CRISPR/Cas system, and protein degrons); Introduction to omics.** Principle of paper discussion

* Discussion paper 1: Protein degrons

Nishimura, K., Fukagawa, T., Takisawa, H., Kakimoto, T. & Kanemaki, M. An auxin-based degron system for the rapid depletion of proteins in nonplant cells. Nat. Methods **6**, 917-922 (2009).

* Following paper: Advanced protein degrons

Yesbolatova, A. et al. The auxin-inducible degron 2 technology provides sharp degradation control in yeast, mammalian cells, and mice. Nat. Commun. **11**, 5701 (2020).

### Week 2 – 1

**Sequence, Omics, and Bioinformatics I**

**Three generations of sequencing; Sequence alignment; Genomics and genome element (GWAS); Transcriptomics (RNA-seq & nascent RNA-seq).** Paper 1 discussion; Hands-on session: Sequence alignment & database searching.

* Discussion paper 2: Loop extrusion model

Fudenberg, G. *et al.* Formation of Chromosomal Domains by Loop Extrusion. *Cell Rep.* **15**, 2038-2049 (2016).

* Additional readings: Review of high throughput sequencing

Reuter, Jason A., Spacek, D. V. & Snyder, Michael P. High-Throughput Sequencing Technologies. Mol. Cell **58**, 586-597 (2015).

### Week 2 – 2

**Sequence, Omics, and Bioinformatics II**

**Functional annotation (GO & KEGG); Epigenomics (ATAC-seq, ChIP-seq, BS-seq); Hi-C; Screening and lineage tracing.** Hands-on session: ChIP-seq analysis using Galaxy.

* Additional readings: Reviews of epigenetic technologies

Rivera, Chloe M. & Ren, B. Mapping Human Epigenomes. Cell **155**, 39-55 (2013).

Mehrmohamadi, M., Sepehri, M. H., Nazer, N. & Norouzi, M. R. A Comparative Overview of Epigenomic Profiling Methods. Front. Cell Dev. Biol. **9** (2021).

### Week 3 – 1

**Transcriptional Regulation I: Enhancers and Transcription factors**

**­Role of enhancers and TFs in transcription initiation, pause-and-release, etc.; Study higher-dimensional architecture of genomes; Model of enhancer-promoter interactions; Enhancer RNA.** Paper 2 discussion

* Discussion paper 3: Cohesin and chromatin structure

Rao, S. S. P. et al. Cohesin Loss Eliminates All Loop Domains. Cell **171**, 305-320.e324 (2017).

* Parallel study: CTCF and chromatin structure

Nora, E. P. et al. Targeted Degradation of CTCF Decouples Local Insulation of Chromosome Domains from Genomic Compartmentalization. Cell **169**, 930-944.e922 (2017).

* Additional readings: Reviews of epigenetic technologies

Haberle, V. & Stark, A. Eukaryotic core promoters and the functional basis of transcription initiation. *Nat. Rev. Mol. Cell Biol.* **19**, 621-637 (2018).

Andersson, R. & Sandelin, A. Determinants of enhancer and promoter activities of regulatory elements. *Nat. Rev. Genet.* **21**, 71-87 (2020).

Li, W., Notani, D. & Rosenfeld, M. G. Enhancers as non-coding RNA transcription units: recent insights and future perspectives. Nat. Rev. Genet. **17**, 207-223 (2016).

### Week 3 – 2

**Transcriptional Regulation II: Epigenetics**

**Epigenetic landscape of the genome; Epigenetic marks of gene activation and silencing; Regulation and dynamics of epigenetics and chromatin remodeling; Epigenetics in development.** Paper 3 discussion

* Discussion paper 4: Phase separation at enhancer regions

Boija, A. et al. Transcription Factors Activate Genes through the Phase-Separation Capacity of Their Activation Domains. *Cell* **175**, 1842-1855.e1816 (2018).

* Additional readings: Reviews of epigenetic regulation & profile of histone marks

Klemm, S. L., Shipony, Z. & Greenleaf, W. J. Chromatin accessibility and the regulatory epigenome. *Nat. Rev. Genet.* **20**, 207-220 (2019).

Barski, A. *et al.* High-Resolution Profiling of Histone Methylations in the Human Genome. *Cell* **129**, 823-837 (2007).

### Week 4 – 1

**Transcriptional Regulation III: Condensates and Bursting?**

**Lipid-lipid phase separation in cells; Novel (yet skeptical) theory of condensates/ clusters/hubs; Dynamics of condensate; Transcriptional bursting.** Paper 4 discussion

* Discussion paper 5: Loop stacking model

Hafner, A. et al. Loop stacking organizes genome folding from TADs to chromosomes. Mol. Cell **83**, 1377-1392.e1376 (2023).

* Additional readings: Perspectives on condensate, loop, and hub model

Lim, B. & Levine, M. S. Enhancer-promoter communication: hubs or loops? *Curr. Opin. Genet. Dev.* **67**, 5-9 (2021).

Furlong, E. E. M. & Levine, M. Developmental enhancers and chromosome topology. *Science* **361**, 1341-1345 (2018).

### Week 4 – 2

**A Case of Study – Estrogen Receptor**

**Estrogen response pathway; Binding of estrogen receptor; Mechanism of estrogen-induced transcription; MegaTrans enhancers.** Hands-on session: Statistical Model of TF-DNA Interactions

* Additional readings: MegaTrans of estrogen-regulated enhancers

Liu, Z. *et al.* Enhancer Activation Requires Trans-Recruitment of a Mega Transcription Factor Complex. *Cell* **159**, 358-373 (2014).

* Additional readings: Phase separation at estrogen-induced enhancers

Nair, S. J. et al. Phase separation of ligand-activated enhancers licenses cooperative chromosomal enhancer assembly. *Nat. Struct. Mol. Biol.* **26**, 193-203 (2019).

### Week 5 – 1

**A Case of Study – Enhancer-Promoter Contact and 3D Organization**

**Global analysis of chromatin folding; Advanced technologies (Micro-C & MERFISH); Shaping of enhancer-promoter contact; Discoveries against canonical theory; Problems of heterogenicity.** Paper 5 discussion

* Additional readings: Reviews of the 3D genome structure

Zheng, H. & Xie, W. The role of 3D genome organization in development and cell differentiation. *Nat. Rev. Mol. Cell Biol.* **20**, 535-550 (2019).

Hafner, A. & Boettiger, A. The spatial organization of transcriptional control. *Nat. Rev. Genet.* **24**, 53-68 (2023).

van Steensel, B. & Furlong, E. E. M. The role of transcription in shaping the spatial organization of the genome. *Nat. Rev. Mol. Cell Biol.* **20**, 327-337 (2019).

### Week 5 – 2\*

**The 4D Genome and New Paradigm**

**Summary of the hierarchical 3D structure of chromatin; Reconstruction of the 3D genome; Reorganization of the genome during development; Structure-determined phenotype and diseases.** Research proposal discussion

* Additional readings: The 4D Genome

Aboelnour, E. & Bonev, B. Decoding the organization, dynamics, and function of the 4D genome. *Dev. Cell* **56**(11), 1562–1573 (2021).

Dekker, J. et al. The 4D nucleome project. *Nature.* **549**, 219–226 (2017).

### Week 6

**Final**

Research proposal presentation

**Q&A**