

## BIBS platform

Maturing In the Time of Coronavirus

Logo BIBS ?

14/09/2020

Magali Hennion

# BIBS : BioInformatics and BioStatistics

## Steering committee (CoPil)

Valérie Lallemand-Mezger

Claire Rougeulle

Olivier Kirsh

Jean-François Ouimette

Florent Hubé

Magali Hennion

Pierre Poulain (IJM)



## Active members

Magali Hennion

Olivier Kirsh

# Call for projects

## **"Projets Pilotes"**

Call initiation : March 3rd

Goal for the platform : building an efficient working environment

- how to communicate with the project holder / between us ?
- how to do the analyses to make them reproducible, reusable, independent of the computer/cluster ?

# Call for projects

## "Projets Pilotes"

Call initiation : March 3rd

Goal for the platform : building an efficient working environment

- how to communicate with the project holder / between us ?
- how to do the analyses to make them reproducible, reusable, independent of the computer/cluster ?

March 30th → 4 projects (Nataliya, Madeleine, Agathe, Guillaume)

CoPil meeting -> decision made public on April 21st

- Try to help everyone
- Start developing a RNA-seq analysis pipeline on Madeleine's data
  - Use it on other datasets, adapt it to other kind of analysis, ...

# Why using a workflow manager?

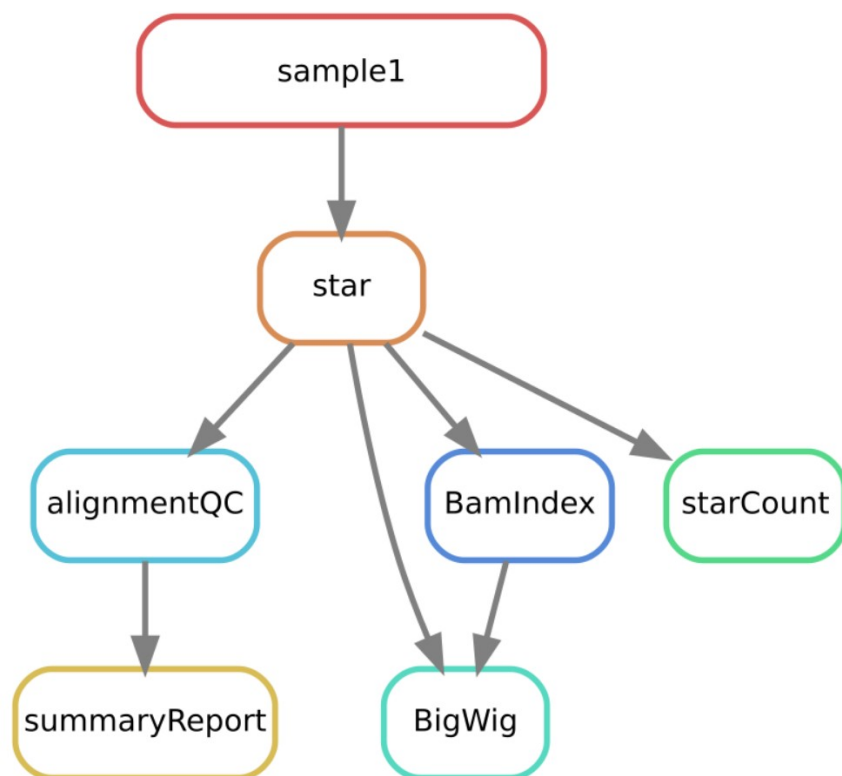


nextflow



**One analysis** : multiple steps, multiple samples

Simple example, one sample

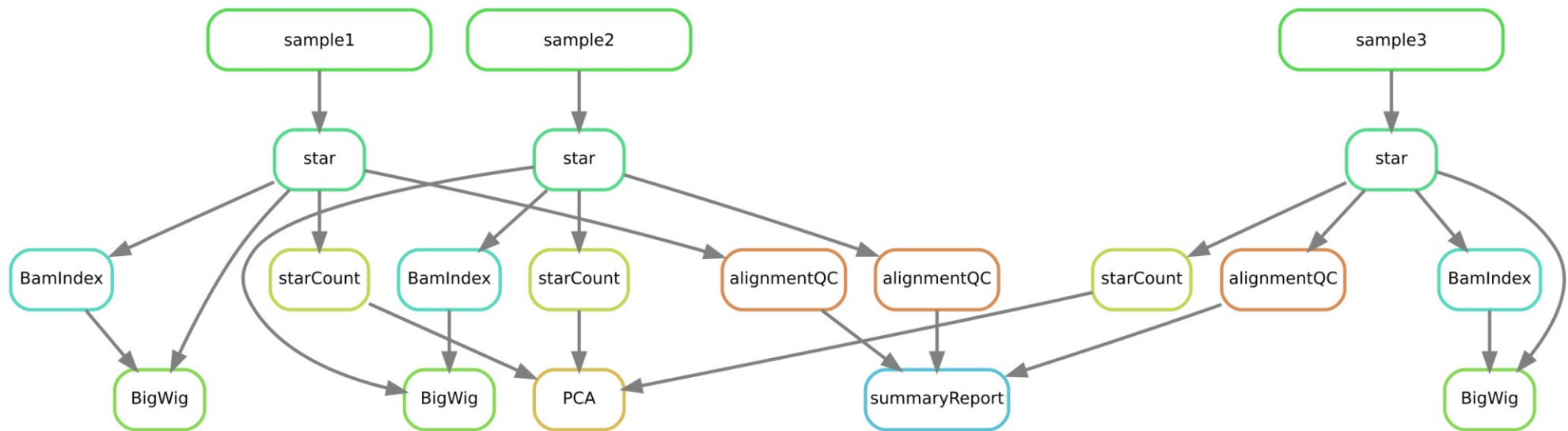


Some tasks can run **in parallel**

Some tasks have to **wait** for one or several others to complete

# Why using a workflow manager?

Simple example, more samples (3)



Some tasks are done **for each sample**

Some **combine** the information from several samples

→ Usually more samples, without workflow manager you may get lost...

	One side						8 V2		
	B1	B2	B3	B4	B5	B6	C1	C2	C3
QC	✓	✓	<del>✓</del> ✓	<del>✓</del> ✓	✓	✓	✓	✓	✓
Napping	✓ <small>dr napping</small>	✓	✓	<del>dr</del> ✓	✓	✓ <sup>2-6</sup>	✓	✓	✓
bw (Feb)	✓	✓	✓	✓	✓	✓ <del>dr</del> ✓	✓	<del>dr</del> <sup>for</sup> <del>dr</del>	✓
Count	✓ <del>dr</del>	✓	✓	✓	✓	✓	✓	✓ <sub>dr</sub>	✓
Edge				<u>dr</u>					
					summary	<u>dr</u>			

- more efficient, faster
- more reproducible
- modular → library of reusable blocks (any language)

- more efficient, faster
- more reproducible
- modular → library of reusable blocks (any language)

# RNA-seq analysis workflow

→ Facilitate primary and secondary analysis of RNA-seq data

Based on a published workflow

Zhang and Jonassen *BMC Bioinformatics* (2020) 21:110  
<https://doi.org/10.1186/s12859-020-3433-x>

BMC Bioinformatics

**SOFTWARE**

**Open Access**

## RASflow: an RNA-Seq analysis workflow with Snakemake

Xiaokang Zhang and Inge Jonassen\* 





# RNA-seq analysis workflow

→ Facilitate primary and secondary analysis of RNA-seq data

Based on a published workflow

Zhang and Jonassen *BMC Bioinformatics* (2020) 21:110  
<https://doi.org/10.1186/s12859-020-3433-x>

BMC Bioinformatics

**SOFTWARE** **Open Access**

## RASflow: an RNA-Seq analysis workflow with Snakemake

Xiaokang Zhang and Inge Jonassen\* 



## Modified

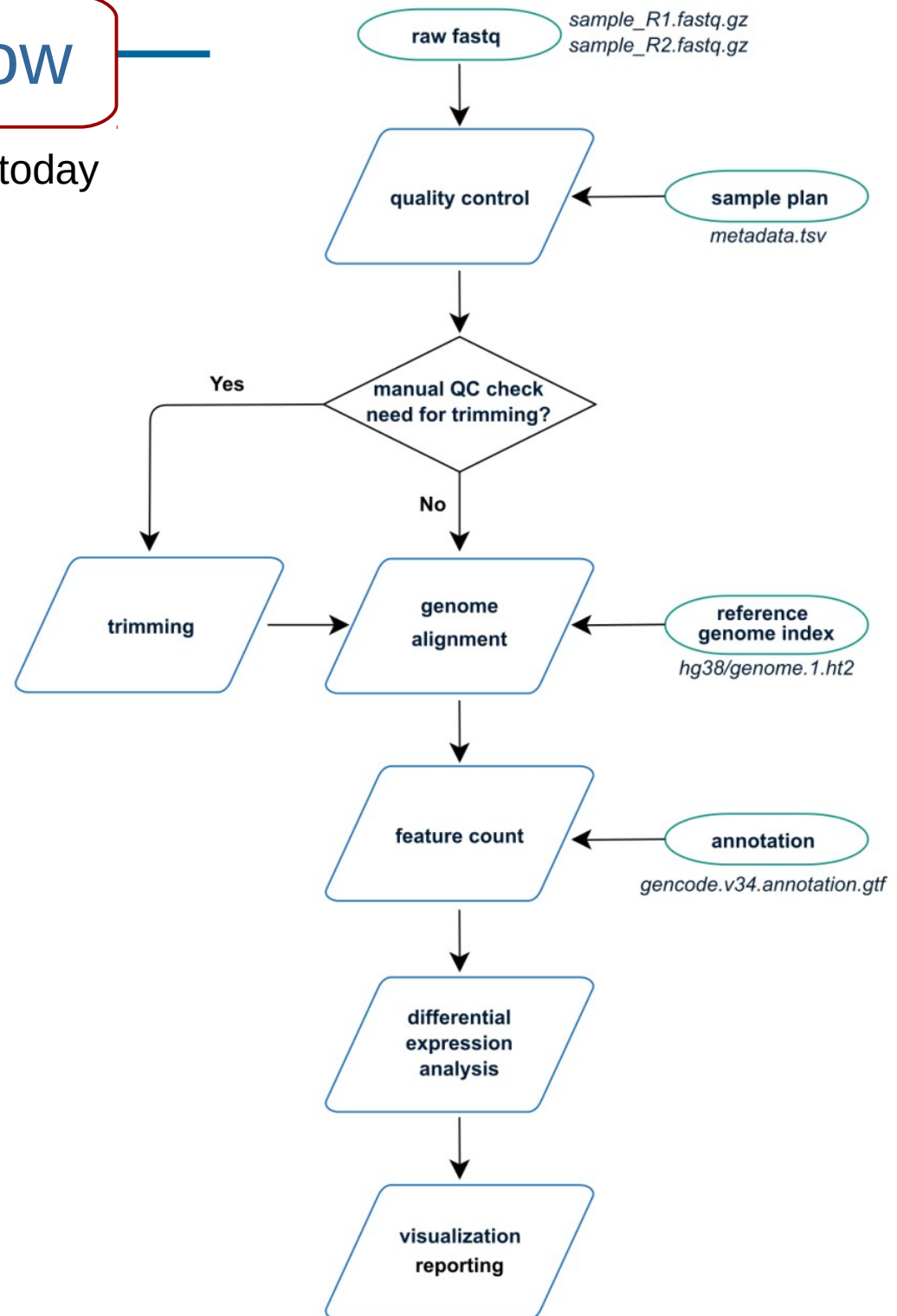
- to run on IFB and RPBS clusters
- to be computationally more efficient
- to include more tools and more parameters

