



Swiss Institute of  
Bioinformatics

INTRODUCTION TO SEQUENCING DATA ANALYSIS

# Read alignment

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Adapted from previous year courses

# Learning Objectives

Understand the concept and purpose of sequence alignment

Distinguish between global and local alignment strategies and their algorithms

Describe how short and long sequencing reads are aligned to reference genomes

Understand indexing strategies (e.g., BWT, suffix arrays) used in fast read alignment tools

In bioinformatics, **alignment** refers to the process of arranging sequences of DNA, RNA, or proteins to identify regions of similarity.

# Alignment types

## Pairwise alignment

- A. Global Alignment - *Needleman-Wunsch Algorithm*
  - DNA, RNA, or protein sequences of similar length
- B. Local Alignment - *Smith-Waterman Algorithm*
  - Protein or RNA/DNA domains

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- Aligns three or more sequences to detect conserved regions.

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## Multiple Sequence Alignment (MSA)

- Aligns three or more sequences to detect conserved regions.

## Read Alignment (e.g. BWA, Bowtie2, minimap2)

- Mapping short reads (DNA/RNA) to a reference genome

# Pairwise alignment: Global

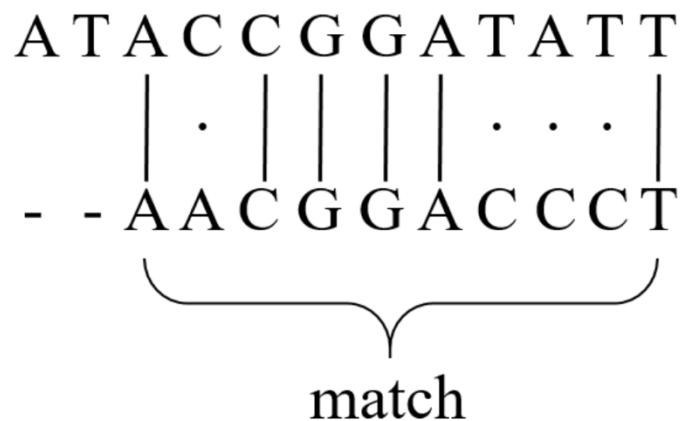
Sequence1 : A T A C C G G A T A T T

Sequence2 : A A C G G A C C C T

# Pairwise alignment: Global

Sequence1 : A T A C C G G A T A T T

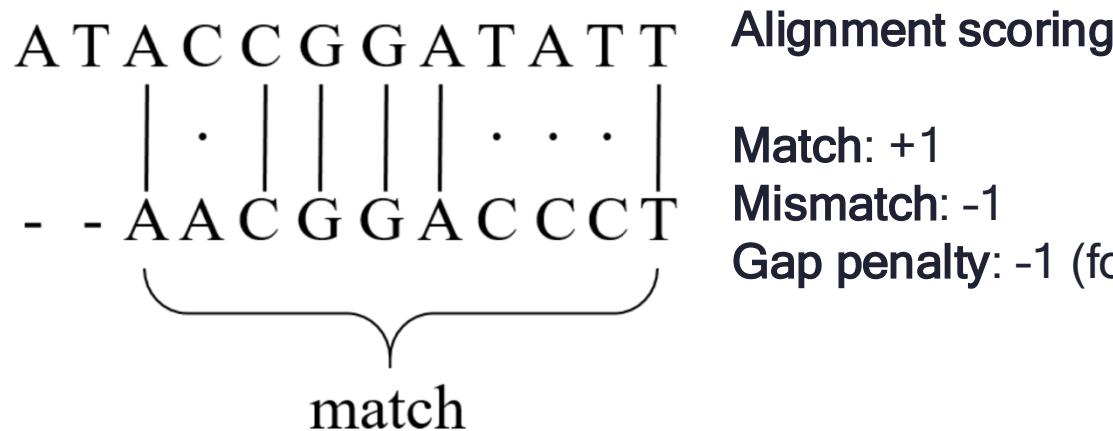
Sequence2 : A A C G G A C C C T



# Pairwise alignment: Global

Sequence1 : A T A C C G G A T A T T

Sequence2 : A A C G G A C C C T



**Match:** +1

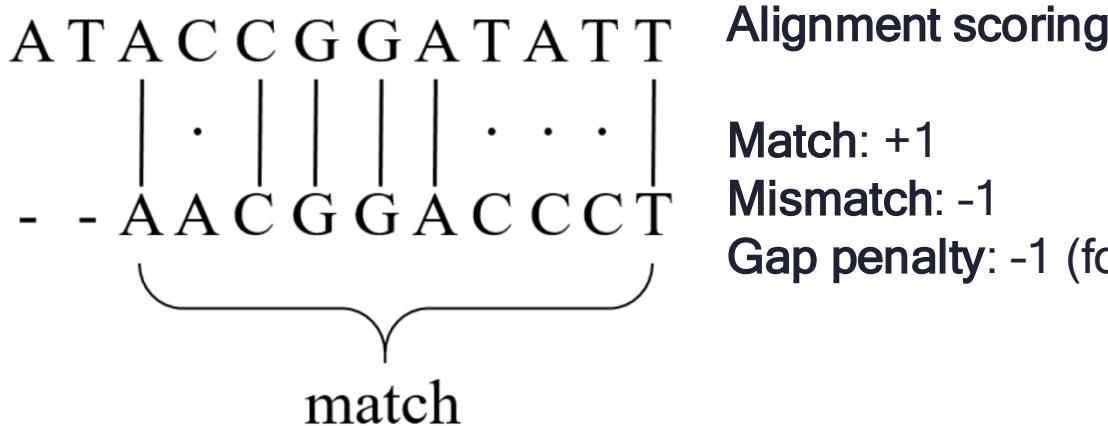
