Cold Spring Harbor Laboratory

PMBIO Module 3:

DNA Alignment

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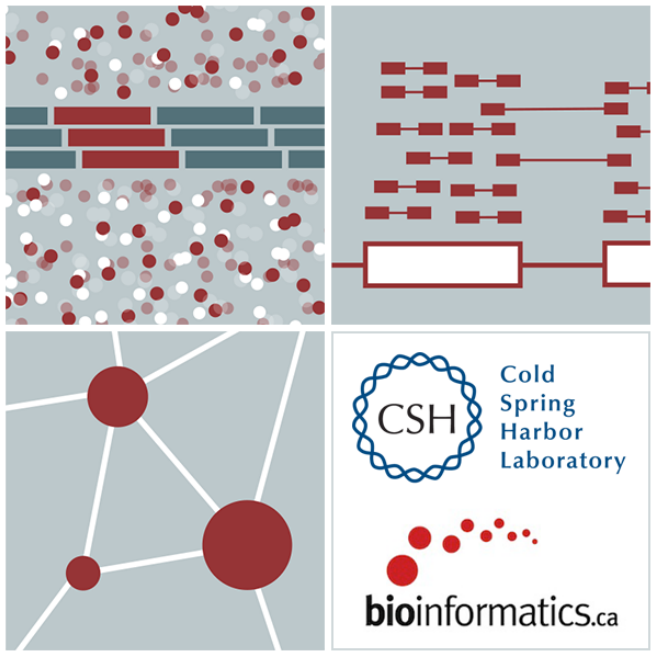
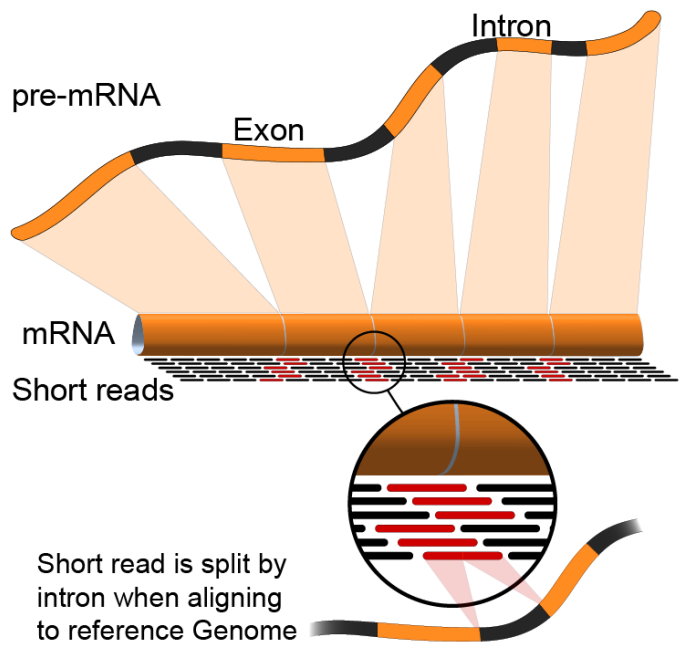




