

RNA-Seq Pipeline

Input folder: **rawdata**

fastq files (zipped)
naming convention for paired end:
sample1.R1.fastq.gz
sample1.R2.fastq.gz

[targets.txt](#)

sample	sample name
file	readcount filename for DE
group	group name of sample
replicate	replicate id

[contrasts.txt](#)

for DE analysis, give one contrast per line, e.g.: KOvsWT=(KO-WT).
 Group names as in targets.txt

[bpipe.config](#)

define workload manager resources
 Set executor="local" if not needed.

Pipeline parameter

+ [essential.vars.groovy](#):
 provide essential parameter
 + [module.name.vars.groovy](#):
 provide module specific parameter (if needed)

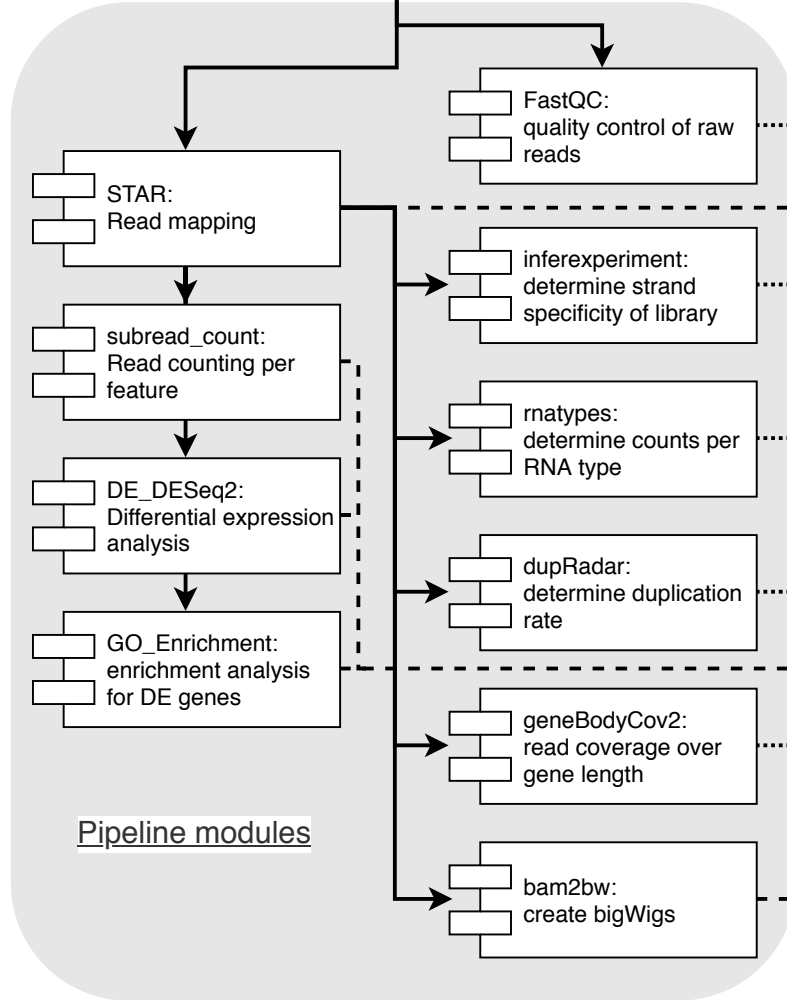
Bioinformatics tools

+ [tool.locations.groovy](#):
 + [tool.versions.groovy](#):
 provide locations and versions of required analysis tools

Reference Genome

+ STAR indexed genome
 + feature definition: gtf
 provide file locations in [essential.vars.groovy](#)

«main pipeline»
[rnaseq.pipeline.groovy](#)
 - editable pipeline definition
 - loads respective pipeline modules
 - loads essential and module-specific parameter
 - executes pipeline



DEreport.html

html output file containing final results of all quality control and analysis steps.

Output folder: **reports**

+ DEreport.Rmd
 + DE.shinyrep.helpers.R
Modify Rmd file with custom analysis parameters and thresholds as needed

shinyReports:
 final report and data visualization

Output folder: **mapped**

bam files

Output folder: **qc**

module output files for qc

Output folder: **results**

count files
 DE results
 GO enrichments

Output folder: **tracks**

bigWig tracks for visualisation

Output folder: **logs**

log-files collected for each module

Legend

bpipe module

pipeline component

table / file

input/output repository

Italics: user input needed
 Different arrow shapes are used for better clarity