**Mismatch:** -1

**Gap penalty:** -1 (for each gap)

# Pairwise alignment: Global

Sequence1 : A T A C C G G A T A T T

Sequence2 : A A C G G A C C C T



Match: +1

Mismatch: -1

Gap penalty: -1 (for each gap)

S1	S2	Score
A	-	-1
T	-	-1
A	A	+1
C	A	-1
C	C	+1
G	G	+1
G	G	+1
A	A	+1
T	C	-1
A	C	-1
T	C	-1
T	T	+1
Total	o	

# Pairwise alignment: Local

Sequence1 : A T A C C G G A T A T T

Sequence2 : A A C G G A C C C T

## Alignment scoring

Match: +1

Mismatch: -1

Gap penalty: -1 (for each gap)

# Pairwise alignment: Local

Sequence1 : A T A C C G G A T A T T

Sequence2 : A A C G G A C C C T

A T A C C G G A T A T T   Alignment scoring



Match: +1

Mismatch: -1

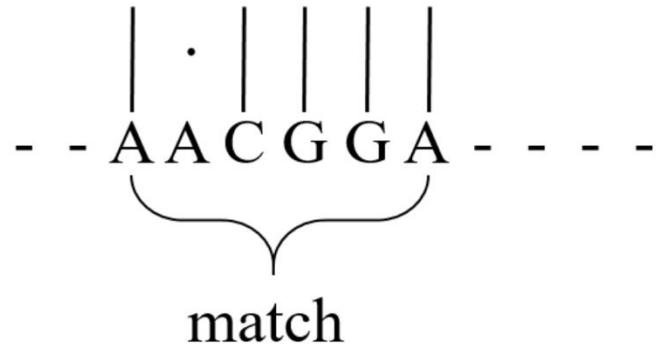
Gap penalty: -1 (for each gap)

# Pairwise alignment: Local

Sequence1 : A T A C C G G A T A T T

Sequence2 : A A C G G A C C C T

A T A C C G G A T A T T   Alignment scoring



Match: +1

Mismatch: -1

Gap penalty: -1 (for each gap)

S <sub>1</sub>	S <sub>2</sub>	Score
A	-	0
T	-	0
A	A	+1
C	A	-1
C	C	+1
G	G	+1
G	G	+1
A	A	+1
T	-	0
A	-	0
T	-	0
T	-	0
Total		4

# Quiz: 21-22

Which of the following algorithms is used for global sequence alignment?

...

In local alignment using the Smith-Waterman algorithm, which of the following best describes the approach?

# Multiple Sequence Alignment

For phylogeny, function prediction, or motif discovery.

Tool: Clustal Omega, MAFFT Command-line example with Clustal Omega

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Tool: Clustal Omega, MAFFT Command-line example with Clustal Omega

A	T	C	G
A	T	G	G
A	C	G	

# Multiple Sequence Alignment

For phylogeny, function prediction, or motif discovery.