Washington University in St. Louis
School of Medicine

Learning Objectives

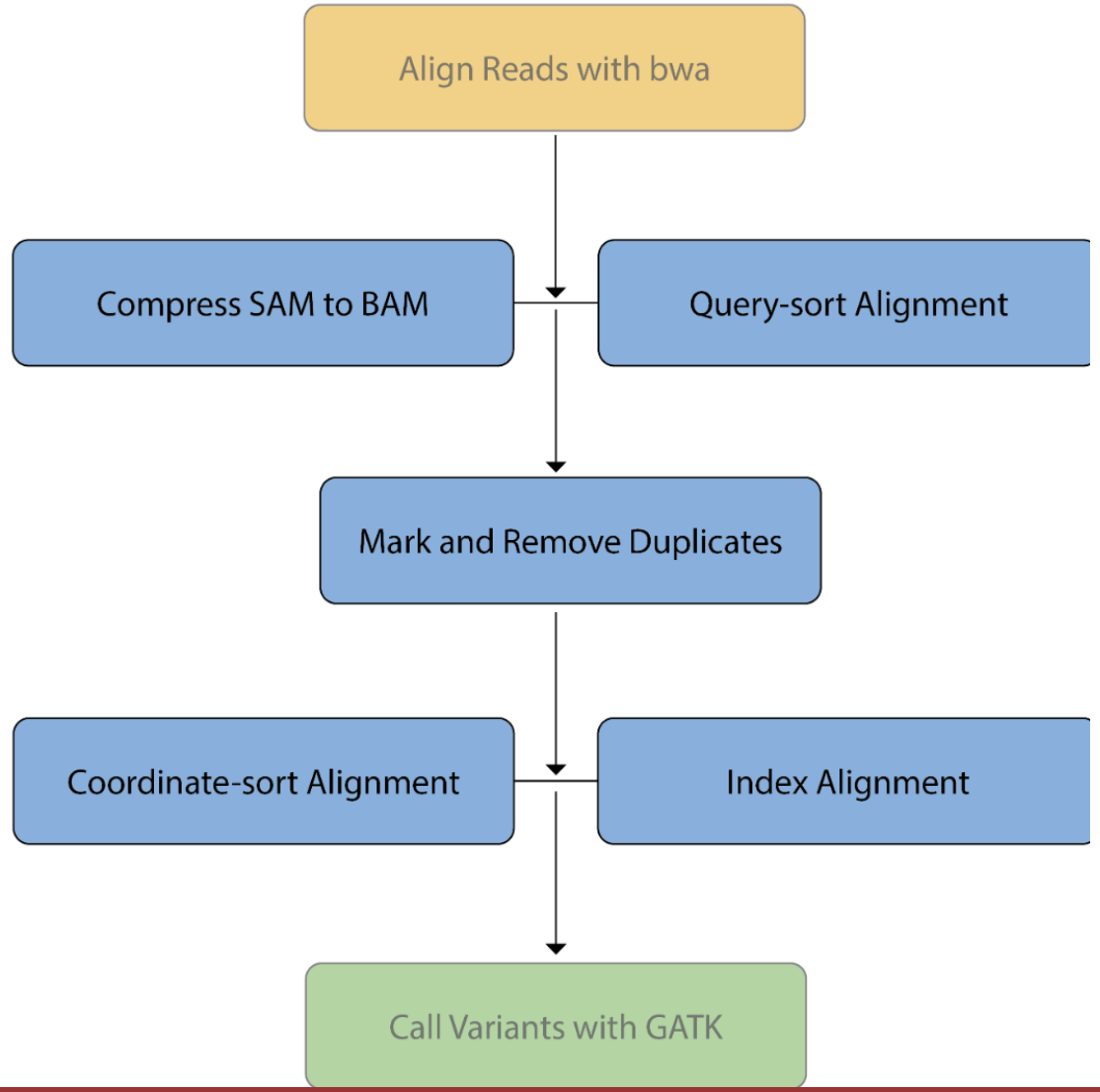
- Understand the process of obtaining and indexing raw DNA sequence data for alignment
- Perform a quality assessment using the aligned data and understand clean-up steps
- Prepare and visualize alignment results using genome viewers

Based on: https://pmbio.org/module-03-align/0003/02/01/Alignment/

For our exercise:

https://gist.github.com/MariamKhanfar/8eae80dfc5bd9011d2ccbab27458ca04

Main QC steps in DNA alignment



Overview of main steps

- Create directories
- Download the data
- Download the reference
- Index Reference with BWA
- Align the data to the reference → Result in SAM
- Convert SAM to BAM
- Sort by query name
- Mark duplicates
- Sort by coordinates
- Index the final BAM
- Generate a flagstat report for the BAM

