

Abstract :

- The Problem is De Bruijn graphs Consume Large memory.
- We have a solution to reduce this complexity by compacting paths with single vertices
- On the other hand, this way has a problem because it requires the graph to be in the memory.
- Bifrost features is a parallel and algorithm which saves the memory ,also.
- Our Parallel algorithm enable the direct construction of the compacted graphs.
- Bifrost has some added features such as indexing, editing, querying and the graph coloring method.

Introduction :

- De Bruijn graph is an abstract data structure used in computational biology as a tool for genome assembly.
- Since 2008, the de Bruijn graph genome assemblers range have been released.
- De Bruijn graph-based methods are nonetheless used to assemble and correct long reads.
- De Bruijn graphs have found widespread by solving a variety of problems such as de novo transcriptome assembly, variant calling, short read compression, short read correction, long read correction, and short read mapping to name a few.
- There is a colored de Bruijn graph which is a variant of the de Bruijn graph.
- The colored de Bruijn graph keeps track of the source of each vertex in the graph.
- The initial application was for assembly, genotyping, pan-genomics, variant calling, and transcript quantification methods.

Related Works:

- **An Eulerian path approach to DNA fragment assembly**

_ The reduction of the fragment assembly to a variation of the classical Eulerian path problem that allows one to generate accurate solutions of large-scale sequencing problems.

_ we do a very counterintuitive (some would say childish) thing: we cut the existing pieces of a puzzle into even smaller pieces of regular shape.

<https://www.pnas.org/content/98/17/9748.short>

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

_ de Bruijn graphs have been proposed as a data structure to facilitate the analysis of related whole genome sequences, in both a population and comparative genomic settings.

_ However, current approaches do not scale well to many genomes of large size (such as mammalian genomes).

<https://academic.oup.com/bioinformatics/article/33/24/4024/2725383?login=true>

- **From squiggle to basepair: computational approaches for improving nanopore sequencing read accuracy**

_ A major limitation of nanopore sequencing is its high error rate, which despite recent improvements to the nanopore chemistry and computational tools still ranges between 5% and 15%.

_ Here, we review computational approaches determining the nanopore sequencing error rate. Furthermore, we outline strategies for translation of raw sequencing data into base calls for detection of base modifications and for obtaining consensus sequences.

<https://link.springer.com/article/10.1186/s13059-018-1462-9>

- **Disk-based compression of data from genome sequencing**

_ We propose overlapping reads compression with minimizers, a compression algorithm dedicated to sequencing reads (DNA only).

_ Our method makes use of a conceptually simple and easily parallelizable idea of minimizers, to obtain 0.317 bits per base as the compression ratio, allowing to fit the 134.0 Gbp dataset into only 5.31 GB of space.

<https://academic.oup.com/bioinformatics/article/31/9/1389/200464?login=true>

