Building the compacted colored de Bruijn Graph

UNIVERSITY OF MARYLAND

Construction of the compacted colored De Bruijn Graph from *reference* sequence

Bioinformatics, 33(24), 2017, 4024–4032
doi: 10.1093/bioinformatics/btw609
Advance Access Publication Date: 21 September 2016
Original Paper

Sequence analysis

TwoPaCo: an efficient algorithm to build the compacted de Bruijn graph from many complete genomes

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Associate Editor: Alfonso Valencia

Received on April 3, 2016; revised on September 1, 2016; accepted on September 16, 2016

TwoPaCo: An efficient algorithm to build the compacted de Bruijn graph from many complete genomes

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8th July 2016

TwoPaCo slides, unless otherwise noted, are from Ilia's presentation

Motivation

- More and more complete genomes
- Pan-genome: analysis within same species
- Mammalian-sized genomes are coming soon

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Key question: what is a handy data structure to represent genomes?

The simplest way: string(s) of characters.

The Linear Representation

Two genomes:

The Linear Representation

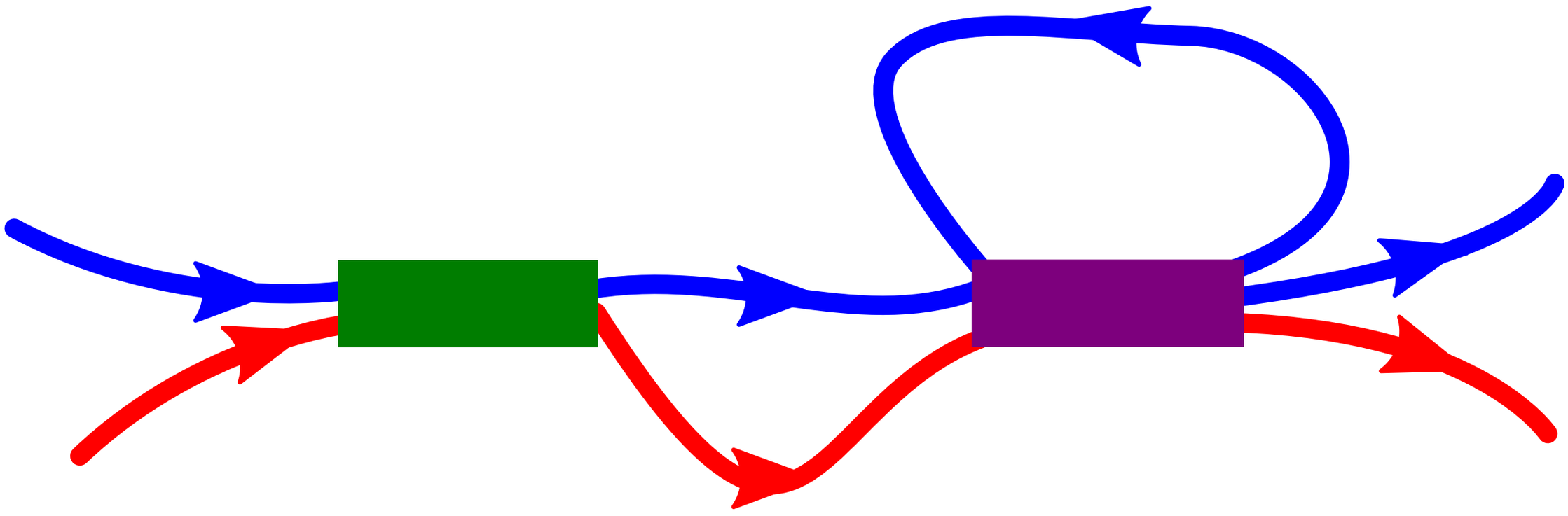
Two genomes:

Issues:

- Homology between genomes?
- Duplications?
- Rearrangements?

Solution: a Graph Representation

What we want to see:



Why de Bruijn graph?

A simple object.

Demonstrated utility in:

- Assembly
- Read mapping
- Synteny identification

The de Bruijn Graph

$k = 2$

TGACGTC

TGACTTC

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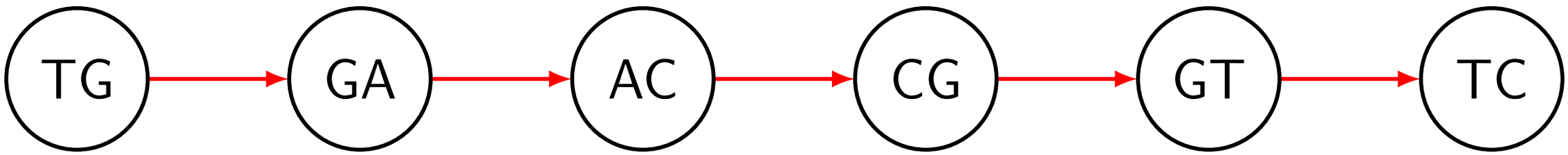

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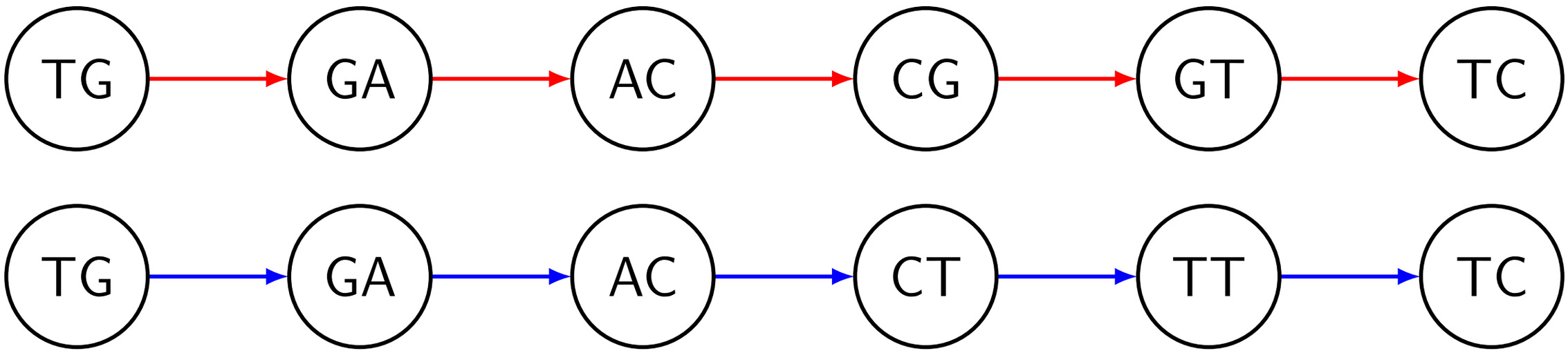