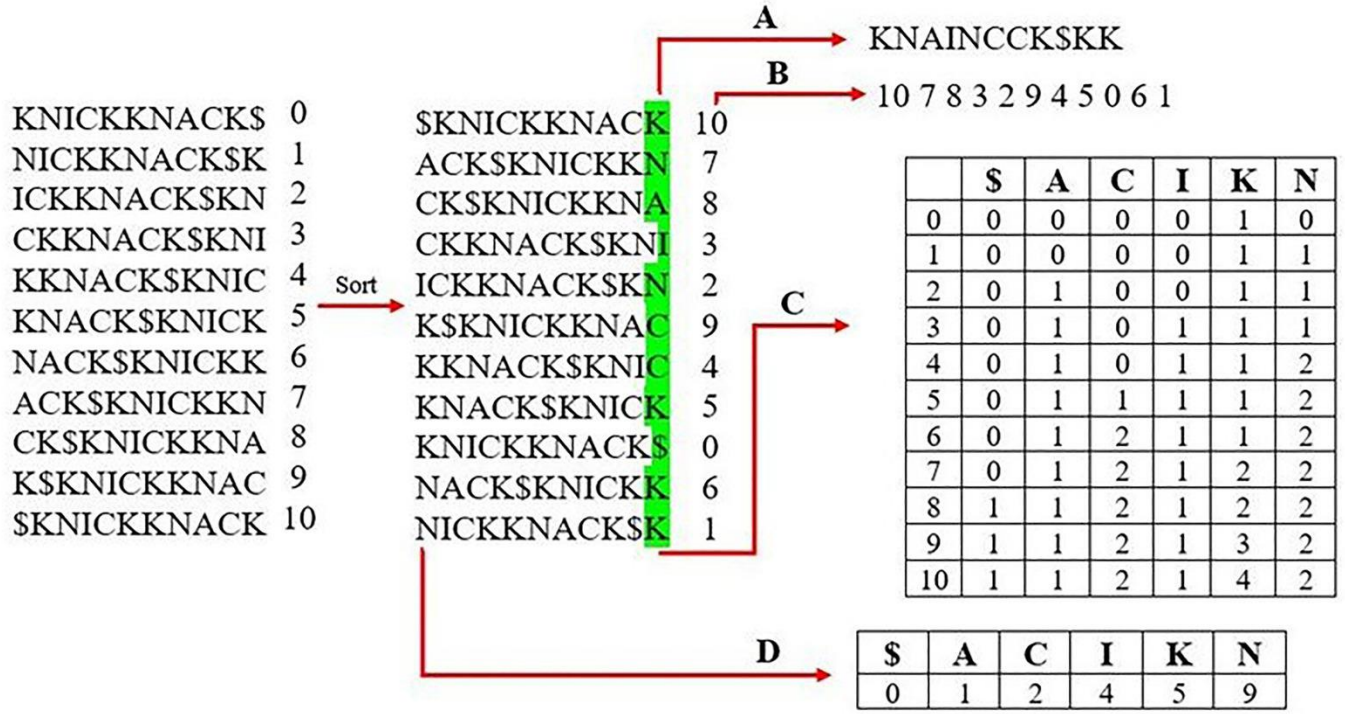
Prepare directories for analysis and obtain data

- Create the main directory "*dna_alignment_lab*"
- Create subdirectories within "*dna_alignment_lab*" for organizing data

- /workspace/dna_alignment_lab/alignment_results
- /workspace/dna_alignment_lab/fastq_files
- /workspace/dna_alignment_lab/reference_sequences

Index reference genome + DNA alignment

- Indexing is like creating a map of the reference genome, allowing the aligner to quickly locate where a sequence might fit; finds information without reading the entire text