→ Modular, highly flexible

# RNA-seq analysis workflow

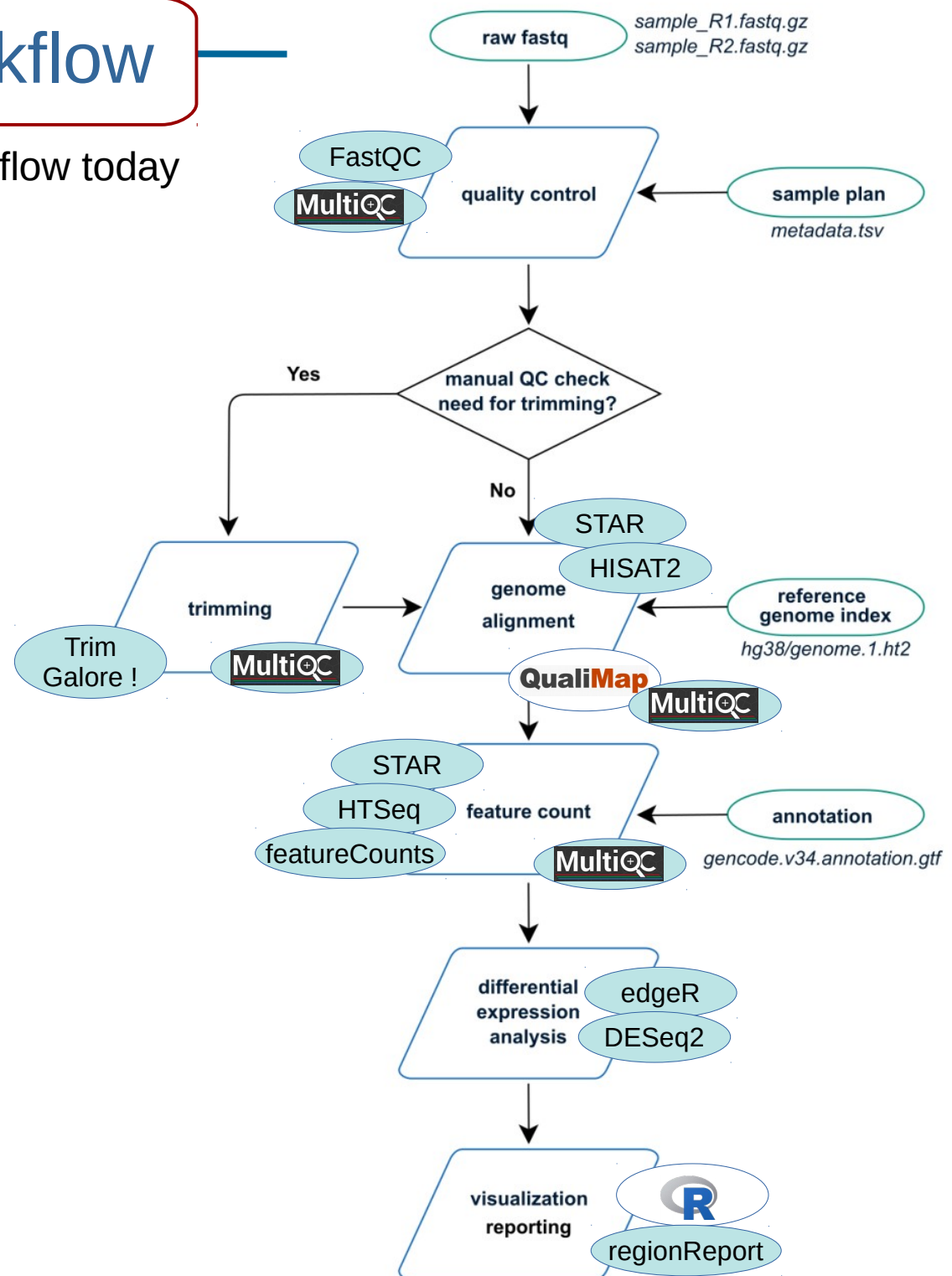


The workflow today



# RNA-seq analysis workflow

The workflow today



# RNA-seq analysis workflow

## Where to compute ?

- IFB core cluster (for now one installation / project)
- RPBS cluster (one installation for everyone)
- personal computer



# RNA-seq analysis workflow

## Where to compute ?

- IFB core cluster (for now one installation / project)
- RPBS cluster (one installation for everyone)
- personal computer



## Prerequisites

- Minimal shell knowledge to copy your files at the right place, navigate between folders...
- Minimal knowledge about the cluster
- Knowledge about the tools (are they adapted to your question?)

→ The platform is here to help you !

**Not needed:** SnakeMake, scripting

**Advanced user:** For now, tools with default parameters

→ add configuration options

# RNA-seq analysis workflow

## How to start your analysis ?

Tutorial and code available on GitHub

The screenshot shows the GitHub repository page for 'parisepigenetics / RASflow\_IFB'. The repository is private and has 3 unwatched items, 0 stars, and 0 forks. The main content area displays a list of files and folders with their commit history. The 'README.md' file is selected, showing its content. The README includes a title 'Tutorial RASflow on IFB core cluster', a maintainer 'Magali Hennion', and a table of contents.

**Files and Folders:**

File/Folder	Commit Message	Time
Tuto_pictures	Update tuto	3 days ago
configs	Update README (STAR usage) - Change name of result directories	last month
scripts	Update README (STAR usage) - Change name of result directories	last month
workflow	Update README (STAR usage) - Change name of result directories	last month
LICENSE	Initial commit	3 months ago
README.md	Update tuto	3 days ago
StarIndex.sh	Update README (STAR usage) - Change name of result directories	last month
Unlock.sh	Make the workflow compatible with RPBS cluster.	2 months ago
Workflow.sh	Update README (STAR usage) - Change name of result directories	last month
cluster.yml	Add STAR	last month
main_cluster.py	Allow the start from count tables	last month

**README.md Content:**

### Tutorial RASflow on IFB core cluster

Maintained by [Magali Hennion](#). Last update : 02/09/2020.

RASflow is a workflow for RNA-seq data analysis originally published by [X. Zhang](#). It has been modified to run effectively on IFB core cluster and to fit our specific needs. Moreover, several tools were added. If you encounter troubles or need additional tools or features, you can create an issue on the [GitHub repository](#), or email directly [Magali](#).

#### Table of content

- Your analysis in a nutshell
- Resources
- Get an account on IFB core cluster and create a project
- Transfer your data
  - FASTQ names
- Connect to IFB core cluster
- RASflow installation and description
- Preparing the run
  - 1. [metadata.tsv](#)
  - 2. [config\\_main.yaml](#)
  - 3. [Workflow.sh](#) [Facultative]
  - 4. [env.yaml](#) [Facultative]
- Running the workflow

**Repository Statistics:**

- About:** Implementation of RASflow on IFB core cluster. License: GPL-3.0.
- Releases:** version 0.3 (Latest) on Jul 28. + 2 releases.
- Packages:** No packages published. [Publish your first package](#).
- Languages:** R 49.5%, Python 27.0%, Shell 16.6%, Java 6.9%.

# RNA-seq analysis workflow

## How to start your analysis ?

Tutorial and code  
available on GitHub

Need for a training session ?

The screenshot shows the GitHub repository page for 'parisepigenetics / RASflow\_IFB'. The repository is private and has 3 unwatched items, 0 stars, and 0 forks. The main content area displays a list of files and folders, including 'Tuto\_pictures', 'configs', 'scripts', 'workflow', 'LICENSE', 'README.md', 'StarIndex.sh', 'Unlock.sh', 'Workflow.sh', 'cluster.yml', and 'main\_cluster.py'. The 'README.md' file is selected, showing its content. The README includes a title 'Tutorial RASflow on IFB core cluster', a maintainer 'Magali Hennion', and a table of contents. The table of contents lists sections such as 'Your analysis in a nutshell', 'Resources', 'Get an account on IFB core cluster and create a project', 'Transfer your data', 'Connect to IFB core cluster', 'RASflow installation and description', 'Preparing the run', and 'Running the workflow'.

parisepigenetics / RASflow\_IFB Private

Unwatch 3 Star 0 Fork 0

Code Issues Pull requests Actions Projects Wiki Security Insights Settings

master 1 branch 3 tags

Go to file Add file Code

Magali Hennion Update tuto 175498e 3 days ago 44 commits

File	Commit	Time
Tuto_pictures	Update tuto	3 days ago
configs	Update README (STAR usage) - Change name of result directories	last month
scripts	Update README (STAR usage) - Change name of result directories	last month
workflow	Update README (STAR usage) - Change name of result directories	last month
LICENSE	Initial commit	3 months ago
README.md	Update tuto	3 days ago
StarIndex.sh	Update README (STAR usage) - Change name of result directories	last month
Unlock.sh	Make the workflow compatible with RPBS cluster.	2 months ago
Workflow.sh	Update README (STAR usage) - Change name of result directories	last month
cluster.yml	Add STAR	last month
main_cluster.py	Allow the start from count tables	last month

README.md

### Tutorial RASflow on IFB core cluster

Maintained by [Magali Hennion](#). Last update : 02/09/2020.

RASflow is a workflow for RNA-seq data analysis originally published by [X. Zhang](#). It has been modified to run effectively on IFB core cluster and to fit our specific needs. Moreover, several tools were added. If you encounter troubles or need additional tools or features, you can create an issue on the [GitHub repository](#), or email directly [Magali](#).

#### Table of content

- Your analysis in a nutshell
- Resources
- Get an account on IFB core cluster and create a project
- Transfer your data
  - FASTQ names
- Connect to IFB core cluster
- RASflow installation and description
- Preparing the run
  - 1. [metadata.tsv](#)
  - 2. [config\\_main.yaml](#)
  - 3. [Workflow.sh](#) [Facultative]
  - 4. [env.yaml](#) [Facultative]
- Running the workflow

About

Implementation of RASflow on IFB core cluster

Readme

GPL-3.0 License

Releases 3

version 0.3 Latest on Jul 28

+ 2 releases

Packages

No packages published

Publish your first package

Languages

R 49.5% Python 27.0% Shell 16.6% Java 6.9%

# RNA-seq analysis workflow

## Your analysis in a nutshell

- Get an [account](#) on IFB core cluster and create a project
- [Transfer your data](#) to the cluster
- [Clone](#) RASflow\_IFB repository
- [Modify](#) `metadata.tsv` and `config_main.yaml`
- Run the [workflow](#) typing `sbatch Workflow.sh`
- Look at the results

### Two files to modify

- sample plan (`metadata.tsv`)

sample	group	subject
T27	J0_WT	1
T28	J0_WT	2
T29	J0_WT	3
T30	J0_KO	1
T31	J0_KO	2
T32	J0_KO	3
T33	J10_WT	1
T34	J10_WT	2
T35	J10_WT	3
T36	J10_KO	1
T37	J10_KO	2
T38	J10_KO	3



# RNA-seq analysis workflow

## Two files to modify

- sample plan
- configuration file

```
# Please check the parameters, and adjust them according to your circumstance

# Project name
PROJECT: EXAMPLE

# ===== Control of the workflow =====

## Do you need to do quality control?
QC: yes # "yes" or "no". If set to "yes", the workflow will stop after the QC to let you decide whether you want to trim

## Do you need to do trimming?
TRIMMED: yes # "yes" or "no"?

## Which mapping reference do you want to use? Genome or transcriptome?
REFERENCE: genome # "genome" or "transcriptome", I haven't implemented transcriptome yet.

## Do you want to do Differential Expression Analysis (DEA)?
DEA: yes # "yes" or "no"

## Do you want to visualize the results of DEA?
VISUALIZE: yes # "yes" or "no"
```

# RNA-seq analysis workflow

## Two files to modify

- sample plan
- configuration file

```
# ===== Shared parameters for some or all of the sub-workflows =====

## key file if the data is stored remotely, otherwise leave it empty
KEY:

## the path to fastq files
READSPATH: /shared/projects/YourProjectName/Raw_fastq

## the meta file describing the experiment settings
METAFILE: /shared/projects/YourProjectName/RASflow_IFB/configs/metadata.tsv

## paths for intermediate and final results
BIGDATAPATH: /shared/projects/YourProjectName/RASflow_IFB/data # for big files
RESULTPATH: /shared/projects/YourProjectName/RASflow_IFB/results

## is the sequencing paired-end or single-end?
END: pair # "pair" or "single"

## number of cores you want to allocate to this workflow
NCORE: 30 # Use command "getconf _NPROCESSORS_ONLN" to check the number of cores/CPU on your machine
```

# RNA-seq analysis workflow

## Two files to modify

- sample plan
- configuration file

```
# ===== Configuration for alignment to genome and feature count =====

## aligner
ALIGNER: hisat2 # "STAR" or "hisat2"

## genome and annotation files
INDEXPATH: /shared/bank/homo_sapiens/hg38/hisat2 # or index/STAR # folder containing index files
INDEXBASE: genome # for hisat2, base of the name of the index files (ie genome.1.ht2)
ANNOTATION: /shared/projects/YourProjectName/RASflow_IFB/gtf/gencode.v34.annotation.gtf # GTF file

## bigwig option
BWSTRANDED: both # "no": bw merging forward and reverse reads, "yes": get 2 bw files, one forward and one reverse

## tool for feature count
COUNTER: featureCounts # "featureCounts" or "htseq-count" or "STARcount" (only with STAR aligner, -

## counting options
ATTRIBUTE: gene_id # the attribute used in annotation file. It's usually "gene_id", but double check
STRAND: "reverse" # "no", "yes", "reverse". For ht-seq counts: For stranded=no, a read is considered
FEATURE: transcript # "exon" or "transcript"
```

[...]

