Group C: Report DNA/RNA Sequencing Course Bread and Cheese

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Abstract

This is the final report of our work during the $2015\ DNA/RNA\ Sequencing\ Course$, which is part of the joint master's programme in Bioinformatics of the Universities of Bern and Fribourg.

We present the result of two tasks: (1) finding novel regulators/targets of TORC1 in genomes of yeast mutants, and (2) building and annotating a de-novo assembly of the genomes of five strains of Lactobacillus Paracasei. For both tasks, we're using free-of-charge and open source software tools which are well known in their respective field. (Quality control, assembling, SNP calling, annotation, visualization).

Using results from the first task, we are able to identify a candidate mutant, which shows interesting SNP mutations that are not yet known to be interferring with the TORC1 pathway.

Our main result from the second task is the newly assembled and annotated genomes of five strains of Paracasei.

Filename	Total Sequences	Sequence length	%GC
$M1_S237_R1_001.fastq.gz$	3493039	35-151	37
$M1_S237_R2_001.fastq.gz$	3493039	35-151	38
$M14_S196_R1_001.fastq.gz$	5619726	35-151	37
$M14_S196_R2_001.fastq.gz$	5619726	35-151	37
$M16_S224_R1_001.fastq.gz$	5311850	35-151	37
$M16_S224_R2_001.fastq.gz$	5311850	35-151	37
$M18_S257_R1_001.fastq.gz$	5483798	35-151	37
$M18_S257_R2_001.fastq.gz$	5483798	35-151	37
$M21_S250_R1_001.fastq.gz$	4089470	35-151	36
$M21_S250_R2_001.fastq.gz$	4089470	35-151	36
$M24_S240_R1_001.fastq.gz$	4029845	35-151	36
$M24_S240_R2_001.fastq.gz$	4029845	35-151	36

Table 1: Yeast Raw Data

Part I

Identification of novel regulators and/or targets of TORC1 in yeast mutants

1 Introduction

What is the goal?

Identify mutations that confer a growth phenotype after rapamycine treatment.

Asses, whether the involved proteins are already known to be involved in the TORC1 pathway.

If there are new candidates, we were trying to characterize their function as new regulators or targets of the TORC1 pathway.

2 Data and Methods

Paired DNA libraries were prepared in advance and handed over to us in gzipped FASTQ file format. Table 1 shows some basic information about the data.

TODO: Mention Reference genome

Data processing was done in multiple stages using free-of-charge and opensourse software tools. The main steps were executed manually on the High Performance Computing Cluster of the University of Bern.

Filtering tewltj

3 Results

We identified the following SNPs to be interesting. Table xy shows these observed mutations.

3.1 Mutant I

Already known to be interferring w/ the TORC1 pathway. (Reference)

3.2 Mutant II

Already known to be interferring w/ the TORC1 pathway. (Reference)

3.3 Mutant III

Shows some significant differences. ...interesting.

4 Analysis and Discussion

Part II

Paracasei

1 Introduction

(Very) Short introduction to cheese making. Collaboration w/ Agroscope. Sequencing of different starters that they have in their library.

What is the goal of our work?

Which genes are associated with the growth phenotype?

What is the biochemical relation between VSC and the growth phenotype?

2 Data and Methods

We had the data from ...?

We used the following Pipeline...

Although the overall quality of the data was acceptable, we used sickle to correct and filter...

3 Results

4 Analysis and Discussion

- A Appendix Some picture of data processing pipeline
- B Appendix Second picture of data processing pipeline