Tool: Clustal Omega, MAFFT Command-line example with Clustal Omega

A	T	C	G
A	T	G	G
A	C	G	

S <sub>1</sub>	S <sub>2</sub>	Score
A	A	+1
A	A	+1
A	A	+1
Total		3

# Multiple Sequence Alignment

For phylogeny, function prediction, or motif discovery.

Tool: Clustal Omega, MAFFT Command-line example with Clustal Omega

A	T	C	G
A	T	G	G
A	C	G	

S <sub>1</sub>	S <sub>2</sub>	Score
A	A	+1
A	A	+1
A	A	+1
Total	3	

S <sub>1</sub>	S <sub>2</sub>	Score
T	T	+1
T	C	-1
T	C	-1
Total	-1	

# Multiple Sequence Alignment

For phylogeny, function prediction, or motif discovery.

Tool: Clustal Omega, MAFFT Command-line example with Clustal Omega

The diagram illustrates three separate local alignments (S1, S2) derived from a common reference sequence (top). Each alignment consists of a 4x4 grid of amino acids and a corresponding table below showing scores and totals.

**Reference Sequence:**

A	T	C	G
A	T	G	G
A	C	G	

**Path 1 (S1):** A → A → A → A

S <sub>1</sub>	S <sub>2</sub>	Score
A	A	+1
A	A	+1
A	A	+1
Total	3	

**Path 2 (S2):** A → T → C → T

S <sub>1</sub>	S <sub>2</sub>	Score
T	T	+1
T	C	-1
T	C	-1
Total	-1	

**Path 3 (S3):** A → C → G → G

S <sub>1</sub>	S <sub>2</sub>	Score
C	G	-1
C	G	-1
G	G	+1
Total	-1	

# Multiple Sequence Alignment

For phylogeny, function prediction, or motif discovery.

Tool: Clustal Omega, MAFFT Command-line example with Clustal Omega

S <sub>1</sub>	S <sub>2</sub>	Score
A	A	+1
A	A	+1
A	A	+1
Total	3	

S <sub>1</sub>	S <sub>2</sub>	Score
T	T	+1
T	C	-1
T	C	-1
Total	-1	

S <sub>1</sub>	S <sub>2</sub>	Score
C	G	-1
C	G	-1
G	G	+1
Total	-1	

S <sub>1</sub>	S <sub>2</sub>	Score
G	G	+1
G	-	-1
G	-	-1
Total	-1	

# Multiple Sequence Alignment

For phylogeny, function prediction, or motif discovery.

Tool: Clustal Omega, MAFFT Command-line example with Clustal Omega

The diagram illustrates four different sequence alignment paths from a reference sequence (top) to four different S1-S2 pairs (bottom). Each path is represented by a green arrow pointing from the reference sequence to its respective score table.

**Reference Sequence:**

A	T	C	G
A	T	G	G
A	C	G	

**Path 1:** A → A → A → Total Score = 3

S <sub>1</sub>	S <sub>2</sub>	Score
A	A	+1
A	A	+1
A	A	+1
Total	3	

**Path 2:** A → T → C → Total Score = -1

S <sub>1</sub>	S <sub>2</sub>	Score
T	T	+1
T	C	-1
T	C	-1
Total	-1	

**Path 3:** A → C → G → Total Score = -1

S <sub>1</sub>	S <sub>2</sub>	Score
C	G	-1
C	G	-1
G	G	+1
Total	-1	

**Path 4:** A → G → - → Total Score = -1

S <sub>1</sub>	S <sub>2</sub>	Score
G	G	+1
G	-	-1
G	-	-1
Total	-1	

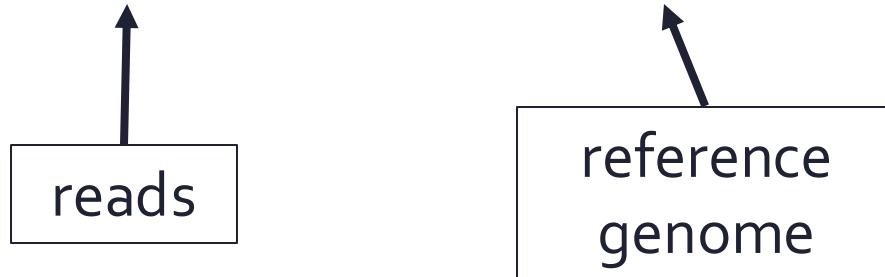
$$\text{MSA score: } 3 - 1 - 1 - 1 = 0$$

# Read Alignment

Mapping millions of short reads to a reference genome.