→ **start the analysis**

# RNA-seq analysis workflow

## Running time

Data from Madeleine, 12 samples (pair-end), 140 GB in total, 4-8 GB / fastq files)

- Data transfer : ~ 1h (from the lab)
- Creating Conda environment : 15-30 min
- QC : 40-60 min
- trimming: 2-3 h
- mapping: 5-7 h
- DEA: 5-10 min
- visualization : 5-10 min

# RNA-seq analysis workflow

## Running time

Data from Madeleine, 12 samples (pair-end), 140 GB in total, 4-8 GB / fastq files)

- Data transfer : ~ 1h (from the lab)

- Creating Conda environment : 15-30 min

- QC : 40-60 min

- trimming: 2-3 h

- mapping: 5-7 h

- DEA: 5-10 min

- visualization : 5-10 min

→ TOTAL : ~ 12h computing + human time

Need  
for a  
human  
brain

# RNA-seq analysis workflow

## Results

### Quality control before and after trimming

[file:///home/mag/Documents/Communication/20200914\\_LabSeminar/report\\_quality\\_control.html](file:///home/mag/Documents/Communication/20200914_LabSeminar/report_quality_control.html)

[file:///home/mag/Documents/Communication/20200914\\_LabSeminar/report\\_quality\\_control\\_after\\_trimming.html](file:///home/mag/Documents/Communication/20200914_LabSeminar/report_quality_control_after_trimming.html)

# RNA-seq analysis workflow

## Results

### Quality control before and after trimming

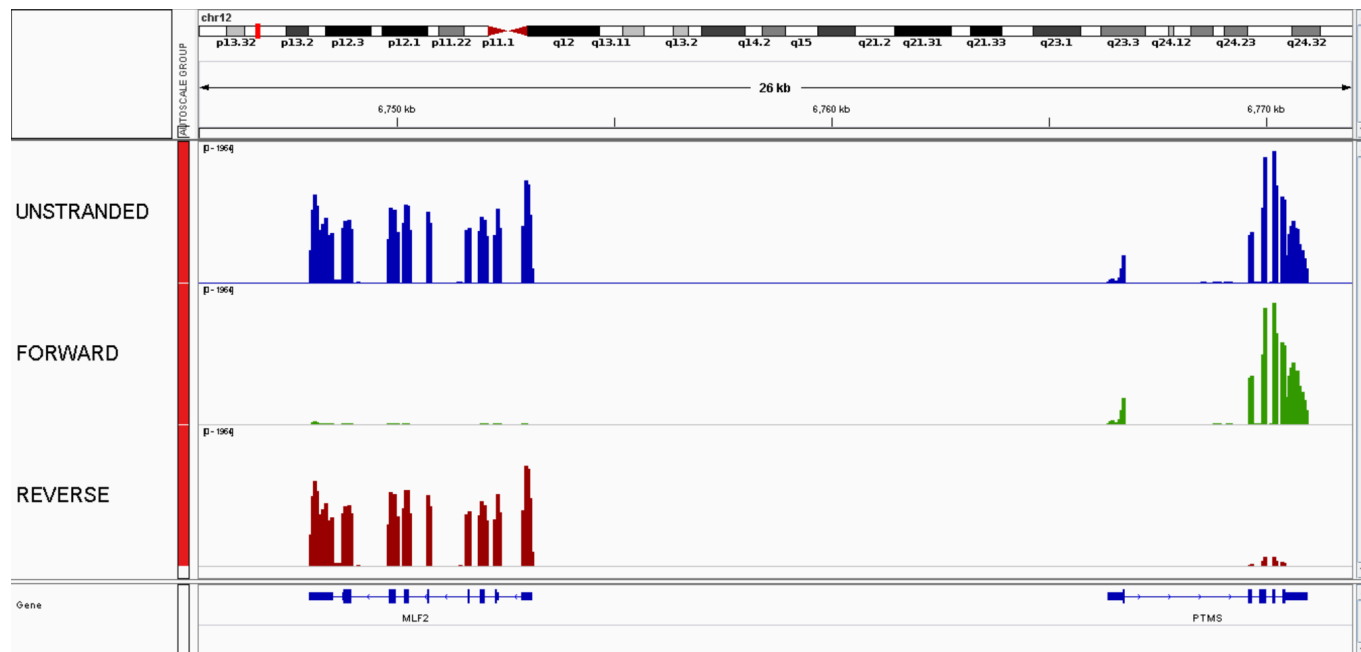
[file:///home/mag/Documents/Communication/20200914\\_LabSeminar/report\\_quality\\_control.html](file:///home/mag/Documents/Communication/20200914_LabSeminar/report_quality_control.html)

[file:///home/mag/Documents/Communication/20200914\\_LabSeminar/report\\_quality\\_control\\_after\\_trimming.html](file:///home/mag/Documents/Communication/20200914_LabSeminar/report_quality_control_after_trimming.html)

### Quality control after mapping and counting

[file:///home/mag/DataMadeleine/TIMING/hisat2/report\\_align\\_count\\_featureCounts.html](file:///home/mag/DataMadeleine/TIMING/hisat2/report_align_count_featureCounts.html)

### BigWig files to look at the data on genome browser



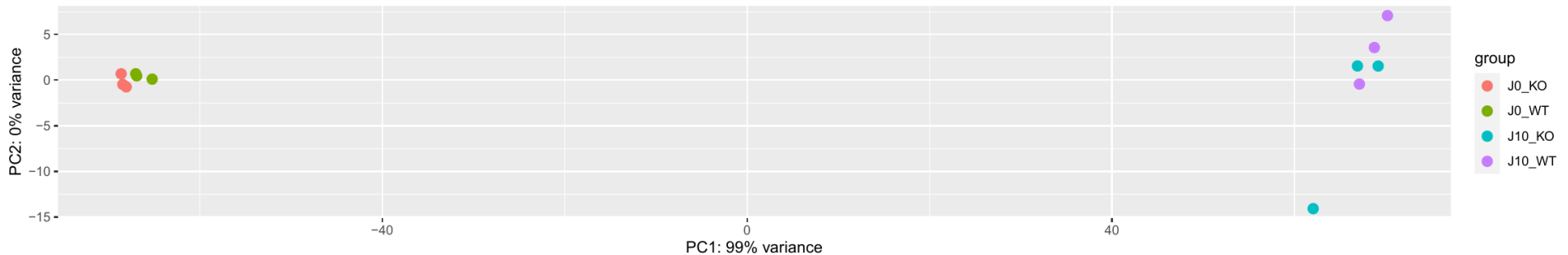
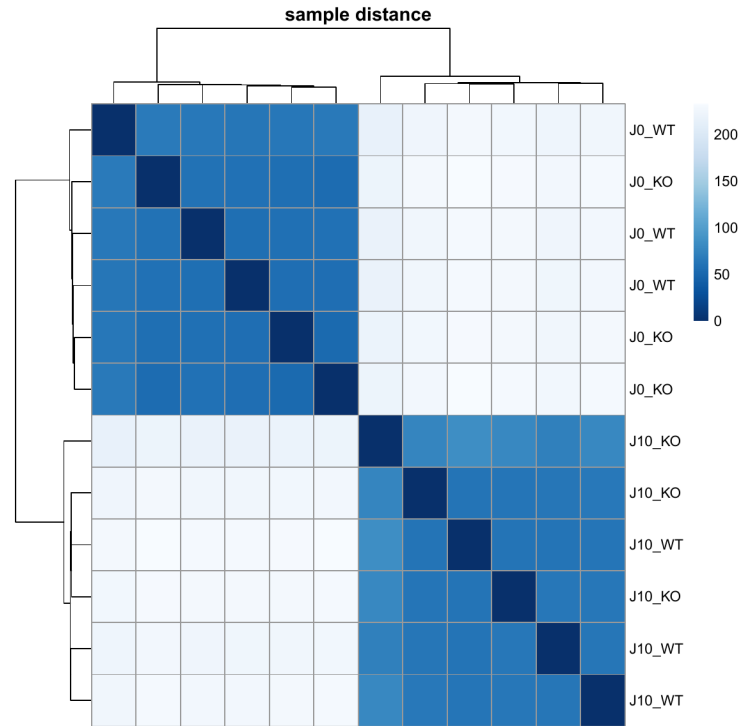
# RNA-seq analysis workflow

## Results

### Raw and normalized count files

D197-D192T35_count.tsv	
ENSG00000223972.5	1
ENSG00000227232.5	263
ENSG00000278267.1	0
ENSG00000243485.5	1
ENSG00000284332.1	0
ENSG00000237613.2	1
ENSG00000268020.3	0
ENSG00000240361.2	0
ENSG00000186092.6	0
ENSG00000238009.6	26
ENSG00000239945.1	0
ENSG00000233750.3	0
ENSG00000268903.1	19
ENSG00000269981.1	13
ENSG00000239906.1	3
ENSG00000241860.7	15
ENSG00000222623.1	0
ENSG00000241599.1	0
ENSG00000279928.2	4
ENSG00000279457.4	517
ENSG00000273874.1	0
ENSG00000228463.10	11
ENSG00000286448.1	0
ENSG00000236679.2	0
ENSG00000236601.2	0
ENSG00000237094.12	75

### Heatmap and PCA of all the samples





# RNA-seq analysis workflow

## Results

### Differential expression analysis (between pairs of conditions)

Tables (all genes / significantly different)

Report

dea_J0_WT_J10_WT.tsv									
baseMean	log2FoldChange	lfcSE	stat	pvalue	padj				
ENSG00000138722.10	28643.5576775341	12.8878559048439	0.334254252259555	38.5570439799108	0	0			
ENSG00000150637.9	44495.3489697712	12.1138628938676	0.225254162861642	53.7786416018794	0	0			
ENSG00000248131.6	23035.2235024924	-11.5102220691977	0.214583147815272	-53.6399162114375	0	0			
ENSG00000236673.5	9410.62778439526	-11.1702993224953	0.284876900393565	-39.2109690433419	0	0			
ENSG0000010671.16	17434.4042351806	11.1368058134479	0.247095134871919	45.0709230645948	0	0			
ENSG00000274090.1	12673.5010973939	-11.0431466139289	0.243020448209864	-45.4412239598557	0	0			
ENSG00000128683.14	38392.4960153405	10.6771649471065	0.165881192381676	64.3663383039795	0	0			
ENSG00000261371.6	21011.2196060373	10.5882903827317	0.202428901612002	52.3062186200387	0	0			
ENSG00000185559.16	9291.39445851381	10.5672720975195	0.264977072858931	39.8799487952127	0	0			
ENSG00000151702.17	27888.8642370209	9.90260964734258	0.151987244928666	65.1542150921303	0	0			
ENSG00000111348.9	12188.4835524031	9.72369598445457	0.186453617244828	52.150750026408	0	0			
ENSG00000077420.16	6537.36530736273	9.72092375413297	0.241392363666131	40.2702206751576	0	0			
ENSG00000165702.14	9343.36263764007	9.59803131814038	0.227418626840593	42.2042444433016	0	0			
ENSG00000162511.8	5696.04146023551	9.51650005466677	0.249728319897167	38.1074123214599	0	0			
ENSG00000255929.5	7598.70343369437	-9.47019261095421	0.19136293672142	-49.4881233179476	0	0			
ENSG00000164512.18	6990.32718933163	9.40628158397668	0.212743801625526	44.2141275661404	0	0			
ENSG00000163737.4	41625.3614482218	9.3987254652196	0.16974012515012	55.3712650848306	0	0			
ENSG00000064201.16	10395.1831045548	9.38771319937487	0.191501351377414	49.0216551050514	0	0			
ENSG00000158578.21	21622.2181893878	9.30066251795876	0.150229090820593	61.9098635767279	0	0			
ENSG00000164035.10	6915.59463518514	9.20058963397617	0.205829692780933	44.7000114981879	0	0			
ENSG00000254277.1	12497.463812679	-9.02174609448843	0.140908284960578	-64.0256610674984	0	0			
ENSG00000182866.17	3254.84007527242	-8.82544806879715	0.222680250843156	-39.6328279467105	0	0			
ENSG00000104903.5	5556.34779269385	8.75726129253258	0.200868468802165	43.5969933198307	0	0			
ENSG00000152208.13	169397.89006299	-8.65256319895797	0.12072759446843	-71.6701366995317	0	0			
ENSG00000268555.2	3961.66735219303	8.312255086070514	0.196847711819128	42.2268096686986	0	0			
ENSG00000095303.17	19954.8652152089	8.281811632252	0.134548170511187	61.5527628565071	0	0			
ENSG00000134138.20	4943.42492987928	8.21668929248445	0.188177877289831	43.6644807074167	0	0			
ENSG00000226792.7	5929.13871166827	-8.19648712538188	0.14617436684561	-56.0733547355745	0	0			
ENSG00000236780.7	13522.677227173	-8.14693091270317	0.120096819744012	-67.8363584486953	0	0			
ENSG00000244342.5	4412.35248366687	-8.10918207356467	0.161825975718805	-50.1105093761678	0	0			
ENSG00000133083.14	5994.21884434801	-8.006668768897	0.141959393523884	-56.4011198565026	0	0			
ENSG00000149781.12	25395.5559785894	7.99738481605514	0.132915719822231	60.1688410276174	0	0			
ENSG00000118308.15	4085.4630243289	7.96054054065835	0.181538060028329	43.8505321661811	0	0			
ENSG00000164076.17	2044.74195604111	-7.9085982226867	0.209760276503151	-37.7030310720815	0	0			
ENSG00000166105.16	2718.07816642691	-7.90155905419819	0.190344071656288	-41.5119787311599	0	0			
ENSG00000172578.12	5552.74698183465	7.87262797383369	0.161041248685017	48.8857857109135	0	0			
ENSG00000165092.13	10490.0915575169	7.86110735462206	0.124118978589746	63.3352565734979	0	0			