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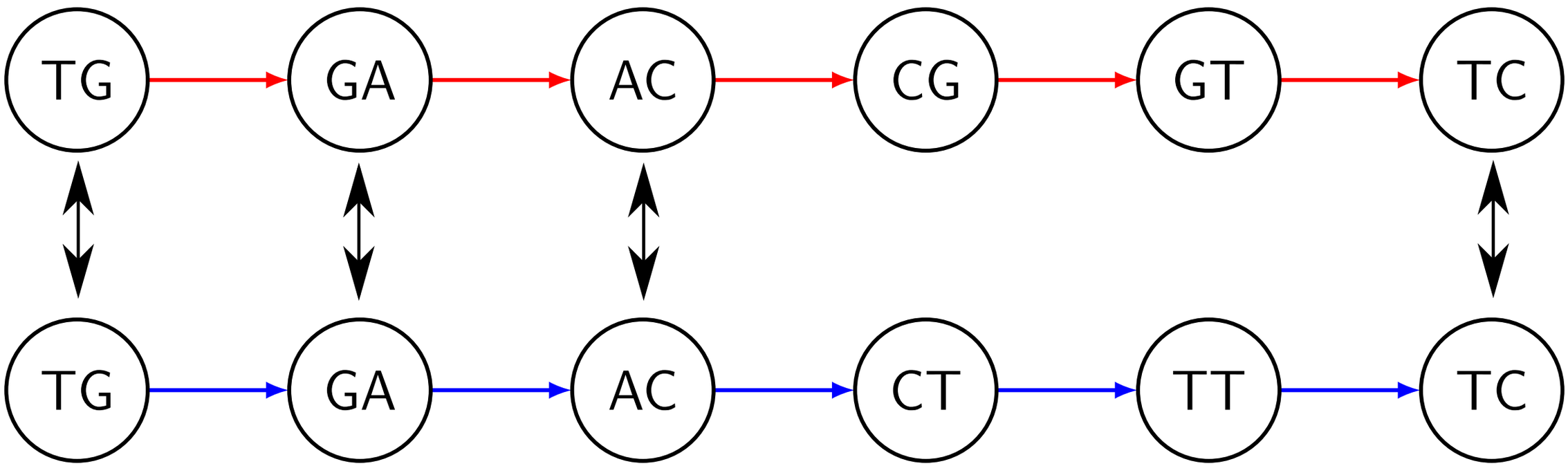
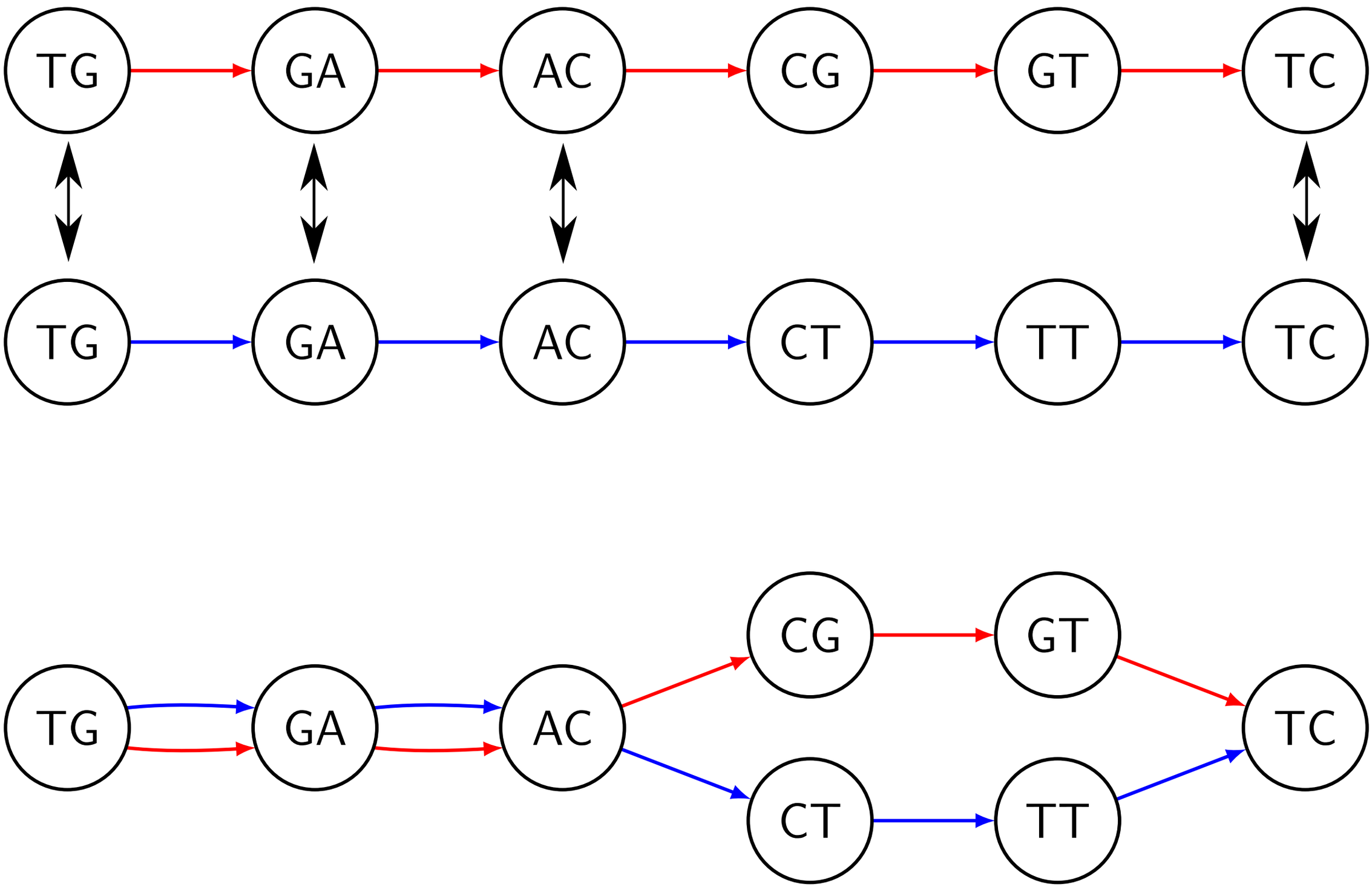
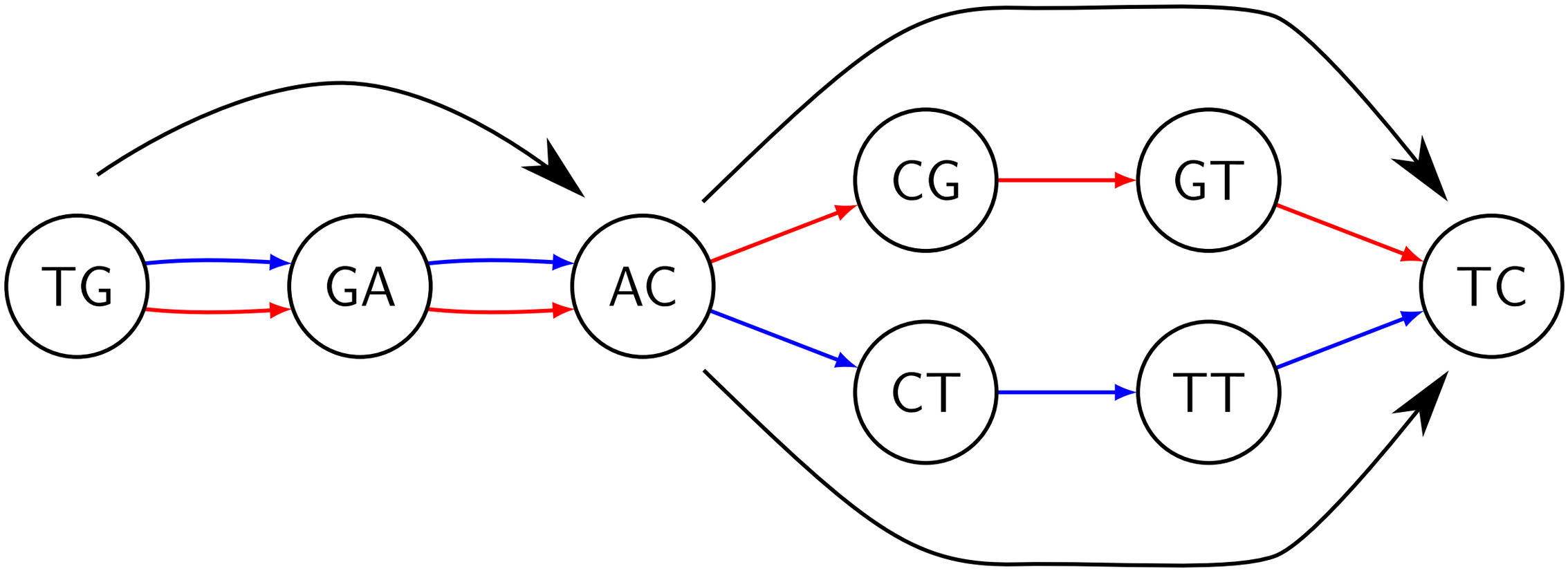


