

### MINIA on a Raspberry Pi, Assembling a 100 Mbp Genome on a Credit Card Sized Computer

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# Minia on Raspberry Pi

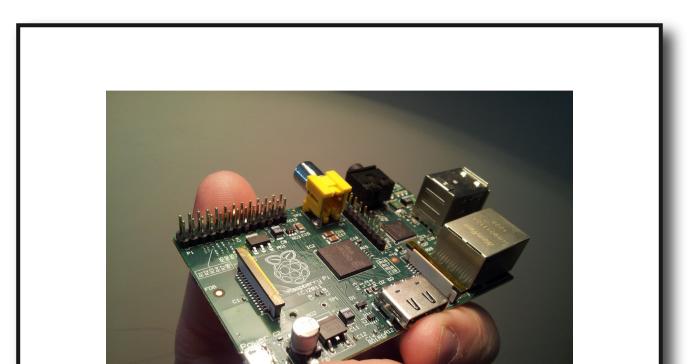
# Assembling a 100 Mbp genome on a Credit Card Sized Computer

Guillaume Collet, Guillaume Rizk, Rayan Chikhi, Dominique Lavenier

# MINIA: contig de novo assembler

This work shows that the genome assembly program MINIA is able to assemble a 100 Mbp genome on a Raspberry Pi. The MINIA software was developed to drastically reduce the memory footprint needed for genome assembly, enabling human genomes to be assembled on a desktop computer. The efficiency of MINIA is based on the DSK k-mer counting [1] and a compact de Bruijn graph data structure [2]. Here we show that it is also able to successfully assemble a genome on a very low-end, low-power system with 512 MB RAM and





a 32 GB flash drive.

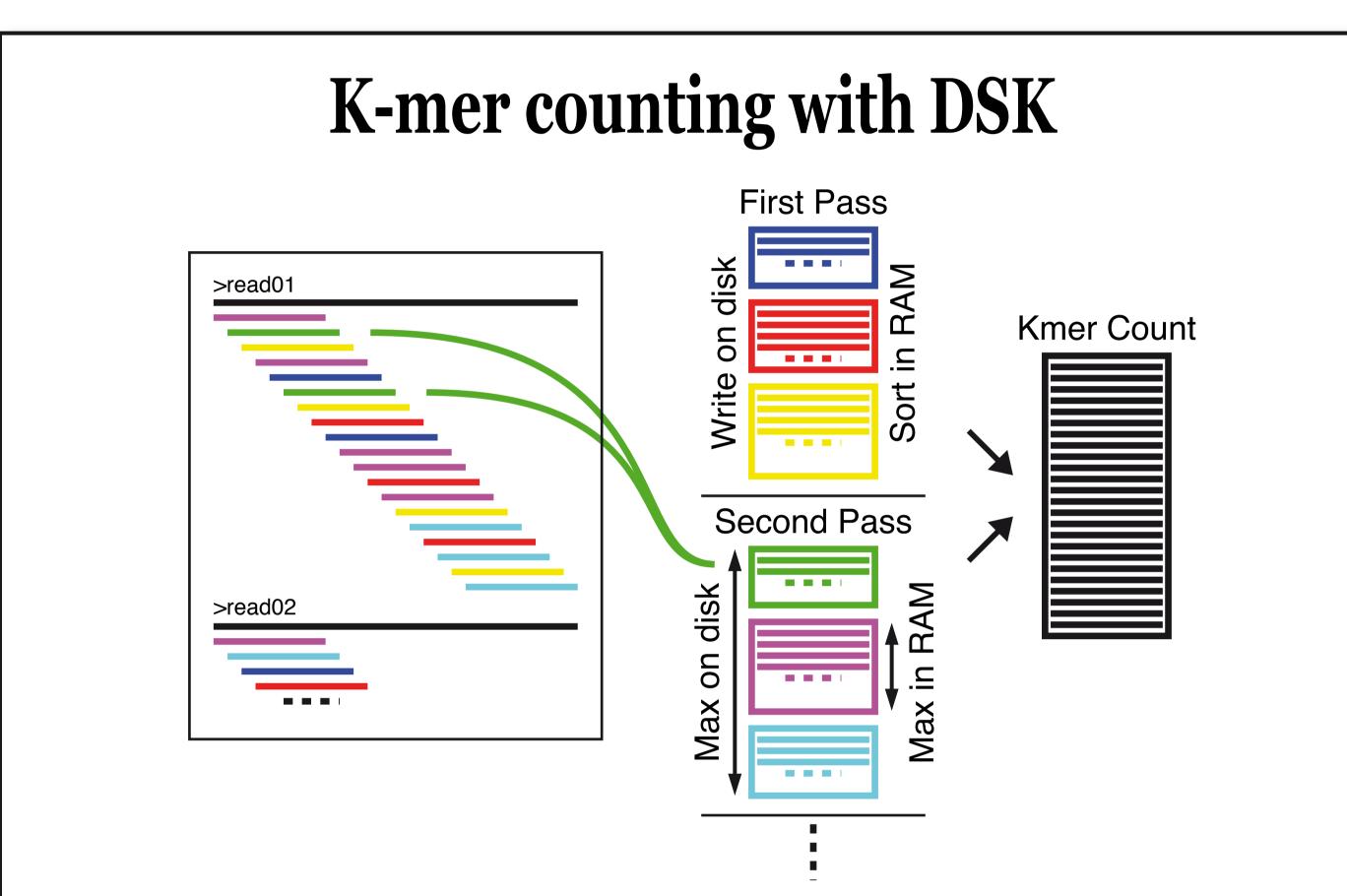
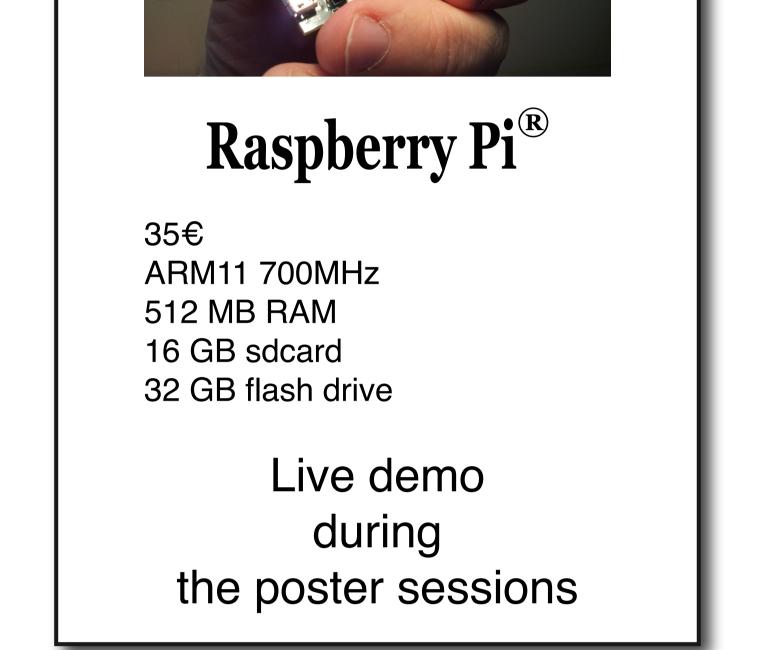