file:///home/mag/DataMadeleine/TIMING/hisat2/dea  
featureCounts/Report\_DESeq2/J0\_WT\_J0\_KO/D  
ESeq2Exploration.html

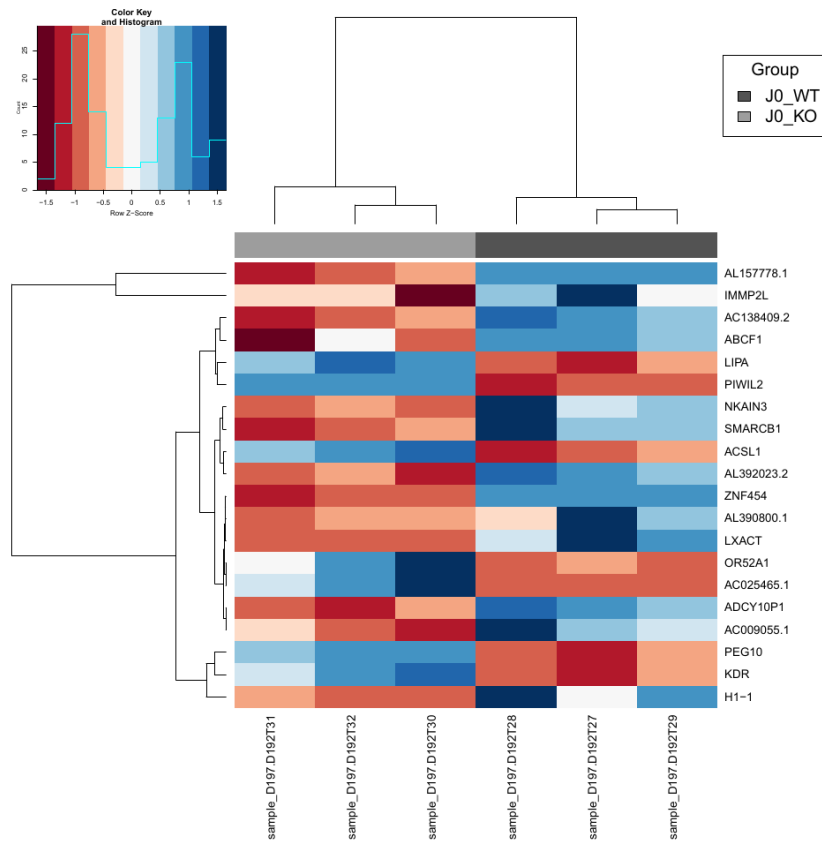
 regionReport  
(with code)

# RNA-seq analysis workflow

## Results

### Differential expression analysis (between pairs of conditions)

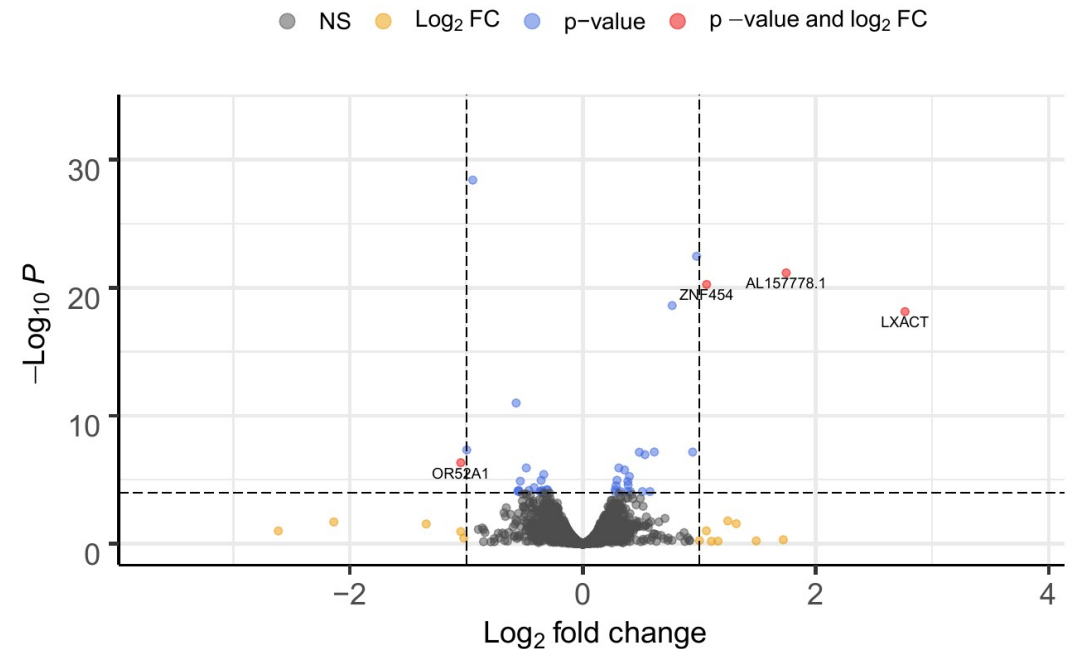
#### Heatmap Top 20 genes



#### Volcano plot

##### DESeq2

J0\_WT vs J0\_KO



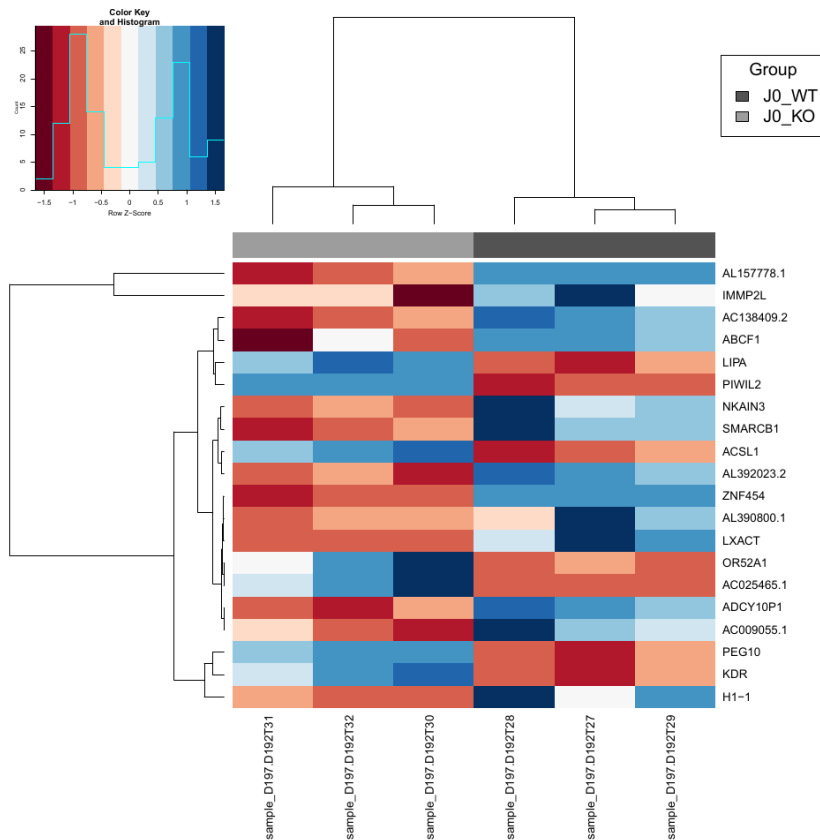
Total = 13583 variables

# RNA-seq analysis workflow

## Results

### Differential expression analysis (between pairs of conditions)

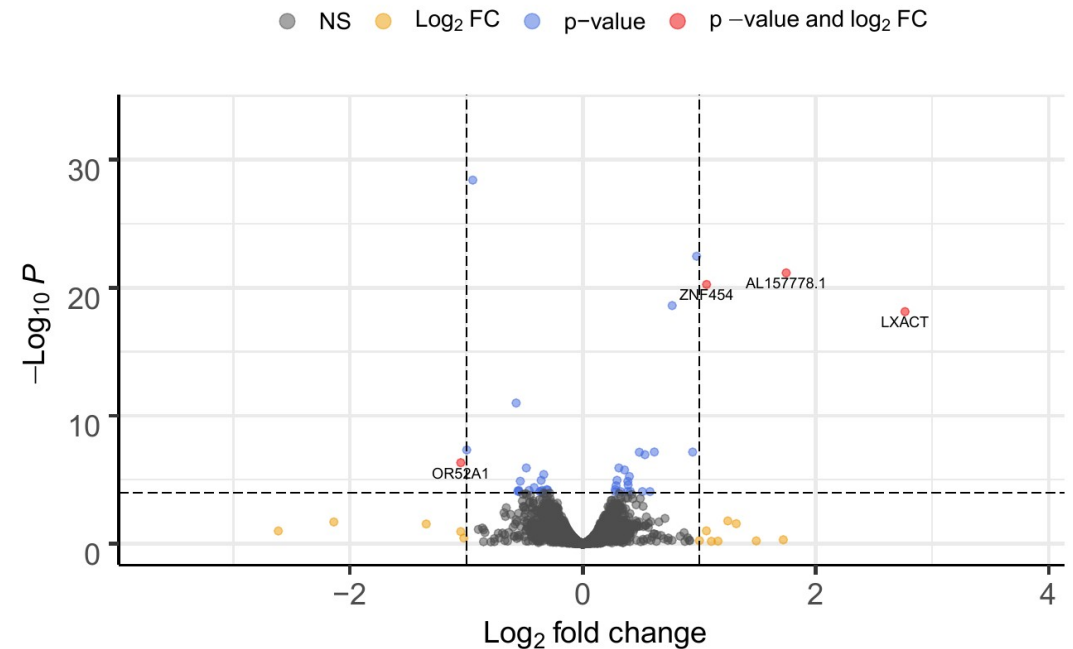
#### Heatmap Top 20 genes



#### Volcano plot

##### DESeq2

J0\_WT vs J0\_KO



Total = 13583 variables



Your analysis is not over ! This is just a starting point...