The de Bruijn Graph



Compaction



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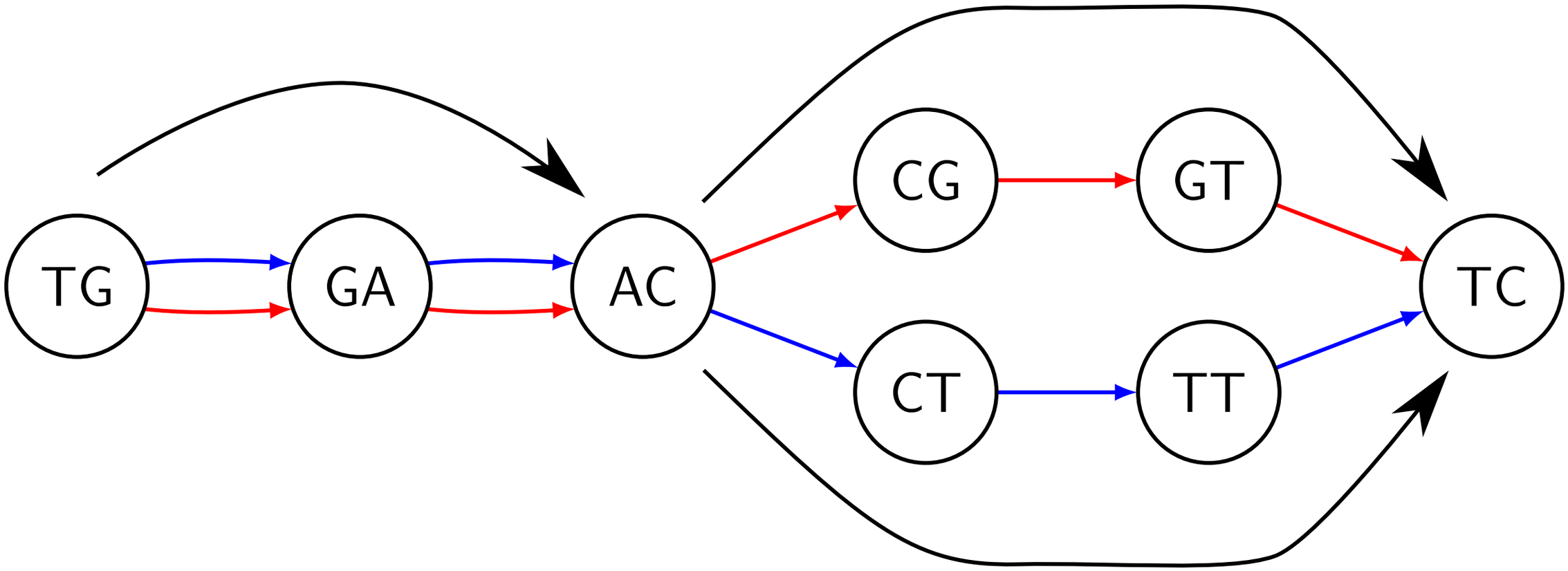
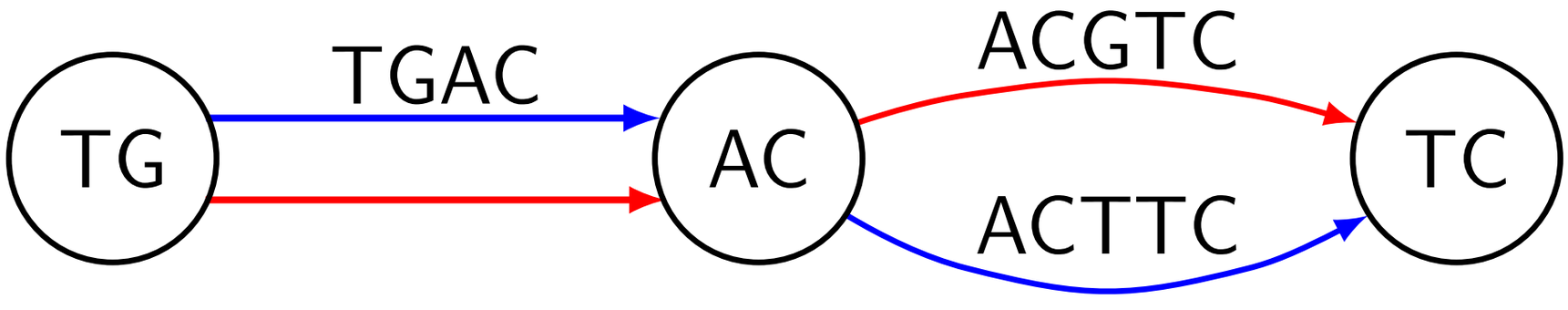


After compaction:



The Challenge

Construct the compacted graph from many large genomes **bypassing** the ordinary graph traverse.

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Earlier work: based on suffix arrays/trees Sibelia & SplitMEM handled > 60 E.Coli genomes.

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Earlier work: based on suffix arrays/trees Sibelia & SplitMEM handled $>$ 60 E.Coli genomes.

A recent advance: 7 Humans in 15 hours using 100 GB of RAM using a BWT-based algorithm by Baier *et al.*, 2015, Beller *et al.*, 2014.

Junctions

A vertex v is a **junction** if:

- v has ≥ 2 distinct outgoing or incoming edges:

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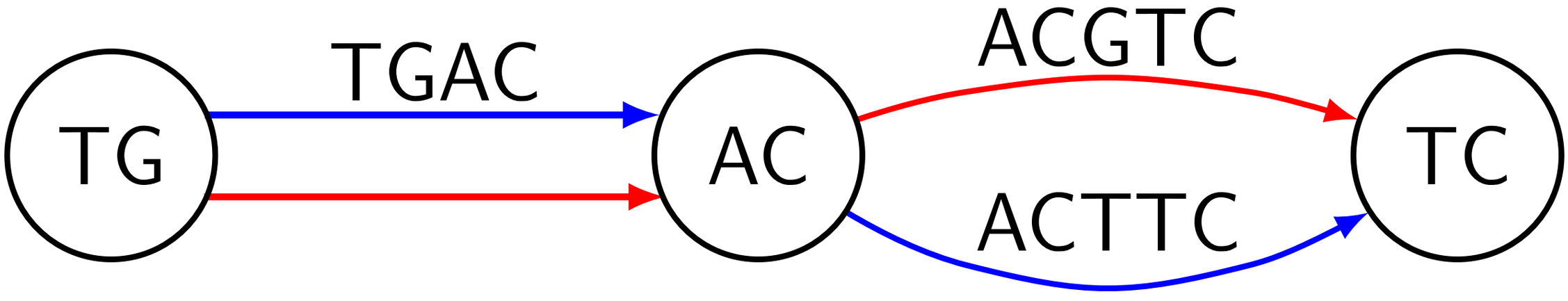
Facts:

- Junctions $=$ vertices of the compacted graph
- Compaction $=$ finding positions of junctions

