

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

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Guillaume Collet, Guillaume Rizk, Rayan Chikhi, Dominique Lavenier. MINIA on a Raspberry Pi, Assembling a 100 Mbp Genome on a Credit Card Sized Computer. JOBIM - Journées Ouvertes en Biologie, Informatique et Mathématiques, Jul 2013, Toulouse, France. 2013. hal-00842027

## HAL Id: hal-00842027 https://inria.hal.science/hal-00842027

Submitted on 6 Jul 2013

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# Minia on Raspberry Pi

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

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## 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.

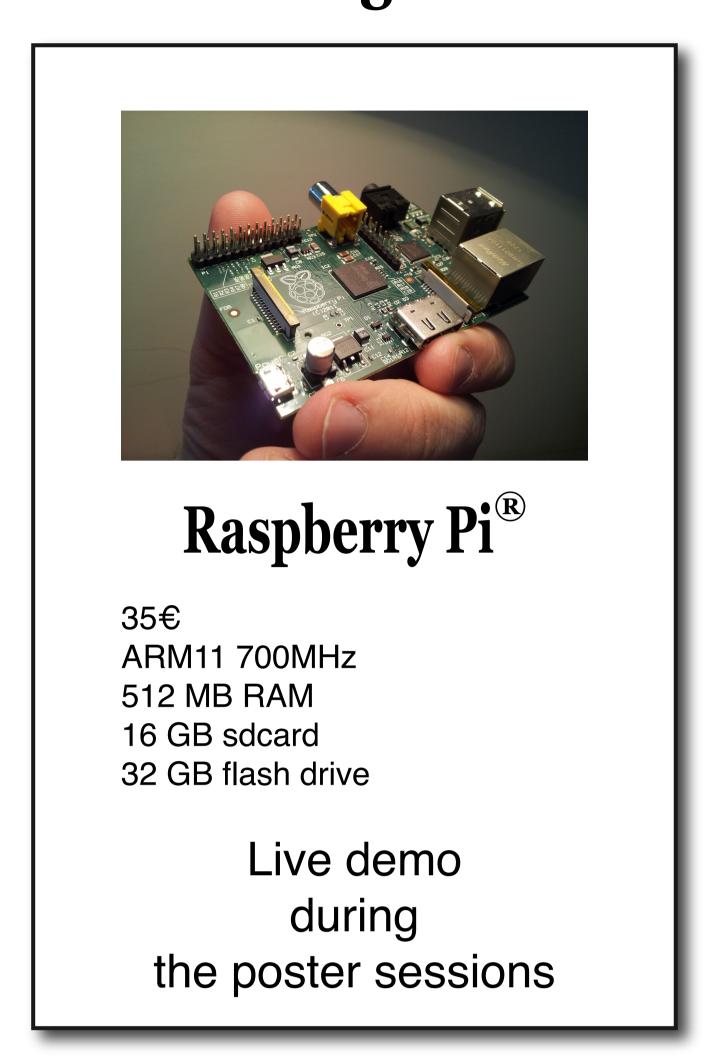
# K-mer counting with DSK First Pass Second Pass

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.

## Compact de Bruijn graph data structure GAG CGC GCT Real Kmer GGA CCG Critical False TGG Positive AAA CTA TCC False Positive TTG Critical False **Bloom Filter Positives** CGC ATT CCG CGA ATC

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.

## **Picontigotron**



## 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.

Method	Minia	SOAPdenovo	Velvet
System	Raspberry Pi	64GB/Xeon E5462	64GB/Xeon E5462
CPU Time (h)	18.9	6.25	13.5
Peak memory (GB)	0.2	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/



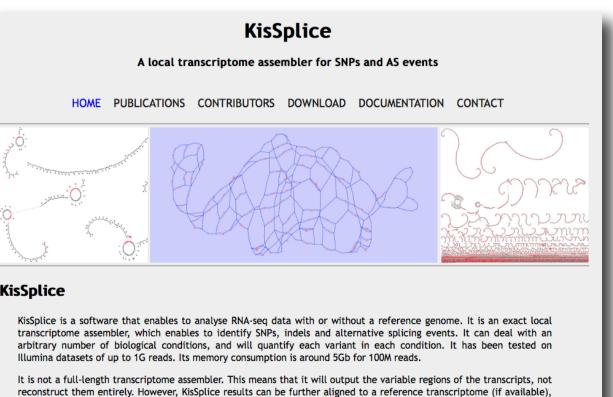
http://minia.genouest.org/



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- [5] G. Alexey et al. (2013). QUAST: quality assessment tool for genome assemblies. Bioinformatics, 29(8), 1072-1075.

http://kissplice.prabi.fr/













or to the output of a full-length transcriptome assembler like Trinity or Oases.