# RNA-seq analysis workflow

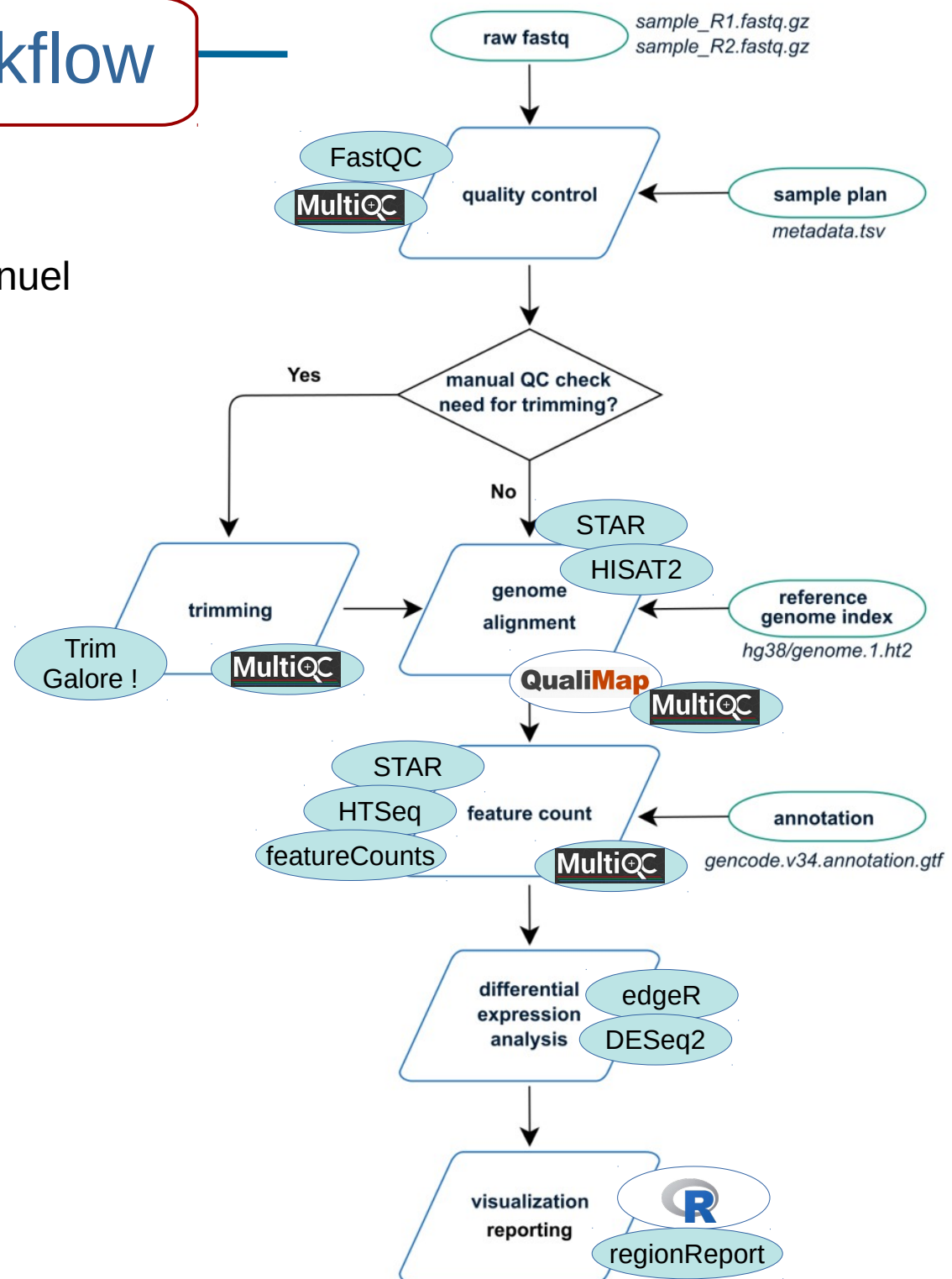
beta-testers : Olivier, Madeleine, Emmanuel

## To be added

- Salmon (pseudo mapping)
- GSEA / GO / KEGG
- export .RData objects
- ask what you need !

## To be adapted to

- genomic repeats
- transcript reconstruction
- ChIP-seq
- ATAC-seq
- WGBS
- ask what you need !



# RNA-seq analysis workflow

## Reproducibility

- Conda environment → define the tools and their version



→ results identical on IFB and RPBS clusters

```
name: rasflow_IFB
channels:
  - conda-forge
  - bioconda
  - r
  - defaults
dependencies:
  - snakemake=5.14.0
  - graphviz=2.42.3
# command tool installs
  - R=4.0
  - python=3.7.6
# r channel installs
  - r-yaml=2.2.1
  - r-statmod=1.4.34
  - r-ggplots=3.0.3
  - r-magick=2.3
  - r-dt=0.13
  - r-sessioninfo=1.1.1
# conda-forge channel installs
  - r-heatmap.plus=1.3
  - r-readr=1.3.1
  - r-hash=3.0.1
  - r-pheatmap=1.0.12
  - r-rcolorbrewer=1.1_2
  - imagemagick=7.0.10
# bioconda channel installs
  - fastqc=0.11.9
  - trim-galore=0.6.5
  - multiqc=1.9
  - salmon=1.2.1
  - hisat2=2.2.0
  - samtools=1.10
  - subread=2.0.1 # featureCounts included
  - htseq=0.12.4 # htseq-count included
  - bioconductor-edger=3.30.0
  - bioconductor-deseq2=1.28.0
  - qualimap=2.2.2a
  - bioconductor-mygene=1.24.0
  - bioconductor-tximport=1.16.0
  - bioconductor-enhancedvolcano=1.6.0
  - bioconductor-biomart=2.44.0
  - deeptools=3.4.3
  - bioconductor-regionreport=1.22.0
  - star=2.7.5a
```



# RNA-seq analysis workflow

## Reproducibility

- reporting

\* logs

\* R libraries (regionReport)

\* MultiQC

### 11 Reproducibility

The input for this report was generated with edgeR (Robinson, McCarthy, and Smyth, 2010; McCarthy, Chen, and Smyth, 2012; Chen, Lun, and Smyth, 2014) and the resulting features were called significantly differentially expressed if their BH adjusted p-values were less than  $\alpha = 0.1$ . This report was generated in path /home/mag/DataMadeleine using the following call to edgeReport():

```
## edgeReport(dge = y, object = lrt, project = "edgeR-example",  
##   intgroup = "group", outdir = "edgeReport-example")
```

Date the report was generated.

```
## [1] "2020-06-22 11:36:02 CEST"
```

Wallclock time spent generating the report.

```
## Time difference of 14.73 secs
```

R session information.

```
## - Session info -  
##  
## setting value  
## version R version 3.6.3 (2020-02-29)  
## os Linux Mint 19.1  
## system x86_64, linux-gnu  
## ui X11  
## language en_US  
## collate en_US.UTF-8  
## ctype en_US.UTF-8  
## tz Europe/Paris  
## date 2020-06-22  
##  
## - Packages -  
##  
## package * version date lib source  
## acepack 1.4.1 2016-10-29 [1] CRAN (R 3.6.3)  
## annotate 1.64.0 2019-10-29 [1] Bioconductor  
## AnnotationDbi 1.48.0 2019-10-29 [1] Bioconductor  
## askpass 1.1 2019-01-13 [1] CRAN (R 3.6.1)  
## assertthat 0.2.1 2019-03-21 [1] CRAN (R 3.6.1)  
## backports 1.1.5 2019-10-02 [1] CRAN (R 3.6.1)  
## base64enc 0.1-3 2015-07-28 [1] CRAN (R 3.6.1)  
## bibtex 0.4.2.2 2020-01-02 [1] CRAN (R 3.6.3)  
## Biobase * 2.46.0 2019-10-29 [1] Bioconductor  
## BiocFileCache 1.10.2 2019-11-08 [1] Bioconductor  
## BiocGenerics * 0.32.0 2019-10-29 [1] Bioconductor  
## BiocManager 1.30.10 2019-11-16 [1] CRAN (R 3.6.3)  
## BiocParallel * 1.20.1 2019-12-21 [1] Bioconductor  
## BiocStyle * 2.14.4 2020-01-09 [1] Bioconductor  
## biomaRt 2.42.0 2019-10-29 [1] Bioconductor  
## Biostrings 2.54.0 2019-10-29 [1] Bioconductor  
## bit 1.1-15.2 2020-02-10 [1] CRAN (R 3.6.3)  
## bit64 0.9-7 2017-05-08 [1] CRAN (R 3.6.3)  
## bitops 1.0-6 2013-08-17 [1] CRAN (R 3.6.3)  
## blob 1.2.1 2020-01-20 [1] CRAN (R 3.6.3)  
## bookdown 0.19 2020-05-15 [1] CRAN (R 3.6.3)  
## BSgenome 1.54.0 2019-10-29 [1] Bioconductor  
## bumphunter 1.28.0 2019-10-29 [1] Bioconductor  
## checkmate 2.0.0 2020-02-06 [1] CRAN (R 3.6.3)  
## cli 2.0.2 2020-02-28 [1] CRAN (R 3.6.1)  
## cluster 2.1.0 2019-06-19 [1] CRAN (R 3.6.3)  
## codetools 0.2-16 2018-12-24 [1] CRAN (R 3.6.3)  
## colorspace 1.4-1 2019-03-18 [1] CRAN (R 3.6.1)  
## crayon 1.3.4 2017-09-16 [1] CRAN (R 3.6.1)  
## crosstalk 1.0.0 2016-12-21 [1] CRAN (R 3.6.1)
```

```
## crosstalk 1.0.0 2016-12-21 [1] CRAN (R 3.6.1)  
## curl 4.3 2019-12-02 [1] CRAN (R 3.6.1)  
## data.table 1.12.8 2019-12-09 [1] CRAN (R 3.6.3)  
## DBI 1.1.0 2019-12-15 [1] CRAN (R 3.6.1)  
## dbplyr 1.4.2 2019-06-17 [1] CRAN (R 3.6.1)  
## DEFormats 1.14.0 2019-10-29 [1] Bioconductor  
## DelayedArray * 0.12.2 2020-01-06 [1] Bioconductor  
## derfinder 1.20.0 2019-10-29 [1] Bioconductor  
## derfinderHelper 1.20.0 2019-10-29 [1] Bioconductor  
## DESeq2 * 1.26.0 2019-10-29 [1] Bioconductor  
## digest 0.6.25 2020-02-23 [1] CRAN (R 3.6.1)  
## doRNG 1.8.2 2020-01-27 [1] CRAN (R 3.6.3)  
## dplyr 0.8.5 2020-03-07 [1] CRAN (R 3.6.3)  
## DT * 0.12 2020-02-05 [1] CRAN (R 3.6.1)  
## edgeR * 3.28.1 2020-02-26 [1] Bioconductor  
## evaluate 0.14 2019-05-28 [1] CRAN (R 3.6.1)  
## fansi 0.4.1 2020-01-08 [1] CRAN (R 3.6.1)  
## farver 2.0.3 2020-01-16 [1] CRAN (R 3.6.1)  
## fastmap 1.0.1 2019-10-08 [1] CRAN (R 3.6.1)  
## foreach 1.5.0 2020-03-30 [1] CRAN (R 3.6.3)  
## foreign 0.8-76 2020-03-03 [1] CRAN (R 3.6.3)  
## Formula 1.2-3 2018-05-03 [1] CRAN (R 3.6.3)  
## genefilter 1.68.0 2019-10-29 [1] Bioconductor  
## geneplotter 1.64.0 2019-10-29 [1] Bioconductor  
## GenomeInfoDb * 1.22.0 2019-10-29 [1] Bioconductor  
## GenomeInfoDbData 1.2.2 2020-03-10 [1] Bioconductor  
## GenomicAlignments 1.22.1 2019-11-12 [1] Bioconductor  
## GenomicFeatures 1.38.2 2020-02-15 [1] Bioconductor  
## GenomicFiles 1.22.0 2019-10-29 [1] Bioconductor  
## GenomicRanges * 1.38.0 2019-10-29 [1] Bioconductor  
## ggplot2 * 3.3.0 2020-03-05 [1] CRAN (R 3.6.3)  
## glue 1.3.1 2019-03-12 [1] CRAN (R 3.6.1)  
## gridExtra 2.3 2017-09-09 [1] CRAN (R 3.6.3)  
## gtable 0.3.0 2019-03-25 [1] CRAN (R 3.6.1)  
## highr 0.8 2019-03-20 [1] CRAN (R 3.6.1)  
## Hmisc 4.3-1 2020-02-07 [1] CRAN (R 3.6.3)  
## hms 0.5.3 2020-01-08 [1] CRAN (R 3.6.1)  
## htmlTable 1.13.3 2019-12-04 [1] CRAN (R 3.6.3)  
## htmltools 0.4.0 2019-10-04 [1] CRAN (R 3.6.1)  
## htmlwidgets 1.5.1 2019-10-08 [1] CRAN (R 3.6.1)  
## httpuv 1.5.2 2019-09-11 [1] CRAN (R 3.6.1)  
## httr 1.4.1 2019-08-05 [1] CRAN (R 3.6.1)  
## IRanges * 2.20.2 2020-01-13 [1] Bioconductor  
## IRdisplay 0.7.0 2018-11-29 [1] CRAN (R 3.6.1)  
## IRkernel 1.1 2020-03-03 [1] Github (IRkernel/IRkernel)  
## iterators 1.0.12 2019-07-26 [1] CRAN (R 3.6.3)  
## jpeg 0.1-8.1 2019-10-24 [1] CRAN (R 3.6.3)  
## jsonlite 1.6.1 2020-02-02 [1] CRAN (R 3.6.1)  
## knitr 1.0.10 2019-09-15 [1] CRAN (R 3.6.3)  
## knitr * 1.28 2020-02-06 [1] CRAN (R 3.6.3)  
## knitrBootstrap 1.0.2 2018-05-24 [1] CRAN (R 3.6.3)  
## labeling 0.3 2014-08-23 [1] CRAN (R 3.6.1)  
## later 1.0.0 2019-10-04 [1] CRAN (R 3.6.1)  
## lattice 0.20-40 2020-02-19 [1] CRAN (R 3.6.1)  
## latticeExtra 0.6-29 2019-12-19 [1] CRAN (R 3.6.3)  
## lifecycle 0.2.0 2020-03-06 [1] CRAN (R 3.6.3)  
## limma * 3.42.2 2020-02-03 [1] Bioconductor  
## locfit 1.5-9.1 2013-04-20 [1] CRAN (R 3.6.3)  
## lubridate 1.7.4 2018-04-11 [1] CRAN (R 3.6.1)  
## magick 2.3 2020-01-24 [1] CRAN (R 3.6.3)  
## magrittr 1.5 2014-11-22 [1] CRAN (R 3.6.1)  
## markdown 1.1 2019-08-07 [1] CRAN (R 3.6.1)  
## Matrix 1.2-18 2019-11-27 [1] CRAN (R 3.6.1)  
## matrixStats * 0.55.0 2019-09-07 [1] CRAN (R 3.6.3)  
## memoise 1.1.0 2017-04-21 [1] CRAN (R 3.6.1)  
## mime 0.9 2020-02-04 [1] CRAN (R 3.6.1)  
## munsell 0.5.0 2018-06-12 [1] CRAN (R 3.6.1)  
## nnet 7.3-13 2020-02-25 [1] CRAN (R 3.6.3)  
## openssl 1.4.1 2019-07-18 [1] CRAN (R 3.6.1)  
## pbdZMQ 0.3-3 2018-05-05 [1] CRAN (R 3.6.1)  
## pheatmap * 1.0.12 2019-01-04 [1] CRAN (R 3.6.3)  
## pillar 1.4.3 2019-12-20 [1] CRAN (R 3.6.1)  
## pkgconfiq 2.0.3 2019-09-22 [1] CRAN (R 3.6.1)
```