Observations



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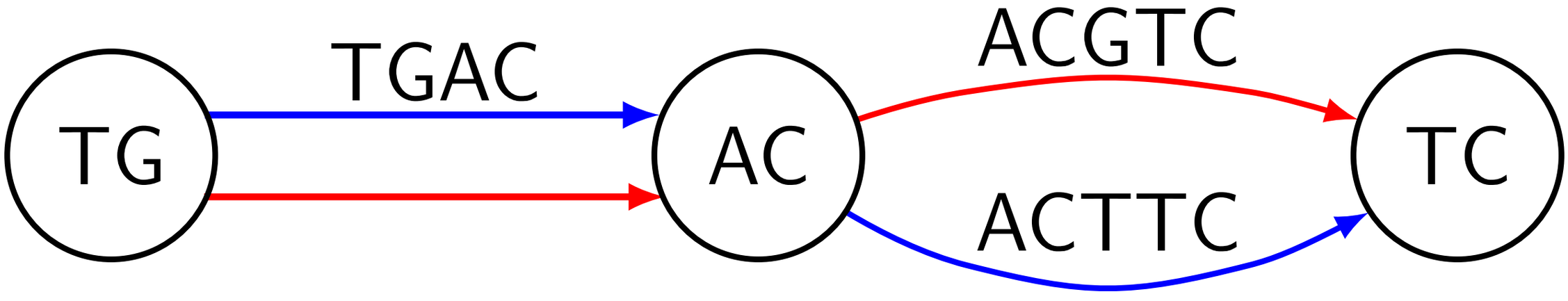
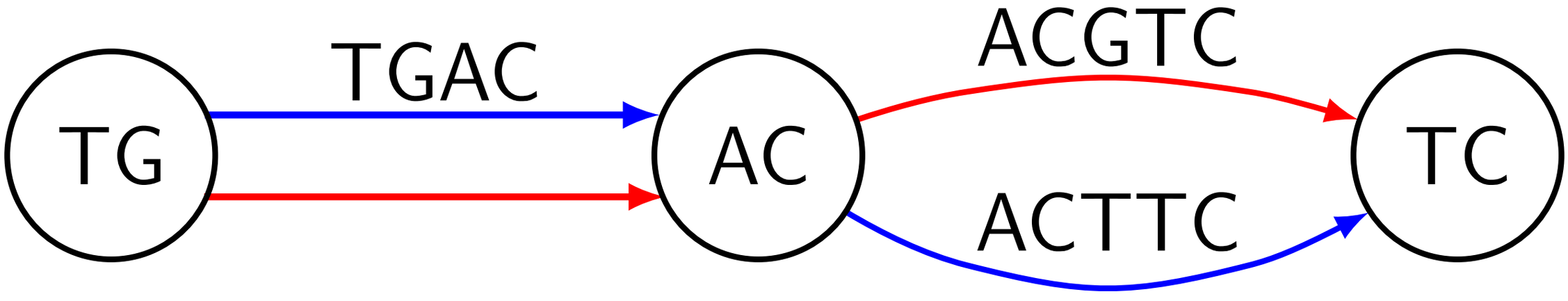


TG GA AC CG GT TC

Observations



TG GA AC CG GT TC

TG $\rightarrow$ AC $\rightarrow$ TC

The Observation

The observation only works when we have complete genomes.

Once we know junctions, construction of the edges is simple.

We can simply traverse input strings and record junctions in the order they appear.

How to identify junctions?

The Naive Algorithm

A naive way:

- Store all $(k+1)$-mers (edges) in a hash table
- Consider each vertex one by one
- Query all possible edges from the table
- If found > 1 edge, mark vertex as a junction

Simple algorithm in more detail

Algorithm 1. *Filter-Junctions*

Input: strings $S = \{s_1, \ldots, s_n\}$, integer k, and an empty set data structure E. A candidate set of marked junction positions $C \supseteq J(S,k)$ is also given. When the algorithm is run naively, all the positions would be marked.
Output: a reduced candidate set of junction positions.

```
1: for s ∈ S do
2:     for 1 ≤ i < |s| − k do
3:         if C[s, i] = marked then                ▷ Insert the two (k+1)-mers containing the k-mer at i into E.
4:             Insert s[i..i + k] into E.
5:             Insert s[i − 1..i − 1 + k] into E.
6: for s ∈ S do
7:     for 1 ≤ i < |s| − k do
8:         if C[s, i] = marked and s[i..i + k − 1] is not a sentinel then
9:             in ← 0                              ▷ Number of entering edges
10:            out ← 0                             ▷ Number of leaving edges
11:            for c ∈ {A, C, G, T} do             ▷ Consider possible edges and count how many of them exist
12:                if v · c ∈ E then               ▷ The symbol · depicts string concatenation
13:                    out ← out + 1
14:                if c · v ∈ E then
15:                    in ← in + 1
16:            if in = 1 and out = 1 then          ▷ If the k-mer at i is not a junction.
17:                C[s, i] ← Unmarked
18: return C
```

The Naive Algorithm

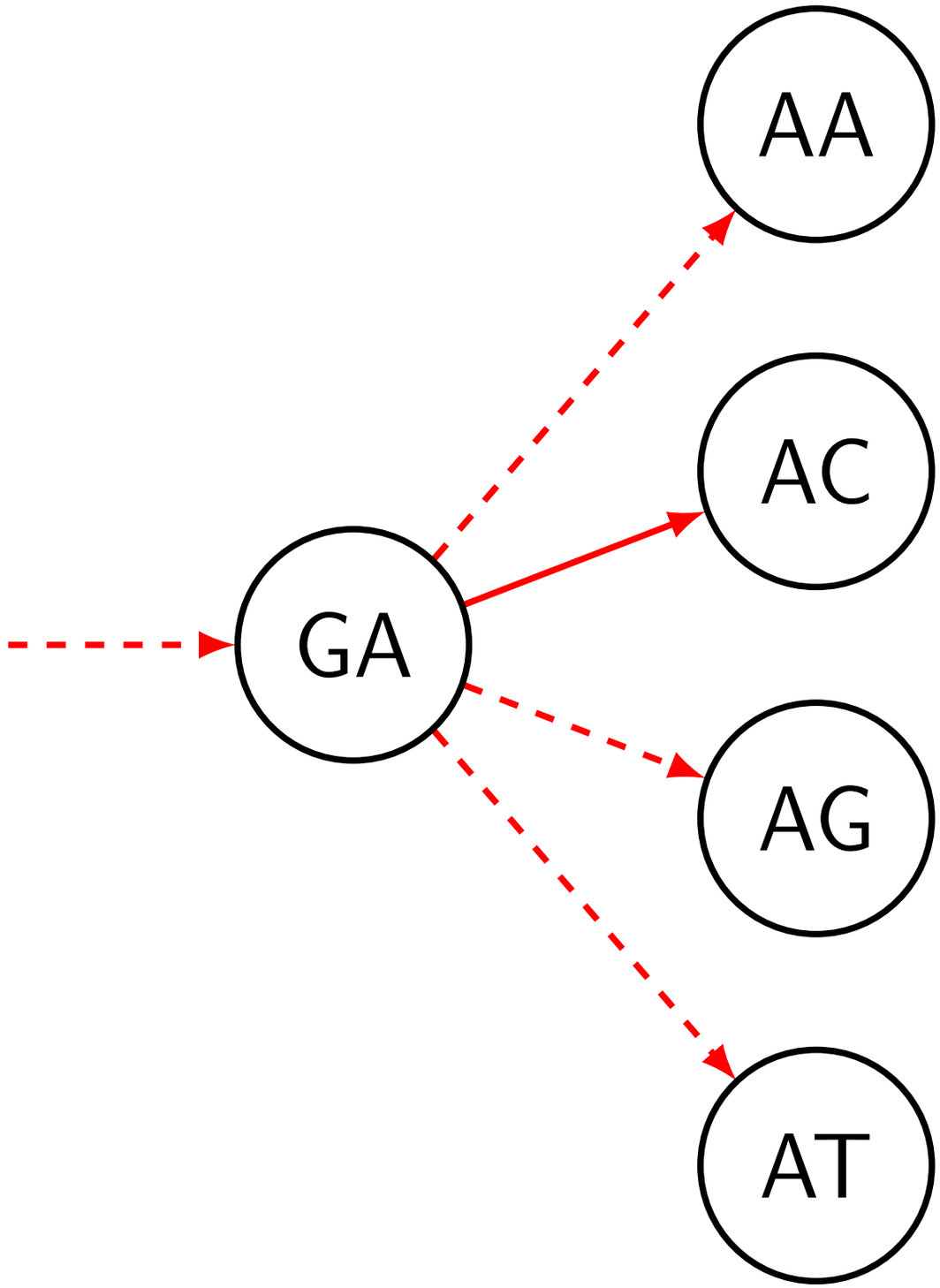
A naive way:

- Store all $(k+1)$-mers (edges) in a hash table
- Consider each vertex one by one
- Query all possible edges from the table
- If found > 1 edge, mark vertex as a junction

Problem: the hash table can be too large.