Read Aligners: **BWA**, **Bowtie2**, **STAR**, etc.

Aim: find **substrings** in **large string**



# Indexing

Aim: generate a ‘phonebook’ for fast searches

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Reference: TAATA\$



EOF

# Indexing

Aim: generate a ‘phonebook’ for fast searches

Reference: TAATA\$

↑  
EOF

0	T	A	A	T	A	\$
1	A	A	T	A	\$	
2	A	T	A	\$		
3	T	A	\$			
4	A	\$				
5	\$					

# Indexing

Aim: generate a ‘phonebook’ for fast searches

Reference: TAATA\$

↑  
EOF

suffix array

0	T	A	A	T	A	\$
1	A	A	T	A	\$	
2	A	T	A	\$		
3	T	A	\$			
4	A	\$				
5	\$					

sort

5	\$					
4	A	\$				
1	A	A	T	A	\$	
2	A	T	A	\$		
3	T	A	\$			
0	T	A	A	T	A	\$

# Querying

Reference: TAATA\$

Query: ATA

5	\$
4	A \$
1	A A T A \$
2	A T A \$
3	T A \$
0	T A A T A \$

# Indexing and querying

Suffix array: large, same sequence stored multiple times

BWT: only first and **last** columns are stored -> still enables fast querying

suffix array

5	\$
4	A \$
1	A A T A \$
2	A T A \$
3	T A \$
0	T A A T A \$

# Indexing and querying

Suffix array: large, same sequence stored multiple times

BWT: only first and **last** columns are stored -> still enables fast querying

suffix array

5	\$					
4	A	\$				
1	A	A	T	A	\$	
2	A	T	A	\$		
3	T	A	\$			
0	T	A	A	T	A	\$

Burrows-Wheeler Transformation

\$	T	A	A	T	A
A	\$	T	A	A	T
A	A	T	A	\$	T
A	T	A	\$	T	A
T	A	\$	T	A	A
T	A	A	T	A	\$

<https://www.youtube.com/watch?v=4WRANhDiSHM&t=1s>

# Global vs local

## Global (end-to-end)

Read:        GACTGGGCGATCTCGACTTCG  
              |||||    ||||||||| |||  
Reference: GACTG--CGATCTCGACATCG

## Local (allows for ‘clipping’)

Read:        ACGGTTGCGTTAA-TCCGCCACG  
              ||||||| |||||  
Reference: TAACTTGCCTAAATCCGCCTGG

examples from: <http://bowtie-bio.sourceforge.net/bowtie2/manual.shtml#end-to-end-alignment-versus-local-alignment>

# Quiz: 23

What is the primary role of the Burrows-Wheeler Transform (BWT) in read alignment?

# Software

Basic alignment:

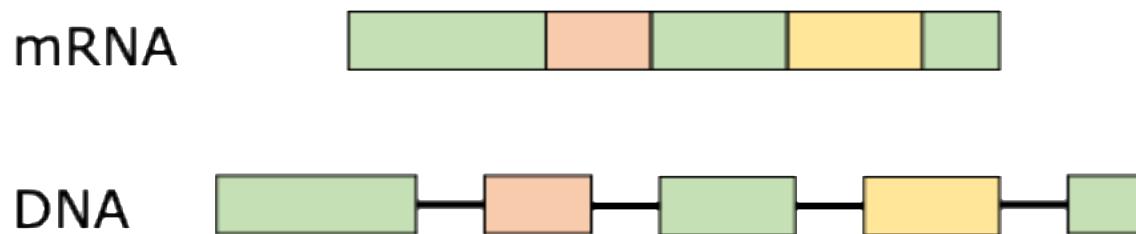
- » bowtie2 (BWT; default = global)
- » bwa-mem (BWT; default = local )

Splice-aware (RNA-seq):