# RNA-seq analysis workflow

## Reproducibility

- reporting

\* logs

\* R libraries (regionReport)

\* MultiQC

Any volunteer to try it?

### 11 Reproducibility

The input for this report was generated with edgeR (Robinson, McCarthy, and Smyth, 2010; McCarthy, Chen, and Smyth, 2012; Chen, Lun, and Smyth, 2014) and the resulting features were called significantly differentially expressed if their BH adjusted p-values were less than  $\alpha = 0.1$ . This report was generated in path /home/mag/DataMadeleine using the following call to edgeReport():

```
## edgeReport(dge = y, object = lrt, project = "edgeR-example",  
##   intgroup = "group", outdir = "edgeReport-example")
```

Date the report was generated.

```
## [1] "2020-06-22 11:36:02 CEST"
```

Wallclock time spent generating the report.

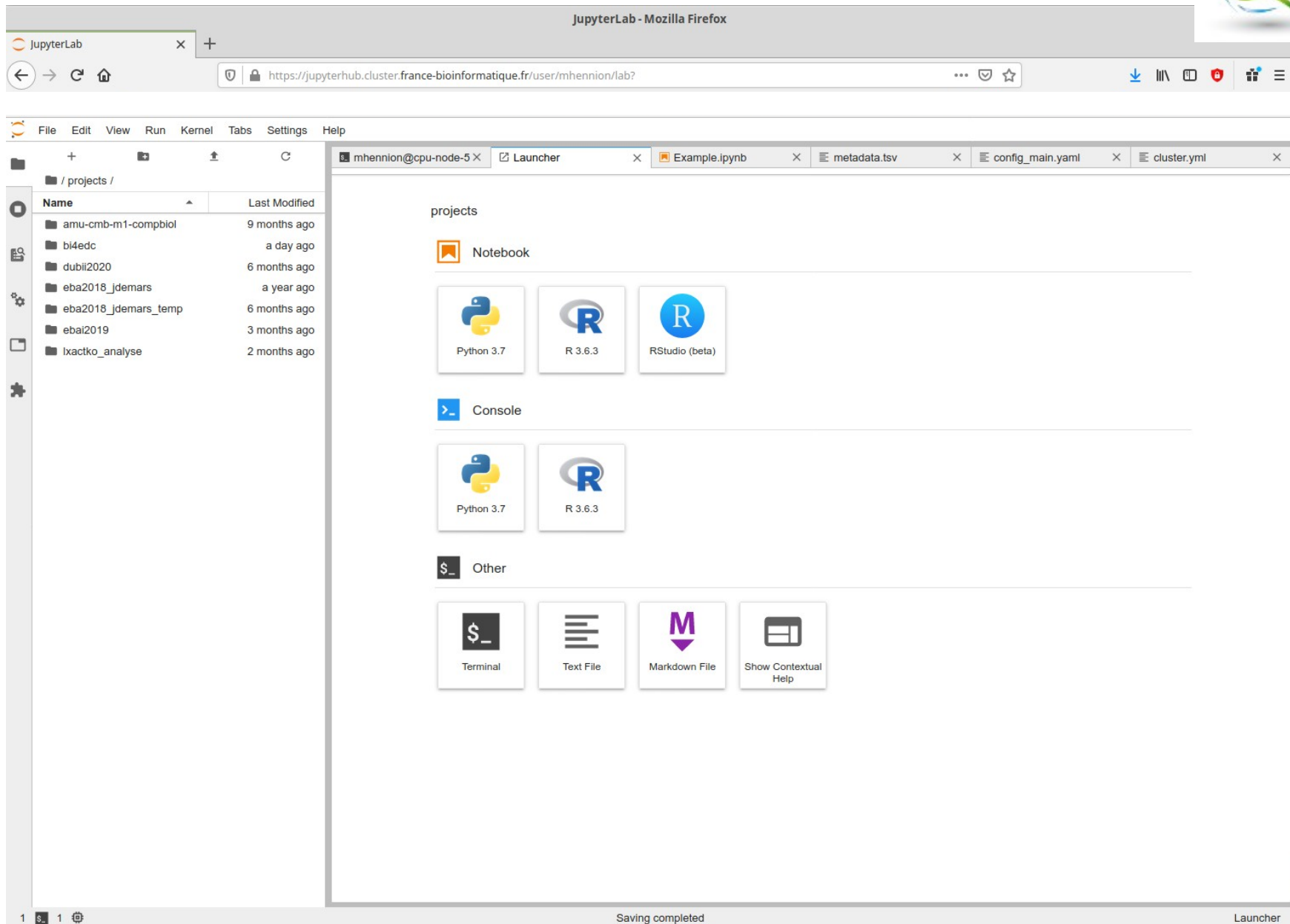
```
## Time difference of 14.73 secs
```