An Example

Hash table $= \{$ GA $\rightarrow$ AC $\}$



What is the Bloom filter

A probabilistic data structure representing a set

Properties:

- Occupies fixed space
- May generate false positives on queries
- False positive rate is low

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Example: Bloom Filter $= \{$ GA $\rightarrow$ AC $\}$

Is GA $\rightarrow$ AC in the set? Yes.

What is the Bloom filter

A probabilistic data structure representing a set

Properties:

- Occupies fixed space
- May generate false positives on queries
- False positive rate is low

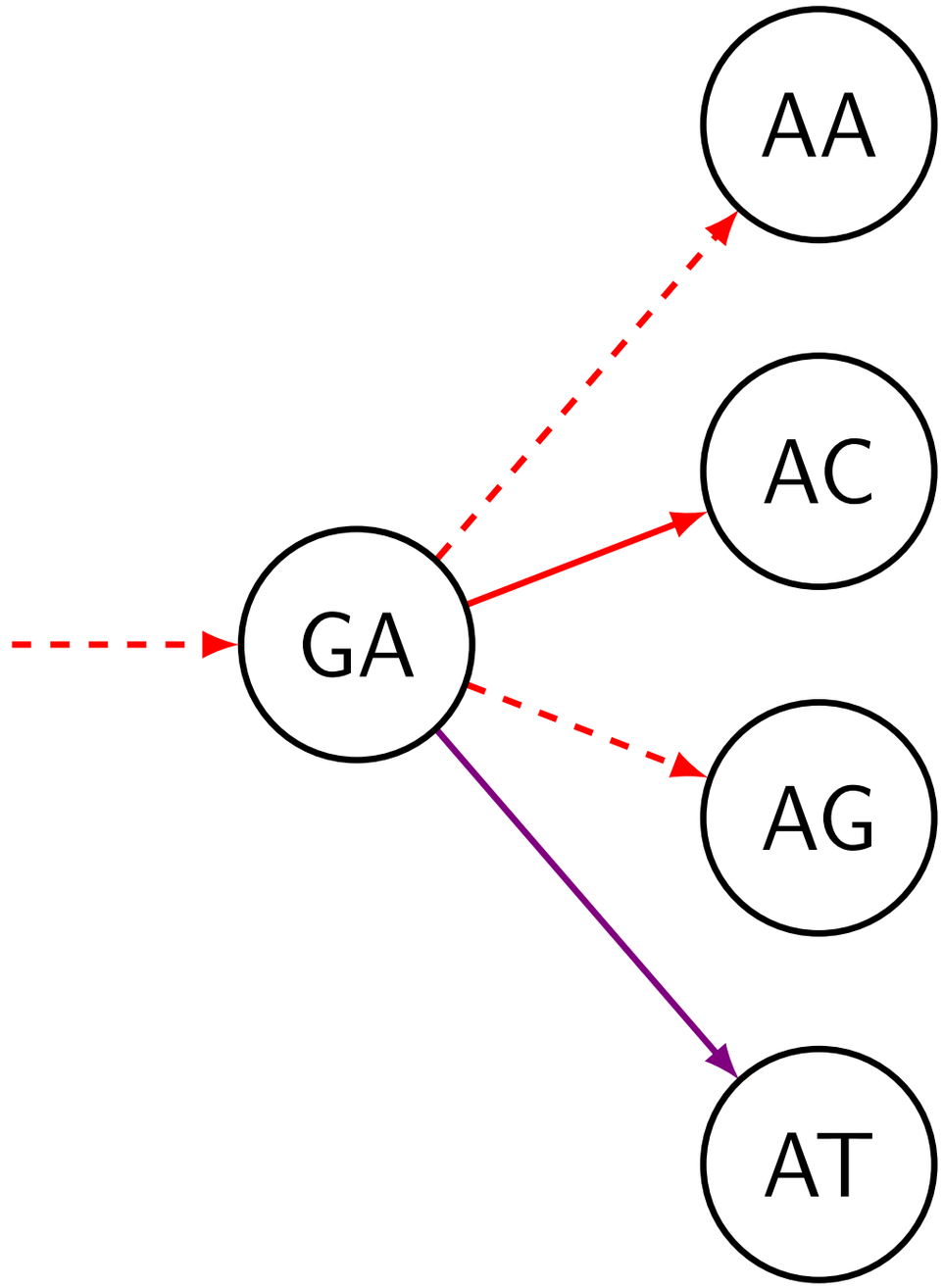
Example: Bloom Filter $= \{$ GA $\rightarrow$ AC $\}$

Is GA $\rightarrow$ AC in the set? Yes.

Is GA $\rightarrow$ AT in the set? **Maybe** no.

An Example

Bloom Filter $= \{$ GA $\rightarrow$ AC, GA $\rightarrow$ AT $\}$



The purple edge is a false positive.

The Two Pass Algorithm

How to eliminate false positives?

The Two Pass Algorithm

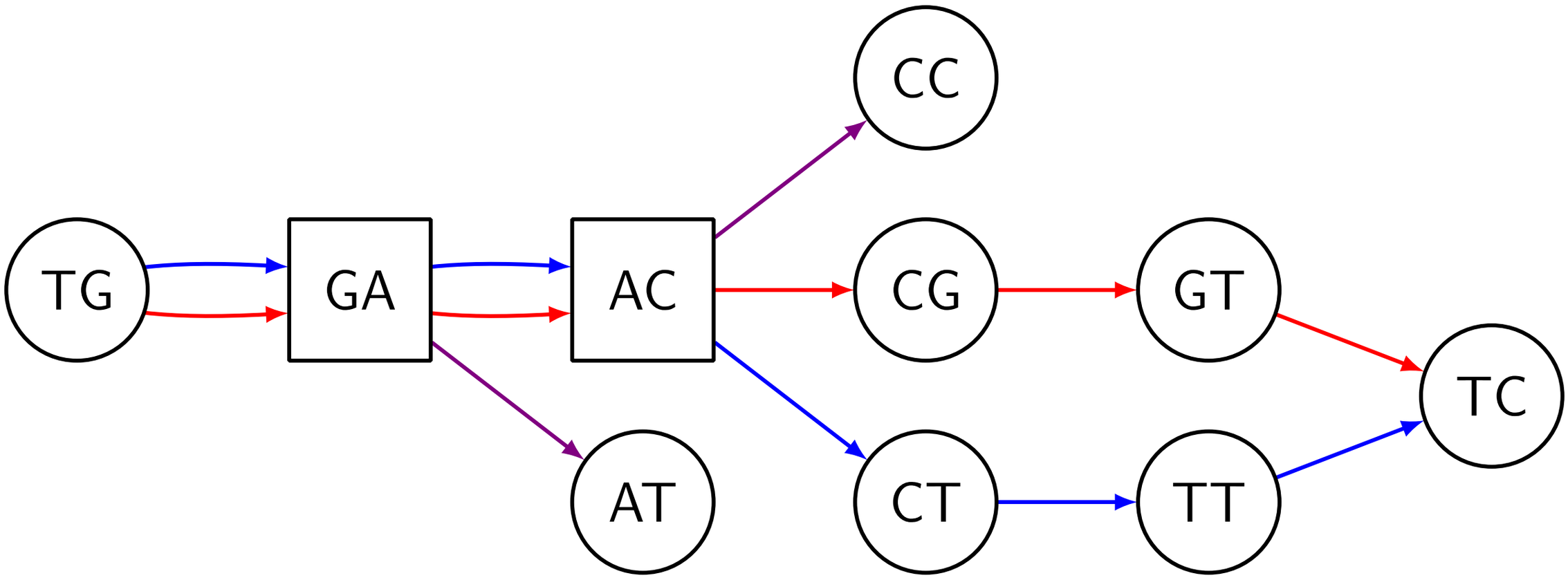
How to eliminate false positives?