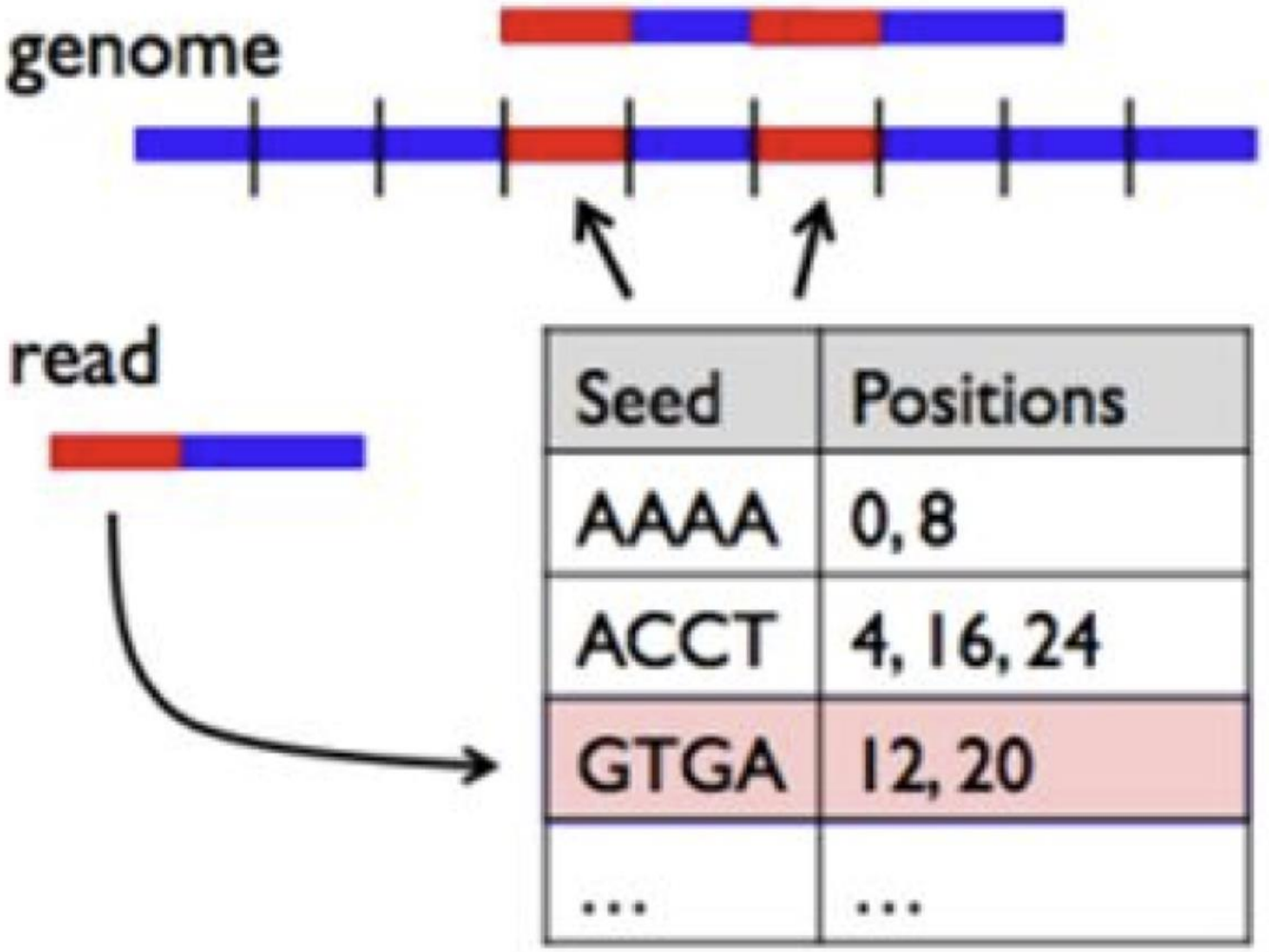


	$	A	C	I	K	N
0	0	0	0	0	1	0
1	0	0	0	0	1	1
2	0	1	0	0	1	1
3	0	1	0	1	1	1
4	0	1	0	1	1	2
5	0	1	1	1	1	2
6	0	1	2	1	1	2
7	0	1	2	1	2	2
8	1	1	2	1	2	2
9	1	1	2	1	3	2
10	1	1	2	1	4	2

