MAP2COV: Convert NGS data guide

MAP2COV is a free converter of NGS alignments file
It has been developed by the Laboratoire Pathogenèse des
Bactéries Anaérobies of the Institut Pasteur.

http://cov2html.sourceforge.net/

« One Click » graphical interface Tcl/Tk



Windows

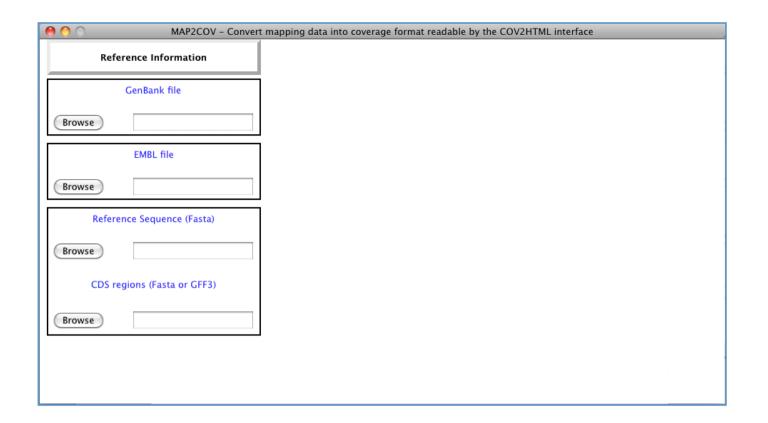


MacOSX

Command line graphical interface Tcl/Tk

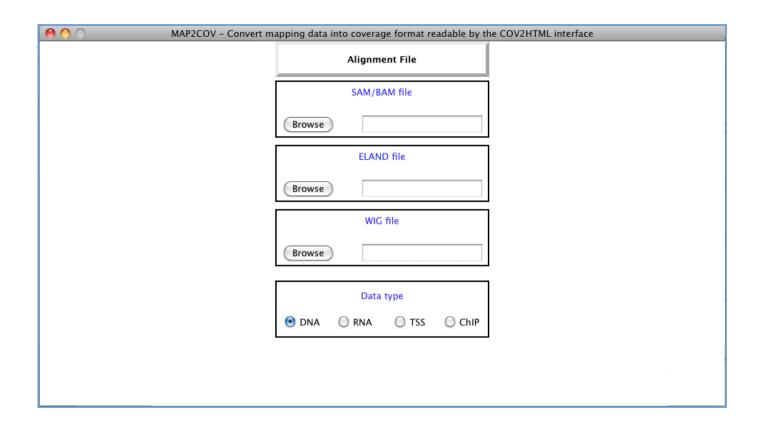


Linux



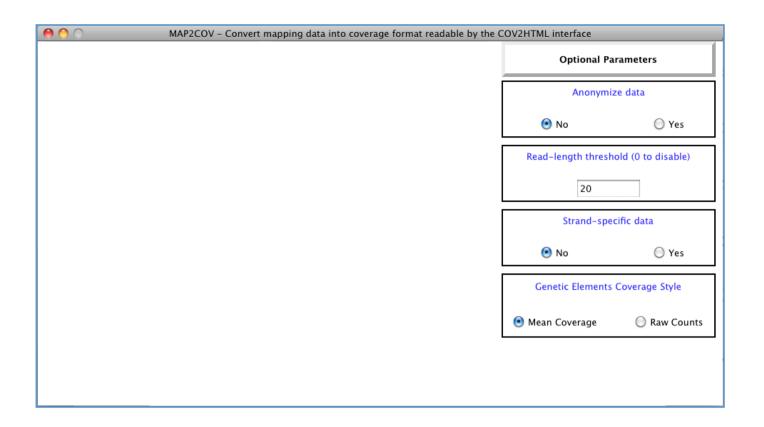
Reference Information

A file containing the annotated reference sequence previously used for the mapping assembly, it can be in GenBank, EMBL format or two files, a file containing the reference sequence in Fasta format associated with a file containing CDS regions annotations in gff3 format or features nucleotide sequence in fasta format.



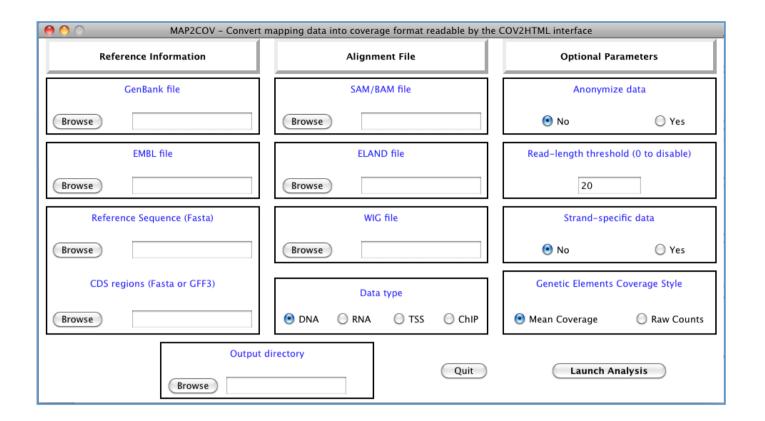
Alignment File

An alignment map file in 'SAM'/'BAM', 'ELAND' or 'WIG' format. Then users have to select the data type contained in the alignment file, 'DNA' (DNA-seq), 'RNA' (RNA-seq), 'TSS' (Transcriptional Start Site) or ChIP (Chromatine ImmunoPrecipitation).



Optional Parameters

Anonymize data (default: **No**): replace ID of genes with unique identifiers numbered from the origin and remove their description. Read-length threshold (default: **20** bp): remove reads from the alignement file shorter than the threshold. Strand-specifc data (default: **No**): generate two strand-specific files from the alignment. Genetic Elements Coverage Style (default: Mean Coverage): coverage of genes and intergenic regions is calculated either as the **Mean coverage** or as the number of read that match on them (**Raw Counts**).



MAP2COV Tcl/Tk graphical interface.

Reference information: GenBank, EMBL file or Genome and Annotation files.

Alignment File: SAM/BAM, Eland or Wig files. Data type: DNA, RNA, TSS, ChIP.

Optional Parameters: Anonymize, Read-length, Strand-specific or Genetic Elements Coverage Style.