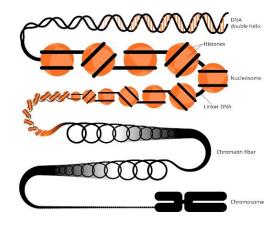
Michael Landi & Andreas Gisel

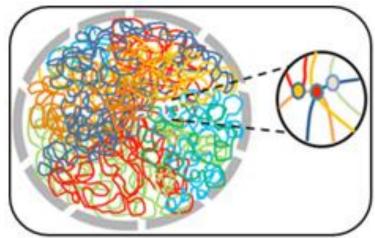
IITA- Bioinformatics Kenya & Nigeria



African Star Apple Assembly workshop 2 – 6 June 2025

Omni-C is a high-throughput genomic and epigenomic technique to capture chromatin conformation.

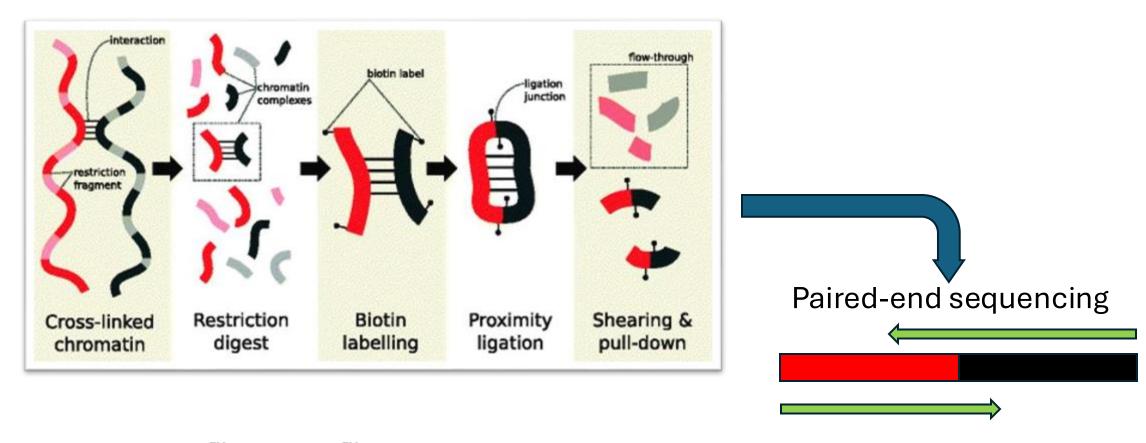




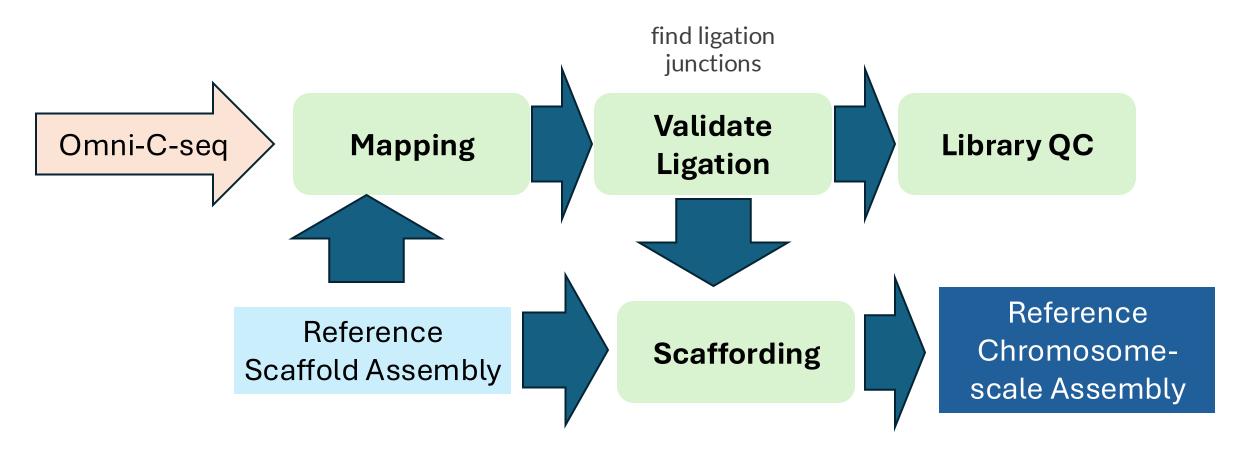
Hi-C comprehensively detects genome-wide chromatin interactions in the cell nucleus.