$	A	C	I	K	N
0	1	2	4	5	9

https://www.frontiersin.org/articles/10.3389/fpls.2021.657240/full

DNA alignment with BWA

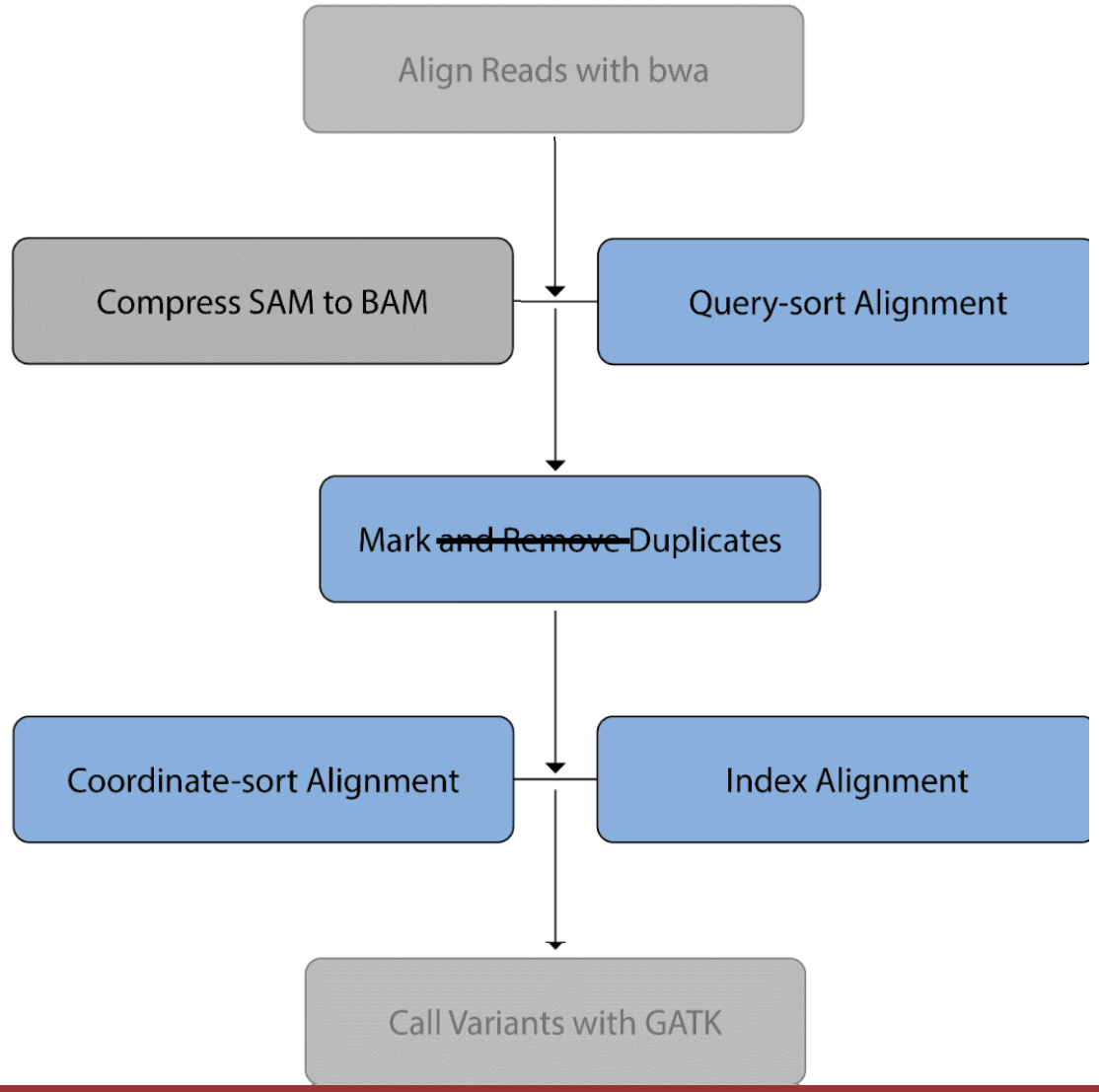


https://rbatorsky.github.io/intro-to-ngs-bioinformatics/lessons/03_Alignment.html