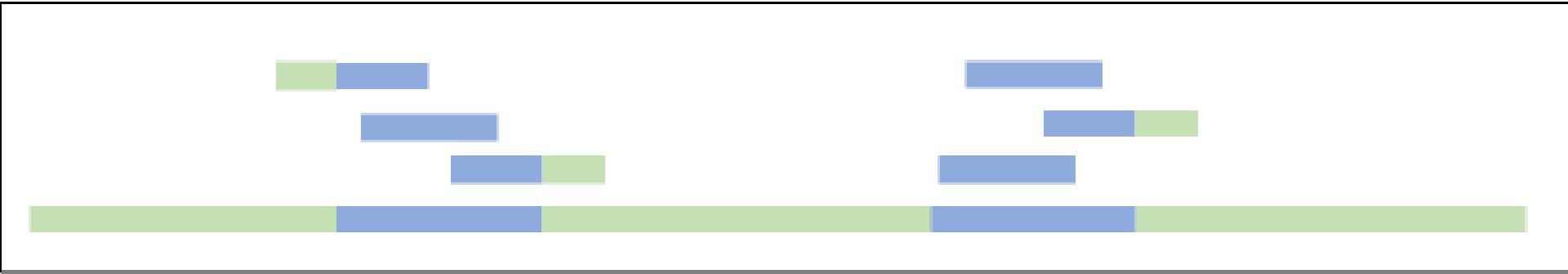
- » hisat2
- » STAR

Long reads + short reads + splice-aware:

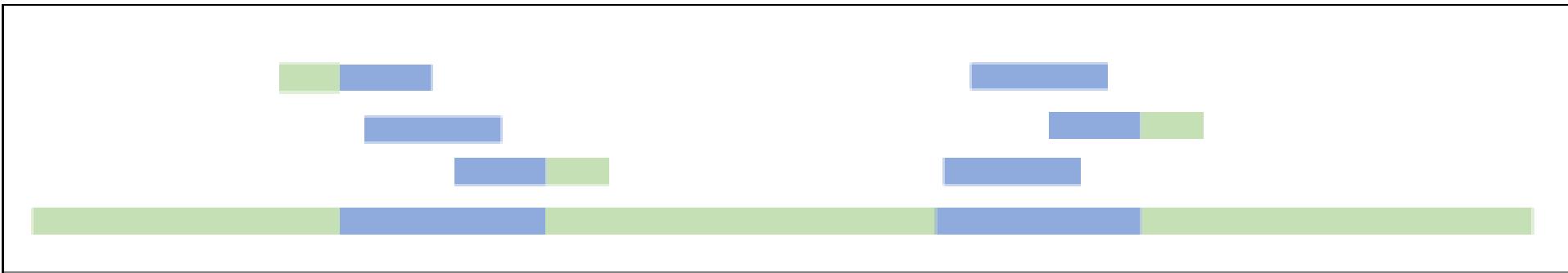
- » minimap2



# Mapping quality



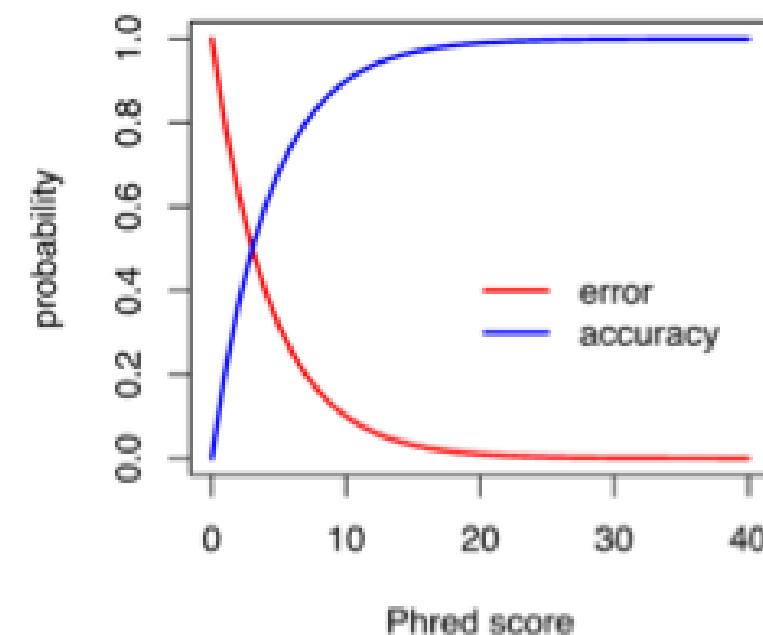
# Mapping quality



$MAPQ$   
 $= -10\log_{10} \Pr\{\text{mapping position is wrong}\}$

$$-10\log_{10} (0.01) = 20$$

$$-10\log_{10} (0.5) = 3$$



# Quiz: 24-25

Which tool is best suited for aligning RNA-seq reads with splice awareness?