R session information.

```
## - Session info -  
## setting value  
## version R version 3.6.3 (2020-02-29)  
## os Linux Mint 19.1  
## system x86_64, linux-gnu  
## ui X11  
## language en_US  
## collate en_US.UTF-8  
## ctype en_US.UTF-8  
## tz Europe/Paris  
## date 2020-06-22  
##  
## - Packages -  
## package * version date lib source  
## acepack 1.4.1 2016-10-29 [1] CRAN (R 3.6.3)  
## annotate 1.64.0 2019-10-29 [1] Bioconductor  
## AnnotationDbi 1.48.0 2019-10-29 [1] Bioconductor  
## askpass 1.1 2019-01-13 [1] CRAN (R 3.6.1)  
## assertthat 0.2.1 2019-03-21 [1] CRAN (R 3.6.1)  
## backports 1.1.5 2019-10-02 [1] CRAN (R 3.6.1)  
## base64enc 0.1-3 2015-07-28 [1] CRAN (R 3.6.1)  
## bibtex 0.4.2.2 2020-01-02 [1] CRAN (R 3.6.3)  
## Biobase * 2.46.0 2019-10-29 [1] Bioconductor  
## BiocFileCache 1.10.2 2019-11-08 [1] Bioconductor  
## BiocGenerics * 0.32.0 2019-10-29 [1] Bioconductor  
## BiocManager 1.30.10 2019-11-16 [1] CRAN (R 3.6.3)  
## BiocParallel * 1.20.1 2019-12-21 [1] Bioconductor  
## BiocStyle * 2.14.4 2020-01-09 [1] Bioconductor  
## biomaRt 2.42.0 2019-10-29 [1] Bioconductor  
## Biostrings 2.54.0 2019-10-29 [1] Bioconductor  
## bit 1.1-15.2 2020-02-10 [1] CRAN (R 3.6.3)  
## bit64 0.9-7 2017-05-08 [1] CRAN (R 3.6.3)  
## bitops 1.0-6 2013-08-17 [1] CRAN (R 3.6.3)  
## blob 1.2.1 2020-01-20 [1] CRAN (R 3.6.3)  
## bookdown 0.19 2020-05-15 [1] CRAN (R 3.6.3)  
## BSgenome 1.54.0 2019-10-29 [1] Bioconductor  
## bumphunter 1.28.0 2019-10-29 [1] Bioconductor  
## checkmate 2.0.0 2020-02-06 [1] CRAN (R 3.6.3)  
## cli 2.0.2 2020-02-28 [1] CRAN (R 3.6.1)  
## cluster 2.1.0 2019-06-19 [1] CRAN (R 3.6.3)  
## codetools 0.2-16 2018-12-24 [1] CRAN (R 3.6.3)  
## colorspace 1.4-1 2019-03-18 [1] CRAN (R 3.6.1)  
## crayon 1.3.4 2017-09-16 [1] CRAN (R 3.6.1)  
## crosstalk 1.0.0 2016-12-21 [1] CRAN (R 3.6.1)
```

```
## crosstalk 1.0.0 2016-12-21 [1] CRAN (R 3.6.1)  
## curl 4.3 2019-12-02 [1] CRAN (R 3.6.1)  
## data.table 1.12.8 2019-12-09 [1] CRAN (R 3.6.3)  
## DBI 1.1.0 2019-12-15 [1] CRAN (R 3.6.1)  
## dbplyr 1.4.2 2019-06-17 [1] CRAN (R 3.6.1)  
## DEFormats 1.14.0 2019-10-29 [1] Bioconductor  
## DelayedArray * 0.12.2 2020-01-06 [1] Bioconductor  
## derfinder 1.20.0 2019-10-29 [1] Bioconductor  
## derfinderHelper 1.20.0 2019-10-29 [1] Bioconductor  
## DESeq2 * 1.26.0 2019-10-29 [1] Bioconductor  
## digest 0.6.25 2020-02-23 [1] CRAN (R 3.6.1)  
## doRNG 1.8.2 2020-01-27 [1] CRAN (R 3.6.3)  
## dplyr 0.8.5 2020-03-07 [1] CRAN (R 3.6.3)  
## DT * 0.12 2020-02-05 [1] CRAN (R 3.6.1)  
## edgeR * 3.28.1 2020-02-26 [1] Bioconductor  
## evaluate 0.14 2019-05-28 [1] CRAN (R 3.6.1)  
## fansi 0.4.1 2020-01-08 [1] CRAN (R 3.6.1)  
## farver 2.0.3 2020-01-16 [1] CRAN (R 3.6.1)  
## fastmap 1.0.1 2019-10-08 [1] CRAN (R 3.6.1)  
## foreach 1.5.0 2020-03-30 [1] CRAN (R 3.6.3)  
## foreign 0.8-76 2020-03-03 [1] CRAN (R 3.6.3)  
## Formula 1.2-3 2018-05-03 [1] CRAN (R 3.6.3)  
## genefilter 1.68.0 2019-10-29 [1] Bioconductor  
## geneplotter 1.64.0 2019-10-29 [1] Bioconductor  
## GenomeInfoDb * 1.22.0 2019-10-29 [1] Bioconductor  
## GenomeInfoDbData 1.2.2 2020-03-10 [1] Bioconductor  
## GenomicAlignments 1.22.1 2019-11-12 [1] Bioconductor  
## GenomicFeatures 1.38.2 2020-02-15 [1] Bioconductor  
## GenomicFiles 1.22.0 2019-10-29 [1] Bioconductor  
## GenomicRanges * 1.38.0 2019-10-29 [1] Bioconductor  
## ggplot2 * 3.3.0 2020-03-05 [1] CRAN (R 3.6.3)  
## glue 1.3.1 2019-03-12 [1] CRAN (R 3.6.1)  
## gridExtra 2.3 2017-09-09 [1] CRAN (R 3.6.3)  
## gtable 0.3.0 2019-03-25 [1] CRAN (R 3.6.1)  
## highr 0.8 2019-03-20 [1] CRAN (R 3.6.1)  
## Hmisc 4.3-1 2020-02-07 [1] CRAN (R 3.6.3)  
## hms 0.5.3 2020-01-08 [1] CRAN (R 3.6.1)  
## htmlTable 1.13.3 2019-12-04 [1] CRAN (R 3.6.3)  
## htmltools 0.4.0 2019-10-04 [1] CRAN (R 3.6.1)  
## htmlwidgets 1.5.1 2019-10-08 [1] CRAN (R 3.6.1)  
## httpuv 1.5.2 2019-09-11 [1] CRAN (R 3.6.1)  
## httr 1.4.1 2019-08-05 [1] CRAN (R 3.6.1)  
## IRanges * 2.20.2 2020-01-13 [1] Bioconductor  
## IRdisplay 0.7.0 2018-11-29 [1] CRAN (R 3.6.1)  
## IRkernel 1.1 2020-03-03 [1] Github (IRkernel/IRkernel@1a45f74)  
## iterators 1.0.12 2019-07-26 [1] CRAN (R 3.6.3)  
## jpeg 0.1-8.1 2019-10-24 [1] CRAN (R 3.6.3)  
## jsonlite 1.6.1 2020-02-02 [1] CRAN (R 3.6.1)  
## knitr 1.0.10 2019-09-15 [1] CRAN (R 3.6.3)  
## knitr * 1.28 2020-02-06 [1] CRAN (R 3.6.3)  
## knitrBootstrap 1.0.2 2018-05-24 [1] CRAN (R 3.6.3)  
## labeling 0.3 2014-08-23 [1] CRAN (R 3.6.1)  
## later 1.0.0 2019-10-04 [1] CRAN (R 3.6.1)  
## lattice 0.20-40 2020-02-19 [1] CRAN (R 3.6.1)  
## latticeExtra 0.6-29 2019-12-19 [1] CRAN (R 3.6.3)  
## lifecycle 0.2.0 2020-03-06 [1] CRAN (R 3.6.3)  
## limma * 3.42.2 2020-02-03 [1] Bioconductor  
## locfit 1.5-9.1 2013-04-20 [1] CRAN (R 3.6.3)  
## lubridate 1.7.4 2018-04-11 [1] CRAN (R 3.6.1)  
## magick 2.3 2020-01-24 [1] CRAN (R 3.6.3)  
## magrittr 1.5 2014-11-22 [1] CRAN (R 3.6.1)  
## markdown 1.1 2019-08-07 [1] CRAN (R 3.6.1)  
## Matrix 1.2-18 2019-11-27 [1] CRAN (R 3.6.1)  
## matrixStats * 0.55.0 2019-09-07 [1] CRAN (R 3.6.3)  
## memoise 1.1.0 2017-04-21 [1] CRAN (R 3.6.1)  
## mime 0.9 2020-02-04 [1] CRAN (R 3.6.1)  
## munsell 0.5.0 2018-06-12 [1] CRAN (R 3.6.1)  
## nnet 7.3-13 2020-02-25 [1] CRAN (R 3.6.3)  
## openssl 1.4.1 2019-07-18 [1] CRAN (R 3.6.1)  
## pbdZMQ 0.3-3 2018-05-05 [1] CRAN (R 3.6.1)  
## pheatmap * 1.0.12 2019-01-04 [1] CRAN (R 3.6.3)  
## pillar 1.4.3 2019-12-20 [1] CRAN (R 3.6.1)  
## pkgconfiq 2.0.3 2019-09-22 [1] CRAN (R 3.6.1)
```

Make it easier



All your analysis from online hub





JupyterLab - Mozilla Firefox

pariseigenetics/RASflow\_IFB Tutorial RASflow on IFB core clus

https://jupyterhub.cluster.france-bioinformatique.fr/user/mhennion/lab?

File Edit View Run Kernel Tabs Settings Help

mhennion@cpu-node-59:sh X Example.ipynb X metadata.tsv X config\_main.yaml X cluster.yml X

pwd  
(base) [mhennion @ cpu-node-59 13:28]\$ ~ : pwd  
/shared/home/mhennion  
(base) [mhennion @ cpu-node-59 13:28]\$ ~ : lx  
(base) [mhennion @ cpu-node-59 13:29]\$ RASflow : squeue

JOBID	PARTITION	NAME	USER	ST	TIME	NODES	MODELIST (REASON)
9874354	bigmem	DNA_pool	echase	PD	0:00	1	(QOSMaxMemoryPerUser)
4369319	fast	_interac	wbendhaf	PD	0:00	1	(launch failed requested held)
13292207	fast	jupyter	mhennion	R	58:39	1	cpu-node-59
13292112	fast	bash	pgladieu	R	2:12:13	1	cpu-node-59
13291947	fast	g165184_	galaxy	R	7:47:22	1	cpu-node-59
13292177	fast	g165296_	galaxy	R	1:27:52	1	cpu-node-59
13292268	bigmem	masurca	rguyot	R	22:49	1	cpu-node-69
13292231	fast	snakejob	lchauvie	R	27:11	1	cpu-node-61
13292244	fast	snakejob	lchauvie	R	23:50	1	cpu-node-63
13292094	fast	align.sh	lchauvie	R	2:21:51	1	cpu-node-63
13279151	long	bcftools	fcharria	R	3-04:41:14	1	cpu-node-13
13292227	fast	snakejob	lchauvie	R	42:55	1	cpu-node-7
13292097	fast	snakejob	lchauvie	R	2:19:27	1	cpu-node-59
13292259	fast	snakejob	lchauvie	R	19:52	1	cpu-node-62
13292269	fast	snakejob	lchauvie	R	14:03	1	cpu-node-59
13292270	fast	snakejob	lchauvie	R	14:03	1	cpu-node-7
13292281	fast	snakejob	lchauvie	R	9:05	1	cpu-node-61
13292256	fast	snakejob	lchauvie	R	21:48	1	cpu-node-6
13292258	fast	snakejob	lchauvie	R	21:46	1	cpu-node-9
13292257	fast	snakejob	lchauvie	R	21:47	1	cpu-node-60
13292288	long	multiple	lherault	R	6:28	1	cpu-node-15
13292287	long	multiple	lherault	R	7:45	1	cpu-node-16
13292286	long	multiple	lherault	R	7:48	1	cpu-node-18
13292285	long	multiple	lherault	R	8:03	1	cpu-node-17
13292284	long	multiple	lherault	R	8:07	1	cpu-node-19
13292283	long	multiple	lherault	R	8:32	1	cpu-node-14
13292282	long	multiple	lherault	R	9:34	1	cpu-node-12
13292040	fast	align.sh	lchauvie	R	4:04:08	1	cpu-node-6
13292291	fast	snakejob	lchauvie	R	2:56	1	cpu-node-13
13292290	fast	snakejob	lchauvie	R	3:06	1	cpu-node-8
13292280	fast	snakejob	lchauvie	R	13:56	1	cpu-node-60
13281751	long	snakejob	fcharria	R	2-04:34:15	1	cpu-node-11
13281743	long	Snake	fcharria	R	2-04:35:33	1	cpu-node-11
13202916	long	prod_S0_	tlasafr	R	9-03:59:25	1	cpu-node-11
13202917	long	prod_S0_	tlasafr	R	9-03:59:25	1	cpu-node-11
13202918	long	prod_S0_	tlasafr	R	9-03:59:25	1	cpu-node-11
13202919	long	prod_S0_	tlasafr	R	9-03:59:25	1	cpu-node-11
13202920	long	surv_S0_	tlasafr	R	9-03:59:25	1	cpu-node-11
13202921	long	surv_S0_	tlasafr	R	9-03:59:25	1	cpu-node-11
13202922	long	surv_S0_	tlasafr	R	9-03:59:25	1	cpu-node-11
13202923	long	surv_S0_	tlasafr	R	9-03:59:25	1	cpu-node-11
13202924	long	surv_S0_	tlasafr	R	9-03:59:25	1	cpu-node-11
13202925	long	surv_S0_	tlasafr	R	9-03:59:25	1	cpu-node-11

1 Saving completed mhennion@cpu-node-59:/shared/projects/lxactko\_analyse/RASflow



JupyterLab - Mozilla Firefox

pariseigenetics/RASflow\_IFB X Tutorial RASflow on IFB core clus X +

https://jupyterhub.cluster.france-bioinformatique.fr/user/mhennion/lab?

File Edit View Run Kernel Tabs Settings Help

mhennion@cpu-node-59:/sh X Example.ipynb X metadata.tsv X config\_main.yaml X cluster.yml X

18 minutes ago 5 minutes ago 3 months ago a month ago 2 minutes ago 4 months ago a month ago a month ago a day ago a minute ago 2 months ago a month ago a month ago 2 months ago 2 months ago a month ago 7 days ago 2 months ago

ifb main\_cluster.py main.py new previous sacct.tsv StarIndex.sh template.sh test.py Test.sh Unlock.sh Workflow.sh

```

1 # Please check the parameters, and adjust them according to your circumstance
2
3 # Project name
4 PROJECT: LXACT-test3
5
6 # ===== Control of the workflow =====
7
8 ## Do you need to do quality control?
9 QC: no # "yes" or "no". If set to "yes", the workflow will stop after the QC to let you decide whether you want to trim your raw data
   or not. In order to run the rest of the workflow, you have to set it to "no".
10
11 ## Do you need to do trimming?
12 TRIMMED: yes # "yes" or "no"
13
14 ## Do you need to do mapping
15 MAPPING: yes # "yes" or "no"
16
17 ## Which mapping reference do you want to use? Genome or transcriptome?
18 REFERENCE: genome # "genome" or "transcriptome", I haven't implemented transcriptome yet.
19
20 ## Do you want to do Differential Expression Analysis (DEA)?
21 DEA: yes # "yes" or "no"
22
23 ## Do you want to visualize the results of DEA?
24 VISUALIZE: yes # "yes" or "no"
25
26 # ===== Shared parameters for some or all of the sub-workflows =====
27
28 ## key file if the data is stored remotely, otherwise leave it empty
29 KEY:
30
31 ## the path to fastq files
32 READSPATH: /shared/projects/lxactko_analyse/Raw_fastq
33
34 ## the meta file describing the experiment settings
35 METAFILE: /shared/projects/lxactko_analyse/RASflow/configs/metadata-test2.tsv
36
37 ## paths for intermediate and final results
38 BIGDATAPATH: /shared/projects/lxactko_analyse/RASflow/data # for big files
39 RESULTPATH: /shared/projects/lxactko_analyse/RASflow/results
40
41 ## is the sequencing paired-end or single-end?
42 END: pair # "pair" or "single"
43
44 ## number of cores you want to allocate to this workflow
45 NCORE: 30 # Use command "getconf _NPROCESSORS_ONLN" to check the number of cores/CPU on your machine
46
47
48 # ===== Configuration for Quality Control =====
49

```

1 1 YAML Saving completed Ln 1, Col 1 Spaces: 4 config\_main.yaml



**RASflow tests**

Following [this tutorial](#), I will analyse RNAseq data obtained from ...