Convert SAM to BAM

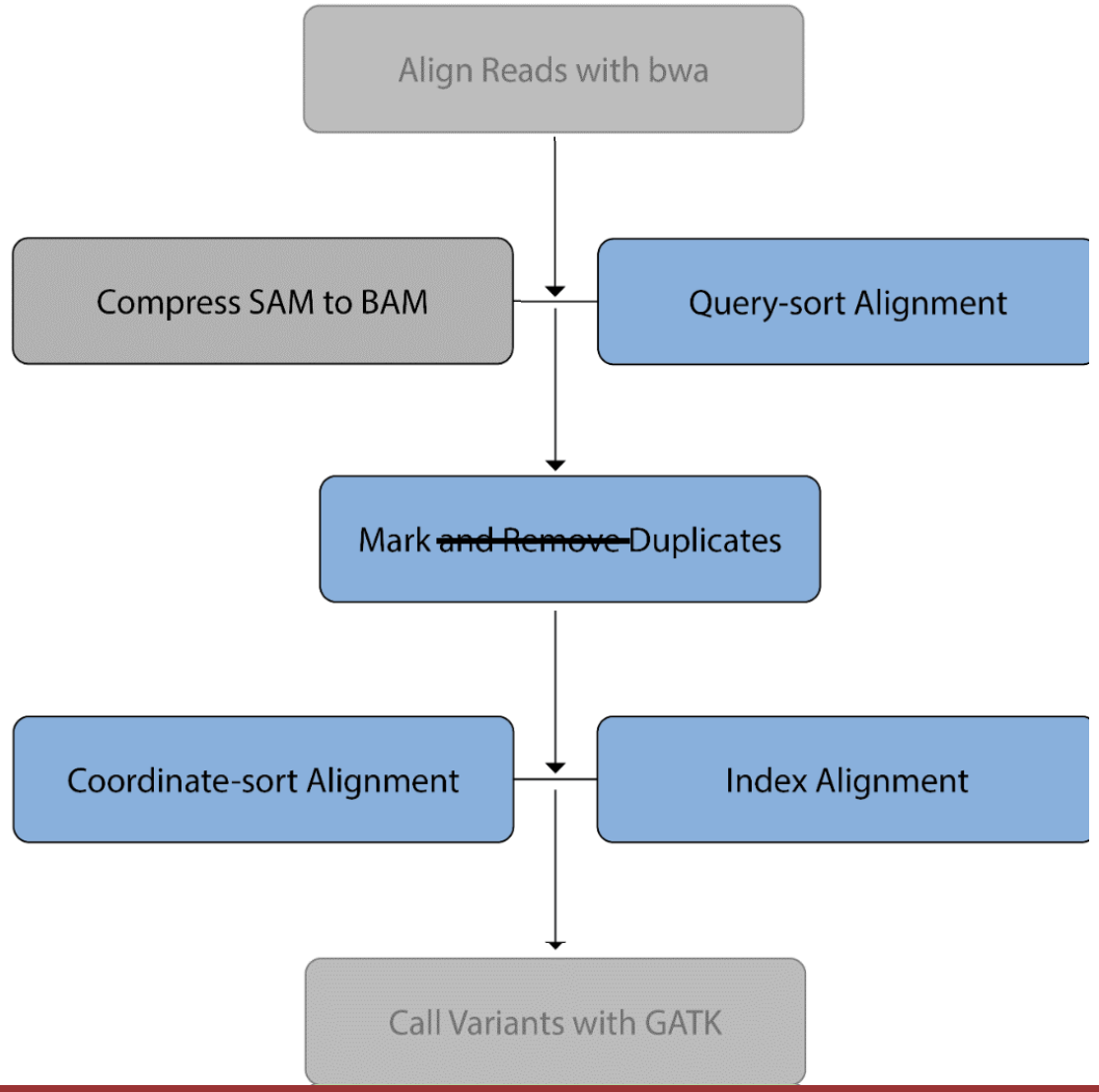
- SAM files are human-readable alignments, but they're large and unwieldy
- BAM files are the binary version of SAM, much smaller and faster for computers to process.

Main QC steps in DNA alignment



https://hbctraining.github.io/variant_analysis/lessons/06_alignment_file_processing.html

Main QC steps in DNA alignment



Can be done on:
SAM, BAM, CRAM

https://hbctraining.github.io/variant_analysis/lessons/06_alignment_file_processing.html

Sort by query name

- Grouping together reads that came from the same DNA fragment (i.e. identifier given to each read during sequencing)
- This is important for the following step; Marking duplicates, as it is essential to account for biases in sequencing where some fragments are overrepresented.

Query-sorted

QNAME	FLAG	RNAME	POS	...
A	[illegible]	chr21	342427	[illegible]
BA	[illegible]	chr4	4653	[illegible]
BBA	[illegible]	chr15	26171	[illegible]
C	[illegible]	chr1	2719101	[illegible]
D	[illegible]	chr15	1748549	[illegible]
E	[illegible]	chr4	2368992	[illegible]

https://hbctraining.github.io/variant_analysis/lessons/06_alignment_file_processing.html

Mark Duplicates

- To assure proper pair alignment, must be used on a name sorted BAM (i.e. mark secondary alignments)
- Locate and tag duplicate reads (both PCR and optical/sequencing-driven) , duplicate reads are defined as originating from the same original fragment of DNA.
- Produces a metrics file indicating the numbers of duplicates for both single- and paired-end reads.

Mark Duplicates

Mark Duplicates →

Sort by position

- Position sorting is performed to organize the reads by their location on the genome
- This is essential for downstream analysis, which require reads to be in order based on their *chromosomal coordinates.*
- Ensure that the subsequent analysis (like variant calling) do not mistakenly count extra copies of the same fragment, which could distort the results.
- Position sorting facilitates the identification of regions with genuine high coverage due to biological reasons rather than technical artifacts.