Hi-C measures the frequency (as an average over a cell population) at which two DNA fragments physically associate in 3D space, linking chromosomal structure directly to the genomic sequence.





The Dovetail™ Omni-C™ library uses a sequence-independent endonuclease (compared to Hi-C, using restriction enzymes) for chromatin digestion prior to proximity ligation and library generation.



**Mapping** 



Reference Contig assembly

For mapping, we need a specific reference index

We use bwa (<a href="https://bio-bwa.sourceforge.net/">https://bio-bwa.sourceforge.net/</a>)

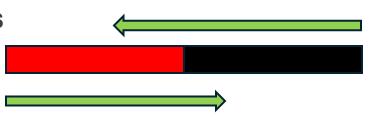
We need to clean the OmniC data

We need to run the mapper



Recording valid ligation events

- pairtools parse



find ligation junctions

Validate Ligation

### Sorting the pairsam file

- pairtools sort

### Removig PCR duplicates

- pairtools dedup

### Generating .pairs and bam files

- pairtools split

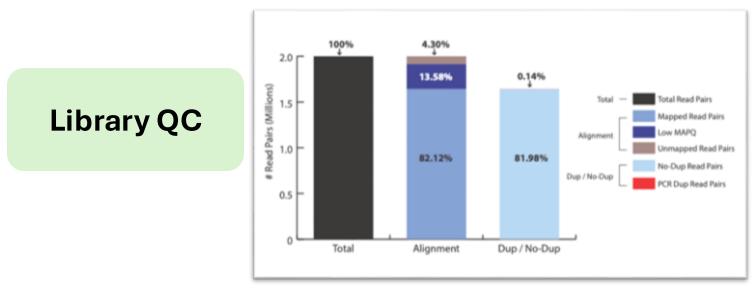


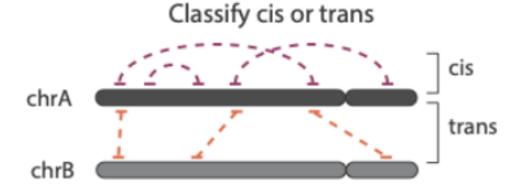
#### **Scaffording**

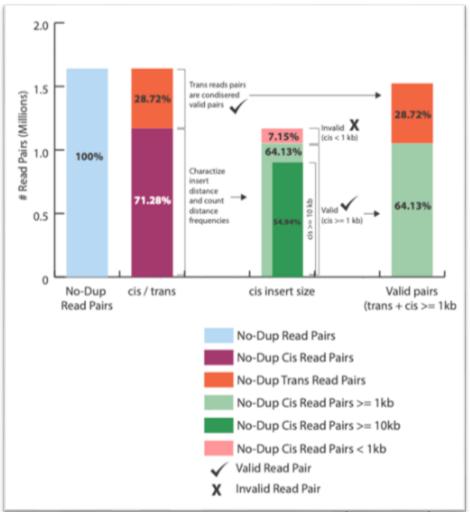
### Generating the final bam file

samtools sort











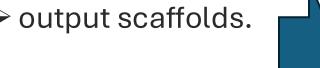
#### **Scaffording**

### YaHS: yet another Hi-C scaffolding tool

Zhou et al 2022, https://doi.org/10.1093/bioinformatics/btac808

### YaHS scaffolding pipelines:

- > map Hi-C reads to input contigs,
- > break contigs where necessary to correct assembly errors,
- build a contact matrix,
- > construct and prune a scaffolding graph
- > output scaffolds.



Reference Chromosomescale Assembly



