Reuben's personal guide to Hi-C analysis

December 14, 2016

Introduction

Data Processing

Download Hi-C files

Files analysed throughout this guide were all obtained form the short read archive (SRA) under the accession .

Sequence Quality

Read mapping and filtering

Generate contact matrix

Use homer package to generate tag dir.

Use homer to convert tag dir to generate raw contact matrix.

Filter rows and columns of contact matrix. Remove rows and columns with poor read mapping.

Use HiCorrector to normalise contact matrices.

Normalising contact matrices

Downstream analysis

Topological domains

Genomic compartments

```
#!/usr/bin/perl

use strict;
use warnings;

for (my $i = 1; $i < 100; $i++) {
   if ($i == 99) {
      print $i." Luftballons reached!\n\n";
   } else {
      print $i." Luftballons...\n";
   }
}
exit;</pre>
```

Listing 2: Nena would be proud.

```
#!/bin/bash
    # Script to perform Hi-C read mapping for paired end reads
       unising hicup software
    # script is written for use on adelaide university phoenix
        cluster
    # Invoked by:
    # READPATH=<path to fastq file dir> GENOME=<name of genome
        assembly> sbatch alignHiC.sh
   #SBATCH −p batch
    \#SBATCH -N 1
    #SBATCH -n 16
    #SBATCH ---time=2-00:00
    #SBATCH ---mem=32GB
    # Notification configuration
   #SBATCH — mail-type=END
#SBATCH — mail-type=FAIL
#SBATCH — mail-user=reuben.buckley@adelaide.edu.au
    #SBATCH ---array=0-1 #result of 'ls *.fastq.qz | wc -l' less
        one.
   module load R
25
    FILES=($(ls $READPATH | rev | cut -c 9- | rev | uniq))
   mkdir ./${FILES[$SLURM_ARRAY_TASK_ID]}
   date > $GENOME.${FILES[$SLURM_ARRAY_TASK_ID]}.log.txt
30
    hicup --bowtie2 /apps/software/Bowtie2/2.2.9-GCC-5.3.0-
        binutils -2.25/bin/bowtie2 --keep --longest 800 --shortest 100 --threads 16 --index ../../genomes/genomeIndex/$GENOME
        /${GENOME}knownChr --digest ../../genomes/genomeDigest/
        $GENOME/DigestKnownChr* --outdir ${FILES[
        $SLURM_ARRAY_TASK_ID]} $READPATH${FILES[
        $SLURM_ARRAY_TASK_ID]}_1.fastq $READPATH${FILES[
        $SLURM_ARRAY_TASK_ID]}_2.fastq &>> $GENOME.${FILES[
        $SLURM_ARRAY_TASK_ID]}.log.txt
    date &>> $GENOME.${FILES[$SLURM_ARRAY_TASK_ID]}.log.txt
```

Listing 1: Bash script used to map reads on phoenix