Sort by query name vs position

Query-sorted

QNAME	FLAG	RNAME	POS	...
A		chr21	342427	
BA		chr4	4653	
BBA		chr15	26171	
C		chr1	2719101	
D		chr15	1748549	
E		chr4	2368992	

Coordinate-sorted

QNAME	FLAG	RNAME	POS	...
C		chr1	2719101	
BA		chr4	4653	
E		chr4	2368992	
BBA		chr15	26171	
D		chr15	1748549	
A		chr21	342427	

https://hbctraining.github.io/variant_analysis/lessons/06_alignment_file_processing.html

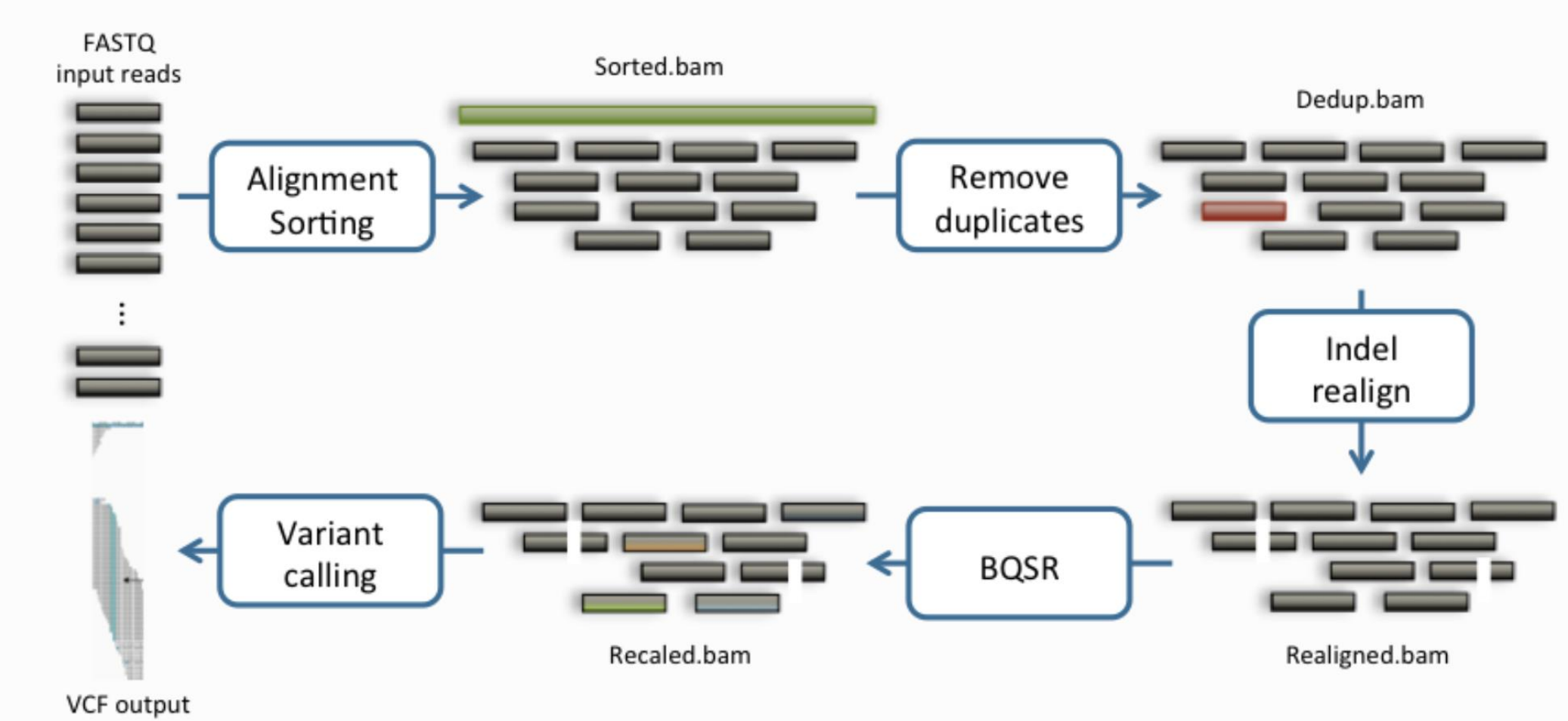
Index your final BAM

- In order to efficiently load and search a bam file, downstream applications typically require an index

SAM Flagstat

- Extract and count the number of reads that are: aligned, primary, not duplicate, and properly paired …etc, based on FLAG field.
  https://samtools.github.io/hts-specs/SAMv1.pdf
  https://broadinstitute.github.io/picard/explain-flags.html

Indel Realign or BQSR?



https://support.sentieon.com/versions/201808.01/manual/DNAseq_usage/dnaseq/