# Long read de novo assembly exercise

#### **Instructions**

Document your work well.

Add Assembly and BUSCO results here:

**Google sheets hyperlink** 

### Set up

- 1. Set up a new directory structure for a de novo genome sequencing projects
- 2. **Create a private github repository** for the project. All directories and file except sequence files should be included.

The repo should have a readme file with a detailed description of what you have done in the project.

3. Add dagahren as a collaborator.

Note! It is important that you throughout your project think of ways of checking what you did.

For example check that the download was complete or that a software ran to completion without errors.

Use top to follow the process and see what the commands are doing.

#### **Dataset**

The dataset is a HiFi Pacbio reads from the yeast Saccharomyces cerevisiae.

The data can be downloaded from NCBI Sequence Read Archive (SRA)

Download the reads for the following page or use sra-tools:

SRR13577846

Make the Data directory readonly!

## **Computational resources**

When using the Course server, max number of CPUs (threads) to be used is 10!

### **Software**

Make a separate directory for each run & investigate

- Quality assessment using fastqc
- de novo assembly (separate dirs if you try more that one!)
- Quast quality assessment
- BUSCO assessment

Make sure that you are documenting all work you do in the README file.

Note the version of each software you use

Time the run of each command so that you can report how fast the assembly was.

Remember to commit your changes and push to Github

Pick an assembler suitable for PacBio Sequencing data or choose freely from publications (but give the proper reference):

For example you can use:

Hifiasm

https://github.com/chhylp123/hifiasm

https://hifiasm.readthedocs.io/en/latest/index.html

## **Quality assessment**

Use the following tools to assess the sequences and the genome assembly using:

- FastQC
- Quast
- BUSCO
- MultiQC
- Other?