Figure 1: K-mer counting is performed by the fixed-memory and fixed-disk space algorithm DSK (Disk Streaming of K-mers). The set of k-mers is divided in partitions (colored boxes). Each k-mer is written only once on disk. Then, each partition is sorted and k-mers are counted. The trade-off between memory, disk-space, and computation allows to use DSK on a very small system.



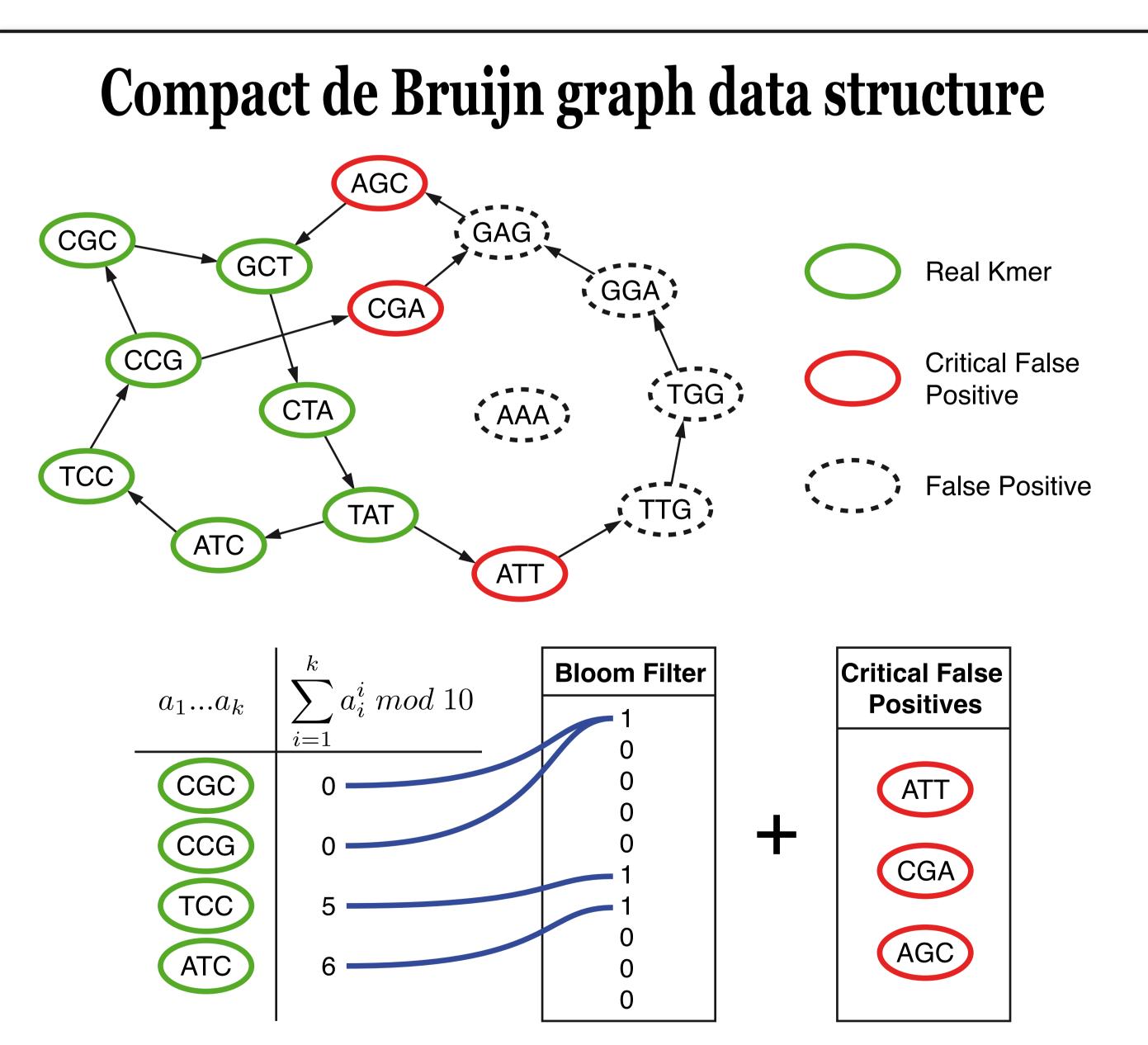
# C. elegans assembly on Raspberry Pi

Our experiment consists in assembling the nematode C. elegans. We used 33 million unfiltered paired-end reads of length 100 bp (SRR065390), covering the genome at about 64x. Paired-end information was not used.

Minia

SOAPdenovo

Velvet



System	Raspberry Pi	64 GB/Xeon E5462	64GB/Xeon E5462
CPU Time (h)	18.9	6.25	13.5
Peak memory (GB)	<b>0.2</b>	29.6	30.6
Number of contigs (K)	29.5	29.5	28.2
Longest contig (Kbp)	75.2	90.9	62.6
Contig N50 (bp)	5741	5975	6031
Sum (Mbp)	86.4	88.3	90.4
Misassemblies	12	7	419
Genome fraction $(\%)$	80.9	82.8	85.0
mismatches (per 100 kbp)	3.2	0.75	25.6

Table 1: De novo C elegant contigs assembled by Minia [2], SOAPdenovo2 [4], and Velvet [3]. Assembly quality was computed using the QUAST software [5]. MINIA and Velvet were single-threaded. For SOAPdenovo2, the CPU time is the sum for each thread.

# **MINIA applications**

Human genome assembly with less than 6 GB RAM



https://colibread.inria.fr/mapsembler2/ https://colibread.inria.fr/read2snps/





Figure 2: The probabilistic de Bruijn graph representation is obtained by inserting all the k-mers in a Bloom filter. Querying the Bloom filter for the membership of a k-mer may return a false positive answer. To avoid false positives, and consequently false branching, we propose to store the critical false positives only, in a separate structure. Thus false positives are not reachable.