Two-pass algorithm:

1. Use the Bloom filter to identify **junction candidates**
2. Use the hash table, but store **only edges that touch candidates**

An Example: the First Step

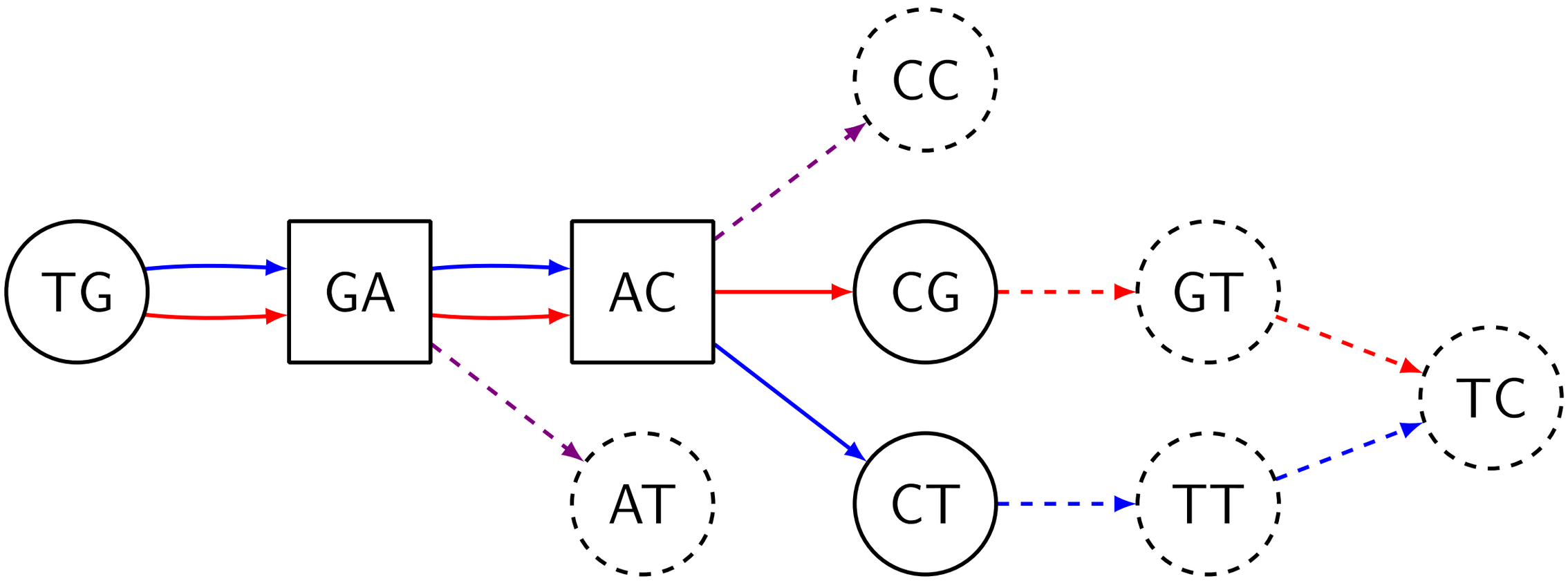
Here edges stored in the Bloom filter, purple ones are false positives:



Junction candidates: GA & AC

An Example: the Second Step

Edges stored in the hash table. We kept only edges touching junction candidates:



Junction: AC

The TwoPass Algorithm

Algorithm 2. Filter-Junctions-Two-Pass

Input: strings $S = \{s_1, \ldots, s_n\}$, integer k, a candidate set of junction positions C_{in}, integer b

Output: a candidate set of junction positions C_{out}

1: $F \leftarrow$ an empty Bloom filter of size b

2: $C_{\text{temp}} \leftarrow \mathit{Filter-Junctions}(S, k, F, C_{\text{in}})$ ▷ The first pass

3: $H \leftarrow$ an empty hash table

4: $C_{\text{out}} \leftarrow \mathit{Filter-Junctions}(S, k, H, C_{\text{temp}})$ ▷ The second pass

5: **return** C_{out}

The TwoPaCo algorithm

Algorithm 3. TwoPaCo

Input: strings $S = \{s_1, \ldots, s_n\}$, integer k, integer ℓ, integer b
Output: the compacted de Bruijn graph $G_c(S, k)$
1: Initialize counters $c_0, \ldots, c_{q-1}$ to zeroes
2: $F \leftarrow$ an empty Bloom filter of size b
3: **for** $s \in S$ **do**
4: **for** $1 \leq i \leq |s| - k + 1$ **do**
5: $h \leftarrow s[i..i + k - 1]$
6: **if** h not in F **then**
7: Insert h into F
8: $c_{f(h)} \leftarrow c_{f(h)} + 1$
9: $T \leftarrow \sum_{0 \leq t < q} c_t / \ell$ ▷ Mean number of k-mers per partition
10: $p_0 \leftarrow 0, p_\ell \leftarrow q$
11: **for** $1 \leq i < \ell$ **do**
12: $p_i \leftarrow$ biggest integer larger than p_{i-1} such that $(\sum_{p_{i-1} \leq j < p_i} c_j) \leq T$, or $\min\{\ell, p_{i-1} + 1\}$ if it does not exist.
13: $C_{\text{init}} \leftarrow$ Boolean array with every position unmarked
14: **for** $1 \leq i \leq \ell$ **do**
15: $C_i \leftarrow$ mark every position of C_{init} that starts a k-mer h with hash value $p_{i-1} \leq f(h) < p_i$
16: $C'_i \leftarrow \text{Filter} - \text{Junctions} - \text{Two} - \text{Pass}(S, k, b, C_i)$
17: $C_{\text{final}} = \cup C'_i$
18: **return** Graph implied by C_{final}, as described in Section 3.