...

What distinguishes global from local alignment in the context of Bowtie2?

# Summary

**Global alignment** aligns full sequences end-to-end

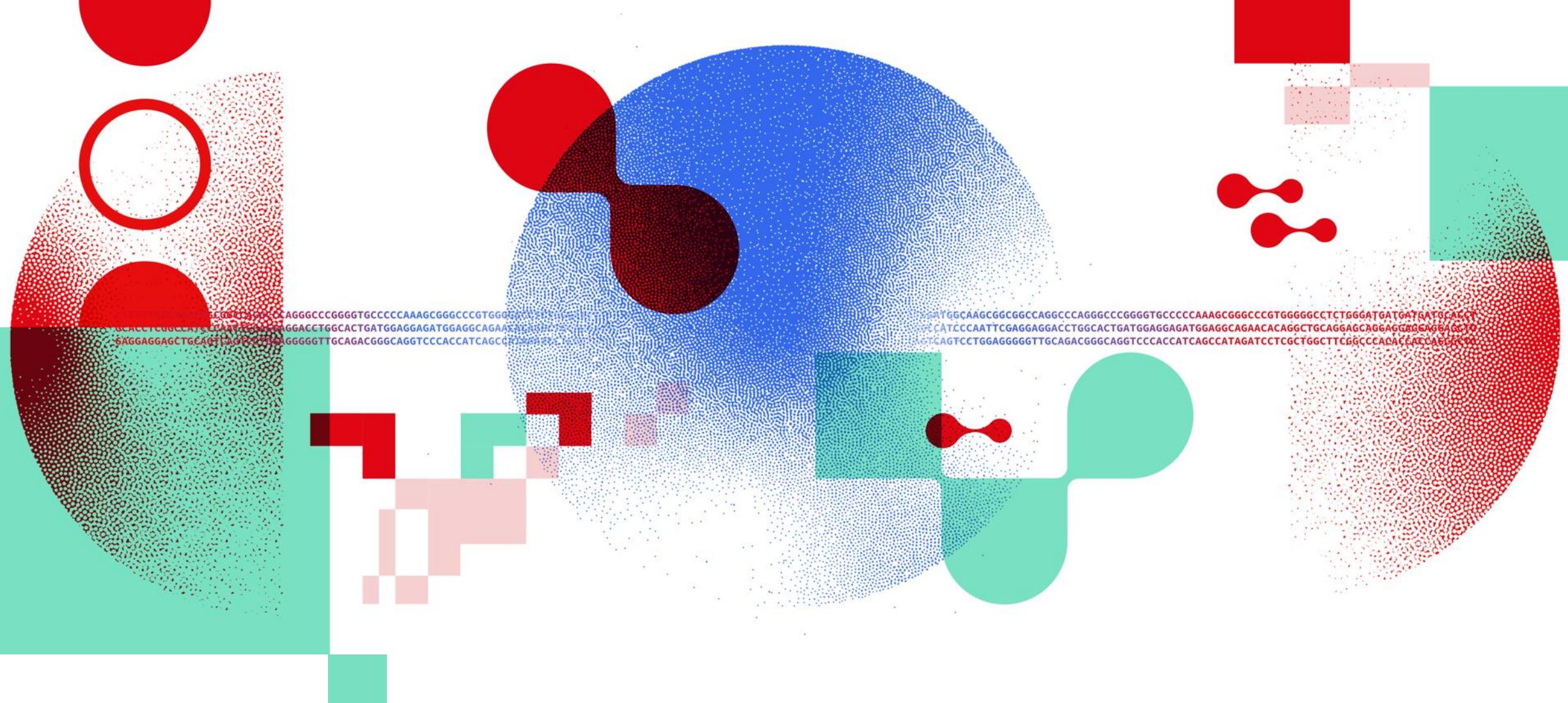
**Local alignment** is suited for partial matches like conserved domains

**MSA** aligns multiple sequences to reveal evolutionary or functional patterns

**Read alignment** maps millions of sequencing reads to a reference genome

**Suffix arrays and BWT** are used to enable fast searching within large genome indexes

Tools like **BWA**, **Bowtie2**, **STAR**, and **minimap2** support various alignment types, with specific strengths for short reads, long reads, or spliced transcripts



# Thank you

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