First I check where I am:

```
[1]: %bash
pwd

/shared/mfs/data/projects/lxactko_analyse/RASflow
```

Then I modify `metadata.tsv` and `config_main.yaml` (in `configs` folder, panel on the left).

Now I can start my run:

```
[3]: %bash
sbatch Workflow.sh
```

Submitted batch job 13292308

I can check that my job is running:

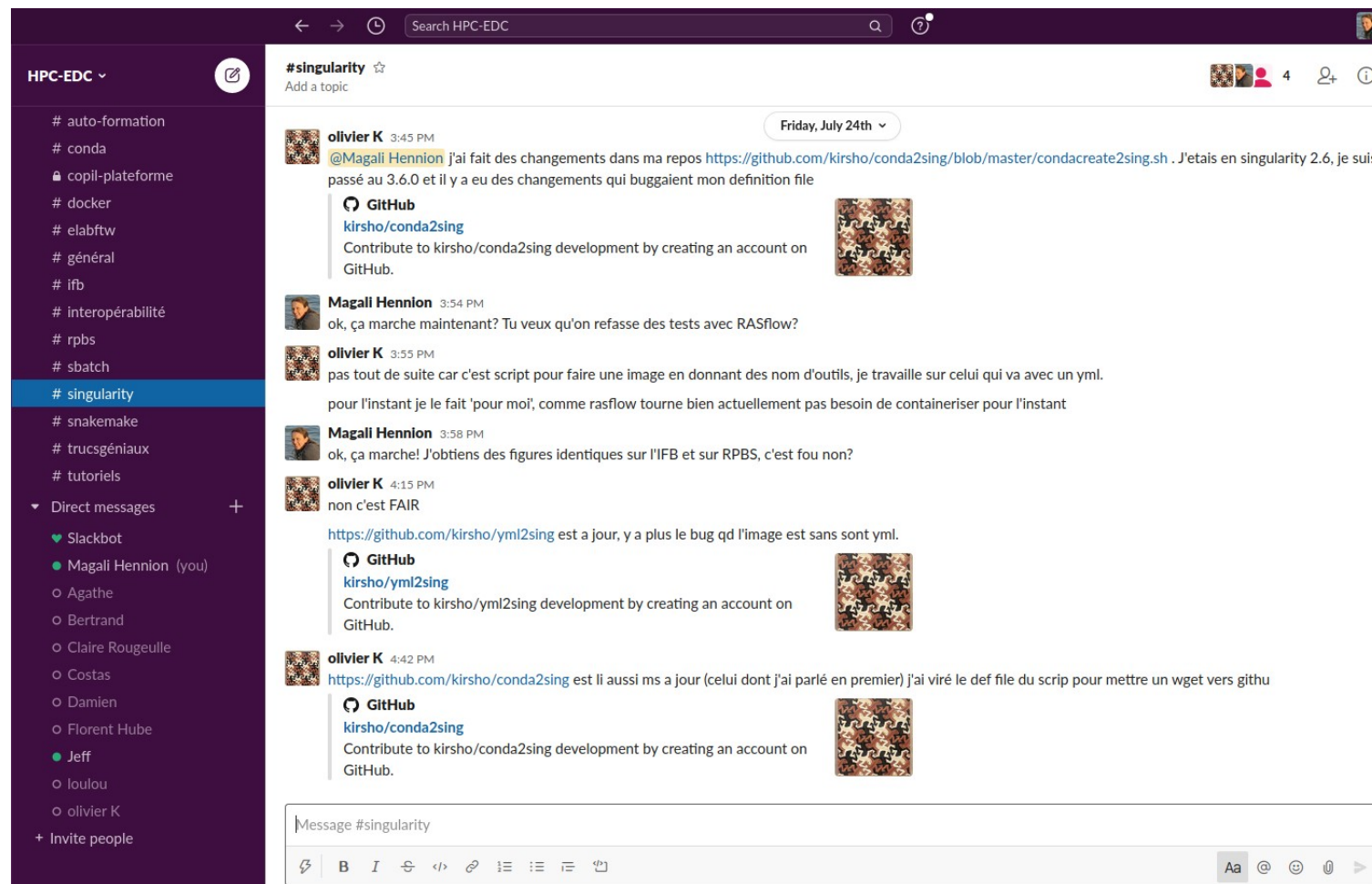
```
[6]: %bash
squeue -u mhennion
```

JOBID	PARTITION	NAME	USER	ST	TIME	NODES	NODELIST(REASON)
13292207	fast	jupyter	mhennion	R	1:10:18	1	cpu-node-59
13292310	fast	trimming	mhennion	R	0:18	1	cpu-node-59
13292308	fast	FromCoun	mhennion	R	1:17	1	cpu-node-59

# Communication

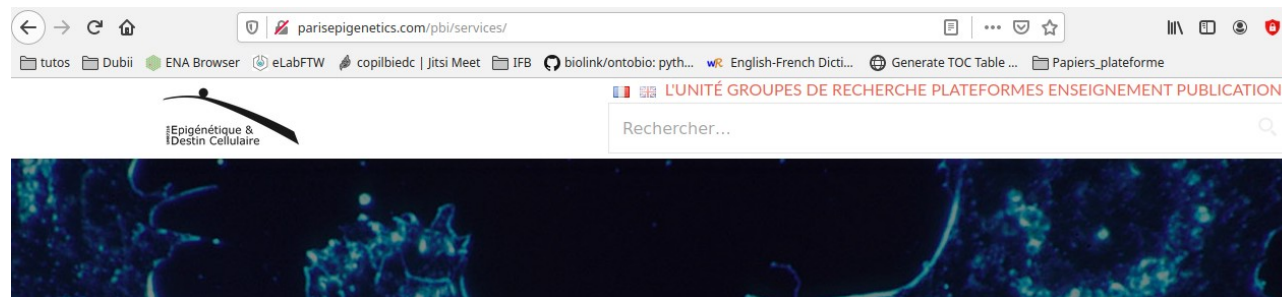
## Slack : [hpc-edc.slack.com](https://hpc-edc.slack.com)

→ Ask your questions, help the others, complain, ...



# Communication

Webpage on EDC website : <http://parisepigenetics.com/pbi/services/>



[Retour](#)

## Services

Bienvenue sur la page de la plateforme BIBS – Bioinformatique et Biostatistiques – de l'unité Epigénétique et Destin Cellulaire.

Rechercher...

## Mission de la plateforme BIBS

La plateforme BIBS de l'UMR7216 développe et met à disposition des protocoles standardisés pour l'analyse de données NGS en épigénomique. Nous maintenons également l'accès à une ressource de calcul locale et nous fournissons aux équipes du laboratoire un éventail de services liés à la bioinformatique. Ces services incluent notamment la veille technologique, des analyses spécifiques, ainsi que des formations personnalisées pour nos utilisateurs.

## Objectifs de la plateforme

- Fournir un accès à une ressource de calcul dédiée à l'analyse NGS.
- Maintenir et disséminer les outils, pipelines et bases de données développés par nos utilisateurs (e.g. RNA-seq, ChIP-seq, Ribosome profiling, etc.)
- Installer et maintenir des images Docker et Singularity des outils NGS.
- Développer des pipelines d'analyse (Snakemake et NextFlow).
- Promouvoir la collaboration et l'échange de connaissances.
- Organiser des formations.

## Comment accéder à la plateforme

Si vous désirez discuter et mettre en place un projet avec la plateforme, merci de remplir le formulaire disponible ici et de l'envoyer à [Magali Hennion](#). Une réunion pour discuter de vos besoins sera organisée dans les jours suivants.

L'accès à nos ressources de calcul est possible sur demande. Ce cluster nécessite l'utilisation de Docker ou de Singularity. Si besoin, la plateforme peut vous former. La documentation est disponible sur le site internet de la plateforme.

Les protocoles, scripts et mode d'emploi du cluster sont hébergés sur [notre dépôt GitHub](#). Une partie des ressources est privée, pour y avoir accès, merci de contacter [Magali Hennion](#) et de lui fournir votre identifiant GitHub.

## Site internet de la plateforme

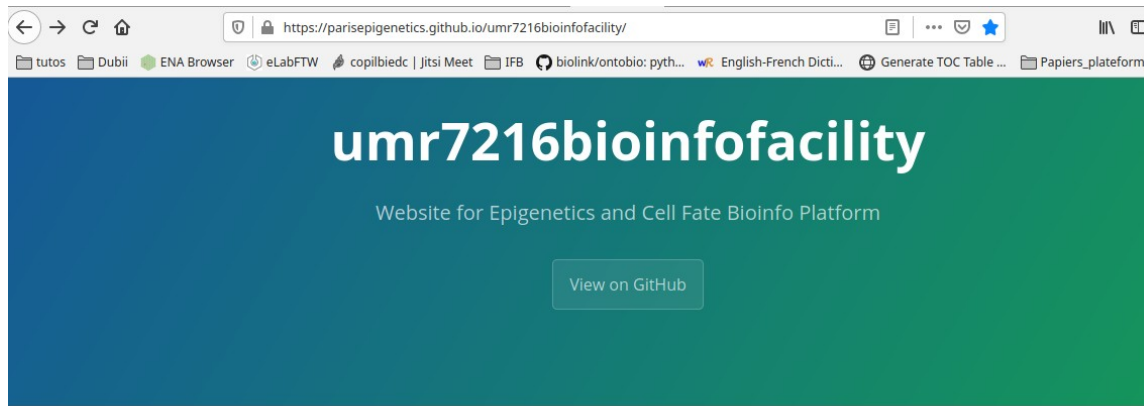
Vous trouverez plus d'information sur le site internet de la plateforme BIBS : <https://parisepigenetics.github.io/umr7216bioinfofacility/>





# Communication

Website : <https://parisepigenetics.github.io/umr7216bioinfofacility/>



## EDC Bioinformatics Core Facility

Welcome to the Epigenetics and Cell Fate (UMR7216) Bioinformatics website.

The UMR7216 bioinformatics platform provides and develops user-friendly, state of the art epigenomics protocols. We also maintain access to a local computing cluster and provide the teams of the UMR7216 a variety of bioinformatics related services such as technology monitoring, project-specific analyses and user-tailored, on-demand training.

This website is the main resource for protocols, scripts and instructions on how to use the UMR7216 cluster.

## How to access the facility

To discuss and set-up a project with the platform, please download and fill-in the form available [here](#) and send it via email to [Magali Hennion](#). A meeting to discuss your needs will be set-up in the following days.

Update on going...

Feedback welcome !

# A word about iPOP-UP

## Mission statement

- Organize a resource for bioinformatics in the U-Paris landscape
- Assist/advise users on specific projects
- Train users to the use of the resource

RPBS cluster will be upgraded and **open** to the community

→ goal : starting production in January 2021

## Technical committee

Pierre Tuffery (BFA, RPBS)  
Julien Rey (BFA, RPBS)  
Jean-Philippe Jais (Necker)  
Pierre Poulain (IJM)  
Christophe Cerin (Paris 13 Nord)  
Magali Hennion (EDC)

Sjoerd DeVries (BFA, RPBS)  
Niclas Setterblad (Saint-Louis)  
Yves Clement (IJM)  
Guillaume Seith (IGBMC, IFB)  
Benjamin Saintpierre (Cochin)

# A word about iPOP-UP

## **Where we are**

- a system administrator (Abdeslam Tahari) has been recruited
- new equipment will be ordered and installed soon
- IR open position (EDC)



# A word about iPOP-UP

## **Where we are**

- a system administrator (Abdeslam Tahari) has been recruited
- new equipment will be ordered and installed soon
- IR open position (EDC)

→ If you have specific needs or great ideas, it's time to let me know!

→ RPBS people want us (and other partners) to be more involved

If you want to be part of the task force that manage the resource and organize the services,... or to do beta-testing, please let me know.

## Acknowledgments

*Olivier*

*Valérie and Claire*

*The CoPil*

*The beta-testers*

*Thank you for your attention!*



ifb  
INSTITUT FRANÇAIS  
DE BIOINFORMATIQUE

**Community Support**



Julien Rey