Results

Datasets:

- 7 humans: 5 versions of the reference +
  2 haplotypes of NA12878 from 1000 Genomes
- 93 simulated humans (FIGG)
- 8 primates available in UCSC genome browser

Results

Format: minutes (GB)

Table 2. Benchmarking comparisons

	DSK+BCALM	Minia	Sibelia	SplitMem	bwt-based from Baier *et al.* (2015)		TwoPaCo	
				Single strand	Single strand	Both strands	1 thread	15 threads
62 *E.coli* ($k = 25$)	6 (1.57)	151 (0.9)	10 (12.2)	70 (178.0)	8 (0.85)	12 (1.7)	4 (0.16)	2 (0.39)
62 *E.coli* ($k = 100$)	13 (2.50)	114 (1.9)	8 (7.6)	67 (178.0)	8 (0.50)	12 (1.0)	4 (0.19)	2 (0.39)
7 humans ($k = 25$)	444 (22.44)	968 (48.09)	–	–	867 (100.30)	1605 (209.88)	436 (4.40)	63 (4.84)
7 humans ($k = 100$)	1347 (221.65)	1857 (222.0)	–	–	807 (46.02)	1080 (92.26)	317 (8.42)	57 (8.75)
8 primates ($k = 25$)	2088 (85.62)	–	–	–	–	–	914 (34.36)	111 (34.36)
8 primates ($k = 100$)	–	–	–	–	–	–	756 (56.06)	101 (61.68)
$(43 + 7)$ humans ($k = 25$)	–	–	–	–	–	–		705 (69.77)
$(43 + 7)$ humans ($k = 100$)	–	–	–	–	–	–		927 (70.21)
$(93 + 7)$ humans ($k = 25$)	–	–	–	–	–	–		1383 (77.42)

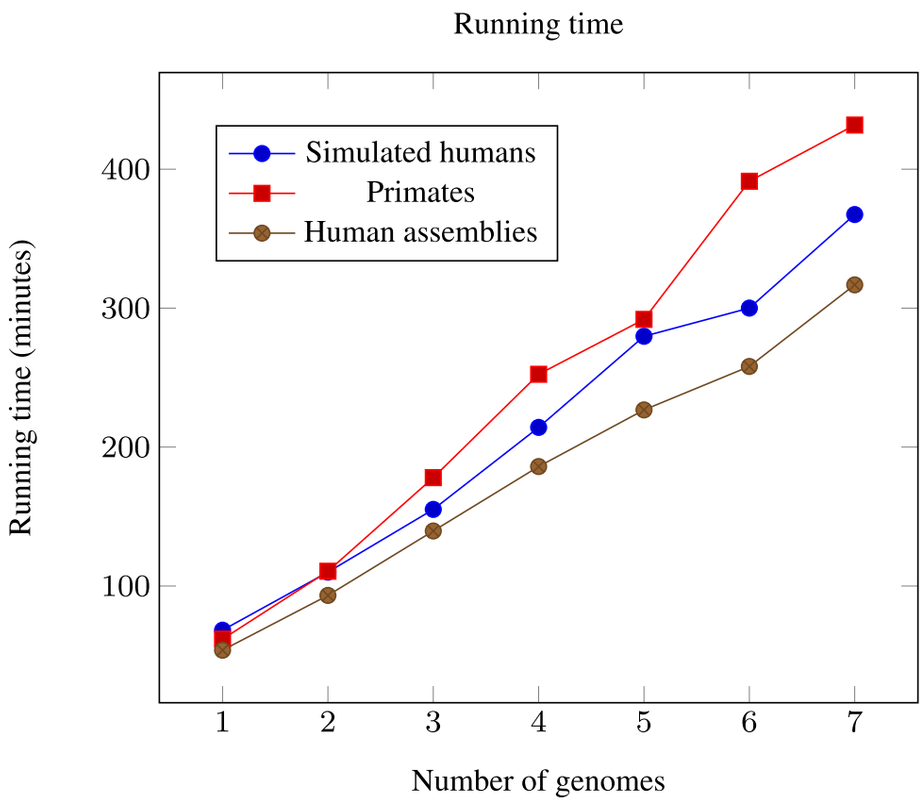
Note: Each cell shows the running time in minutes and the memory usage in parenthesis in gigabytes. TwoPaCo was run using just one round, with a Bloom filter size $b = 0.13$ GB for *E.coli*, 4.3 GB for 7 humans with $k = 25$, $b = 8.6$ GB with $k = 100$, $b = 34$ GB for primates, and $b = 69$ GB for $(43 + 7)$ and larger human dataset. A dash in the SplitMem and bwt-based columns indicates that they ran out of memory, a dash in the Sibelia column indicates that it could not be run on such large inputs, a dash in the minia column indicates that it did not finish in 48 h, a dash in the BCALM column indicates that it ran out of disk space (4 TB). A double dash indicates that the software had a segmentation fault. An empty slot indicates that the experiment was not done.

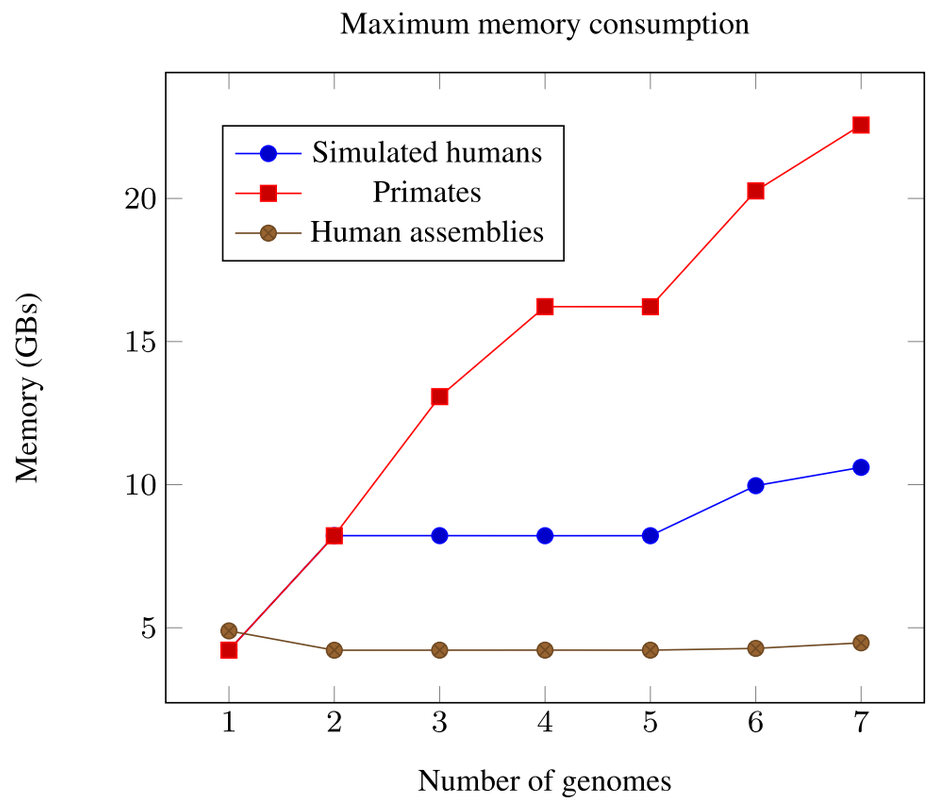
Conclusion & Future Work

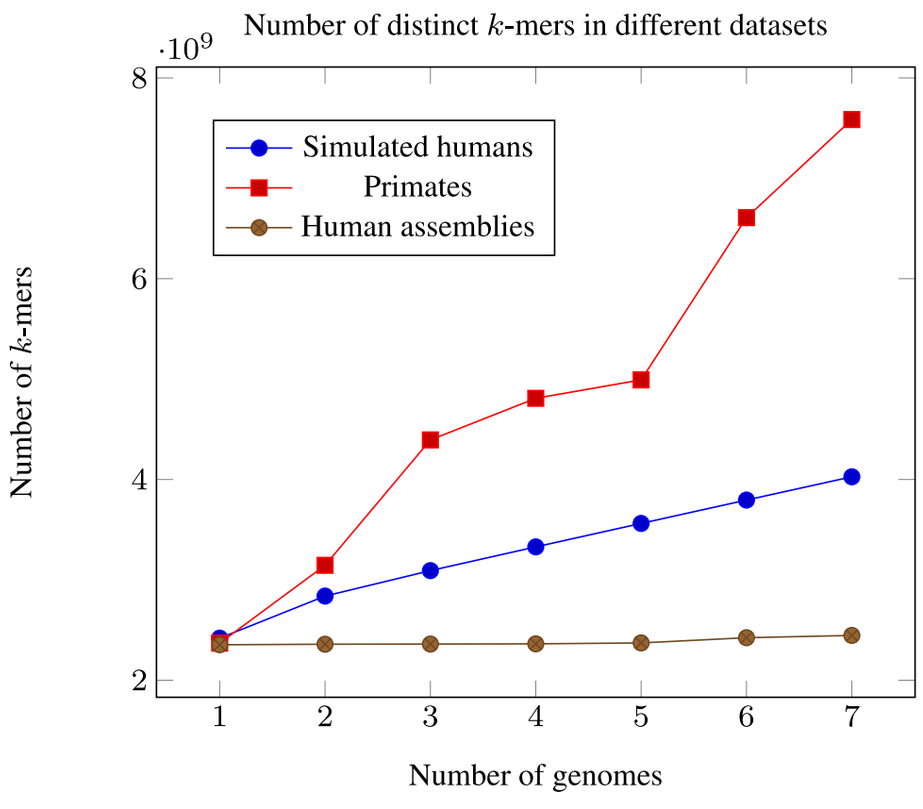
Can potentially facilitate:

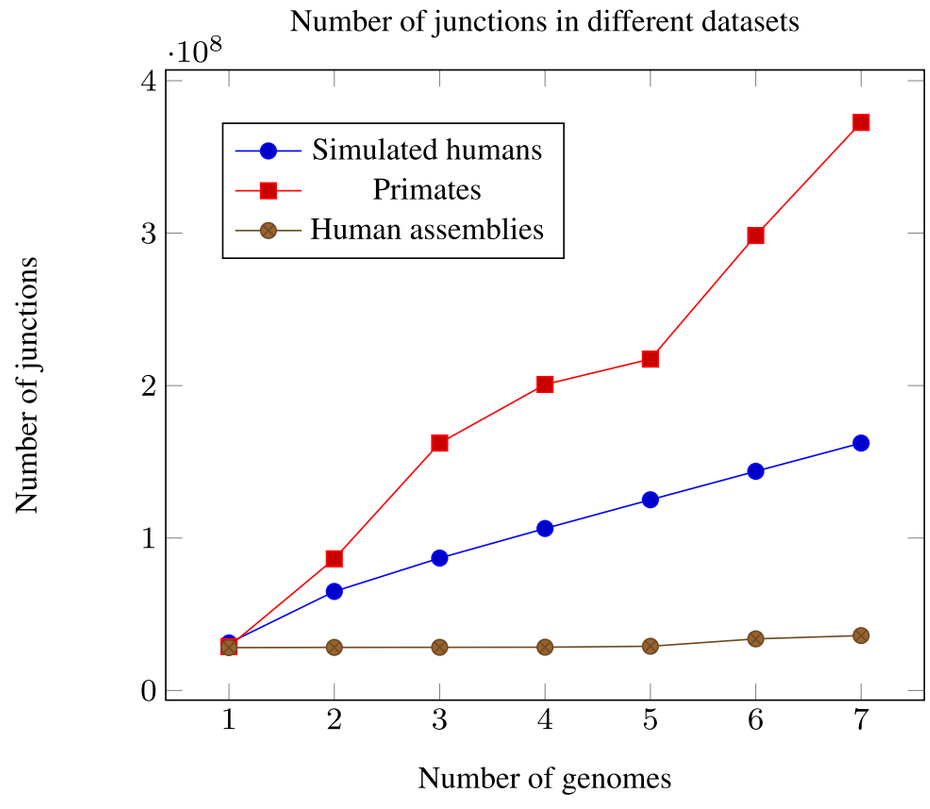
- Visualization
- Synteny mining (Sibelia)
- Structural variations analysis
- ...

Input Size vs. Performance

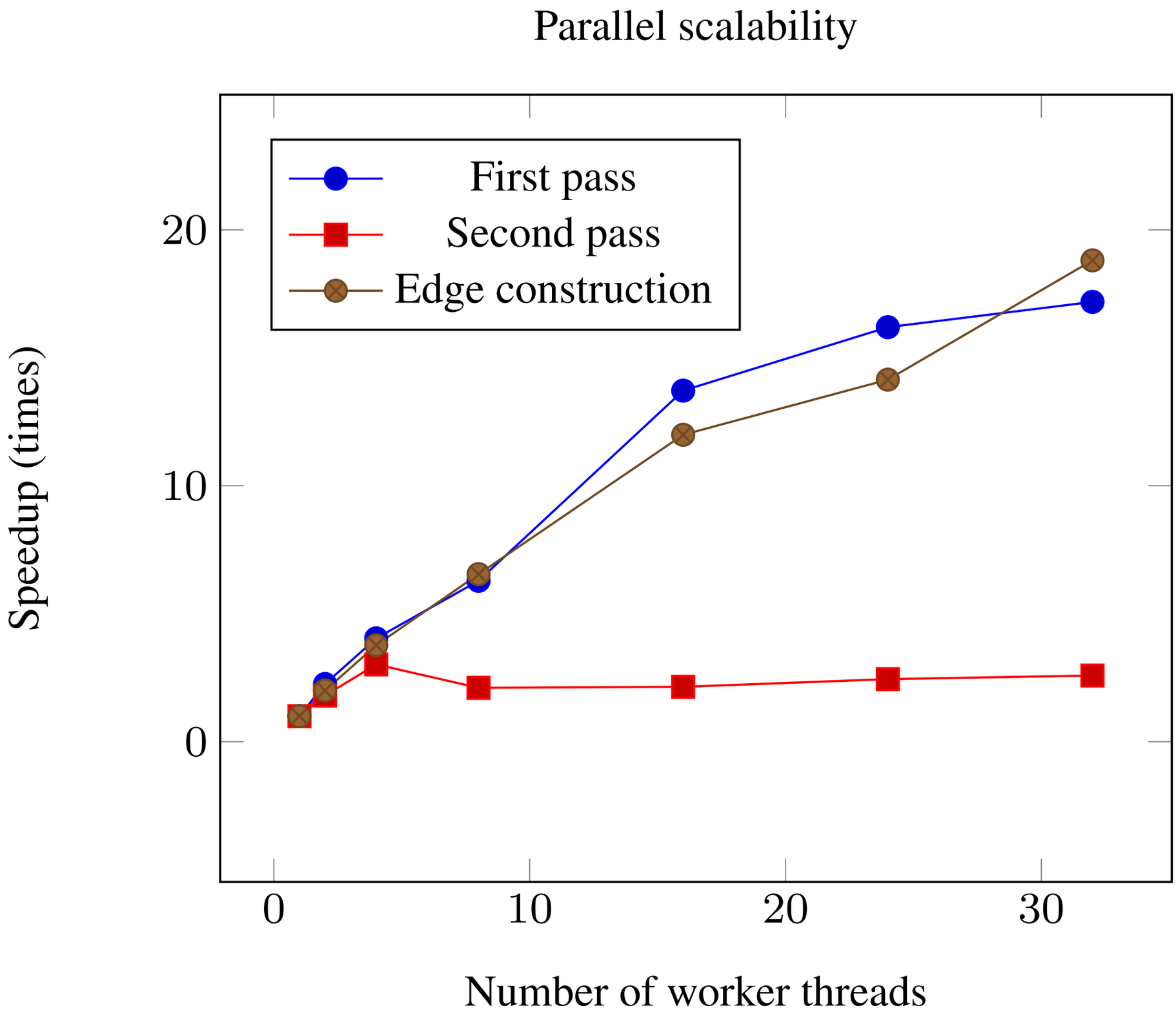








Parallel Scalability



Splitting

Table 1: The minimal number of rounds it takes to compress the graph without exceeding a given memory threshold.

Memory threshold	Used memory	Bloom filter size	Running time	Rounds
10	8.62	8.59	259	1
8	6.73	4.29	434	3
6	5.98	4.29	539	4
4	3.51	2.14	665	6