## References

- [1] G. Rizk, D. Lavenier, and R. Chikhi (2013). DSK: k-mer counting with very low memory usage. Bioinformatics, 29(5), 652-653.
- [2] R. Chikhi and G. Rizk (2012). Space-efficient and exact de Bruijn graph representation based on a Bloom filter. In Lecture Notes in Computer Science (Ed.), wabi (Vol. 7534, pp. 236–248).
- [3] D. Zerbino and E. Birney (2008). Velvet: algorithms for de novo short read assembly using de Bruijn graphs. Genome Res., 18, 821-829.
- [4] R. Li et al. (2012). SOAPdenovo2: an empirical improved memory-efficient short-read de novo assembler. GigaScience, 1(1), 1-6.
- [5] G. Alexey et al. (2013). QUAST: quality assessment tool for genome assemblies. Bioinformatics, 29(8), 1072-1075.



#### **KisSplice**

A local transcriptome assembler for SNPs and AS events

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#### KisSplice

KisSplice is a software that enables to analyse RNA-seq data with or without a reference genome. It is an exact local transcriptome assembler, which enables to identify SNPs, indels and alternative splicing events. It can deal with an arbitrary number of biological conditions, and will quantify each variant in each condition. It has been tested on Illumina datasets of up to 1G reads. Its memory consumption is around 5Gb for 100M reads

It is not a full-length transcriptome assembler. This means that it will output the variable regions of the transcripts, not reconstruct them entirely. However, KisSplice results can be further aligned to a reference transcriptome (if available), or to the output of a full-length transcriptome assembler like Trinity or Oases.









# **MARISA**