

Sequana: a set of Snakemake NGS pipelines

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September 25th 2017, Institut Curie, Paris

Introduction

Motivation

Development driven by the Biomics Pole at Pasteur Institute, which involves many aspects of NGS including :

https://research.pasteur.fr/en/team/biomics/

- De novo and targeted sequencing of viruses, prokaryotes and eukaryotes
- Variant (SNP, indel, large rearrangements) detection
- Human and Mouse SNP detection by array
- Transcriptional analysis (RNA-Seg) for both prokaryotes and eukaryotes
- 16S and deep-sequencing metagenomic studies (mouse, human, and other environments)
- Bottom-up whole proteomic analysis and quantification
- Analysis of a wide range of post-translational modifications
- Determination of the dynamics of protein complexes.
- Epigenetics (CHIP-Seq, methylation studies)
- Projects involving two or more techniques (i.e. proteogenomics, single-cell DNA/RNA analysis)



A glue language, a scientific language



A pipeline framework mixing Python and Makefile Köster, Johannes and Rahmann, Sven. Snakemake - A scalable bioinformatics workflow engine. Bioinformatics 2012.



Dedicated standalone such as genome coverage characterisation.

D. Desvillechabrol, C. Bouchier, S. Kennedy, T. Cokelaer
Detection and characterization of low and high genome coverage
regions BioRxiv https://doi.org/10.1101/092478

Snakemake

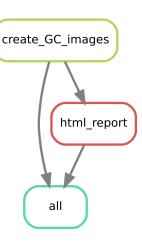
What is snakemake?



- Clusters can be used with minimum efforts (no intrusive code)
- Workflows can be run from or up to a given rule
- Nice logging system to follows the status
- Suspend / Resume
- Various code can be integrated: R, bash, and of course Python

Pipeline example: minimalist (GC content)

```
# list of FastO files without extension
files = ["A", "B", ....]
rule all:
    input: expand("{data}.png", data=files)
        "index.html"
rule create_GC_images:
    input: "{data}.fastq.gz"
    output: "{data}.png"
    run: # CODE TO COMPUTE IMAGES
rule html_report:
    input: expand("{data}.png", data=files)
    output: "index.html"
    run: # CODE TO CREATE HTML
```



Execution

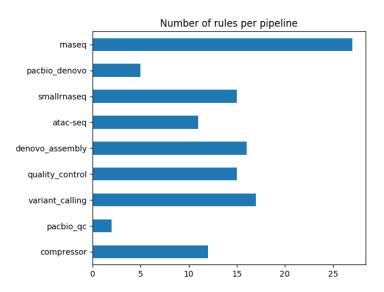
From a shell:

snakemake -s gc_minimalist.rules

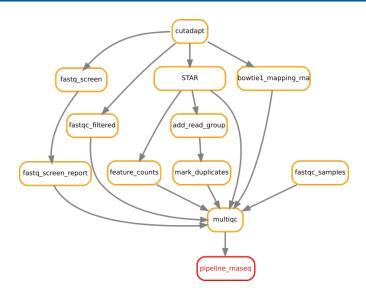
More options if a configuration file is required, or execution is on a cluster, or ... something goes wrong.

Sequana

Pipelines available in Sequana



Pipeline RNA-seq



Markduplicates rule

```
rule mark duplicates:
    """DOCSTRING"""
    input:
        mark duplicates input
    output:
        bam = mark duplicates output,
        metrics = mark duplicates metrics
    log:
        out = __mark_duplicates__log_std,
        err = __mark_duplicates__log_err
    params:
        remove = config["mark_duplicates"]["remove"],
        tmpdir = config["mark_duplicates"]["tmpdir"]
    shell:
        (picard MarkDuplicates I={input} O={output.bam} \
            M={output.metrics} REMOVE_DUPLICATES={params.remove} \
            TMP_DIR={params.tmpdir} && samtools index {output.bam}) \
            > {log.out} 2> {log.err}
        11 11 11
```

Snakefile

YAML configuration file

```
mark duplicates (picard-tools) allows to mark PCR duplicate in
 BAM files
 : Parameters:
  - do: if unchecked, this rule is ignored. Mandatory for RNA-SeQC
       tool.
  - remove: If true do not write duplicates to the output file
          instead of writing them with appropriate flags set.
          Default value: false. This option can be set to
          'null' to clear the default value.
          Possible values: {true, false}
   tmpdir: write tempory file on this directory
          (default /tmp/)
mark duplicates:
   do: ves
   remove: no
   tmpdir: "/local/scratch/"
```

Using command line

One command line to initiate the pipeline

- The sequana executable creates a directory with the project name
- The directory contains all the necessary files (config, snakefile)

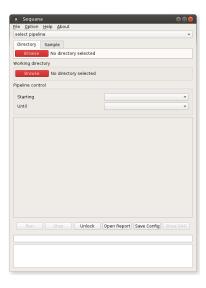
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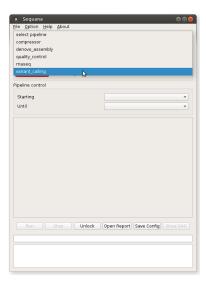
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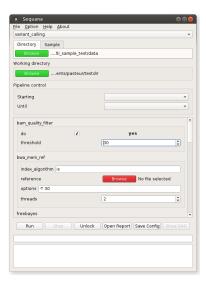
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- Sequana developers want to expose their pipelines dynamically
- Snakemake developers want to use Sequanix ;-)



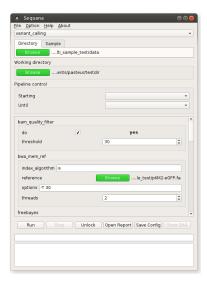
- Interface developed with PyQT5 and Python
- Wrap our snakemake pipelines to ease the usage
- Usable on our cluster, which allows X11



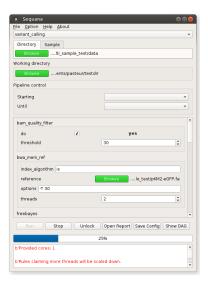
1 Choose a pipeline



- Choose a pipeline
- Set input and output



- Choose a pipeline
- Set input and output
- 3 Fill the config form



- Choose a pipeline
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- Fill the config form
- 4 Run the pipeline



- Choose a pipeline
- Set input and output
- Fill the config form
- 4 Run the pipeline
- Finished!

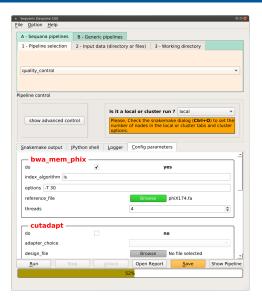
Automatic tooltips

BWA used to remove a contaminant

Parameters:

- · do: if unchecked, this rule is ignored
- reference_file: the name of the reference file to be found in the analysis directory. If set to phiX174.fa, it is downloaded automatically from Sequana, otherwise you will need to copy it yourself in the working directory.
- index_algorithm: the BWA index algorithm
- options: any options recognised by BWA tool
- threads: number of threads to be used

Last version of Sequanix



Desvillechabrol, D., Legendre, R., Rioualen, C., Bouchier, C., van Helden, J., Kennedy, S., & Cokelaer, T. (2017). Sequanix: A Dynamic Graphical Interface for Snakemake Workflows. bioRxiv, 162701.

Continuous Integration

Versioning, Test and Documentation



https://github.com/sequana/sequana



Continuous Integration on Travis with 100 tests with 75% coverage



Uses Sphinx (RST syntax) to document the source code and provides user guide.



Updated after each commits on sequana.readthedocs.io