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# User feedback of a Galaxy workflow conception

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# How have I built the Workflow?

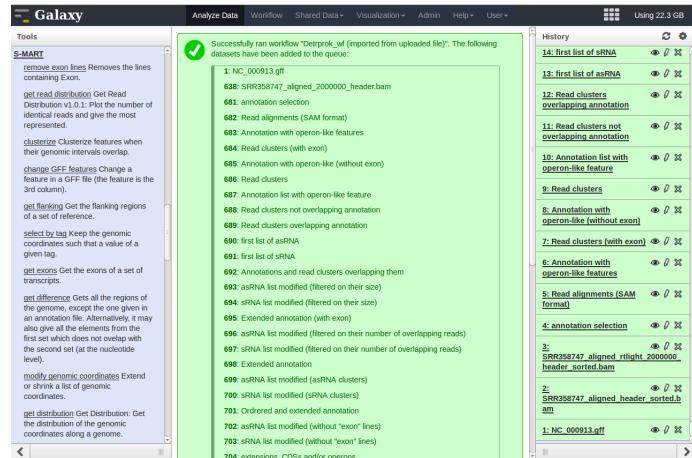
step-by-step, biologist

Data = README file

s-mart tools, home-made scripts

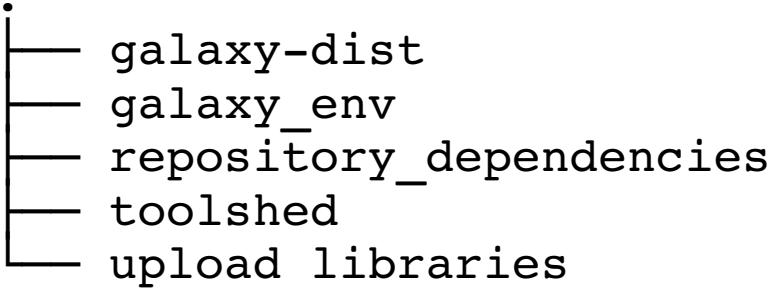
```
### 1- select reads outside "gene" :
# select reads (-i VIBSP_chrl_smart.gff3) outside
# "annotation" (-j NC_011744_annot.gff3 -c -x), with
# keeping reads overlapping ATG (-e 10)
for i in "NC_011744" "NC_011753" ; do
    compareOverlapping.py -i ${i}_RNAseq_smart.gff3 -f gff3
    -j ${i}_annot.gff -g gff3 -e -10 -x -c -o ${i}_cis_ReadsOutGene ;
done ;
### 2- clusterize reads before seed selection :
# 2 clusterization steps: strict + d < 20
for i in "NC_011744" "NC_011753" ; do
    clusterize.py -i ${i}_cis_ReadsOutGene.gff3 -f gff3 -c -o ${i}_cis_e10_clusters ;
    clusterize.py -i ${i}_cis_e10_clusters.gff3 -f gff3 -c -d 20 -o ${i}_cis_e10_clusters20 ;
### 3- crossing cluster with seed :
for i in "NC_011744" "NC_011753" ; do
    seedGff.pl -i ${i}_annot.gff -p 25 -e 15 -o ${i}_cis_seed.gff ; #
seed creation in -15 -25 bp from ATG
    compareOverlapping.py -i ${i}_cis_e10_clusters20.gff3 -f gff3 -j ${i}_cis_seed.gff -g gff3 -c -o ${i}_cis_e10_cluster20InSeed2515 ; #
crossing cluster with seed
done ;
#
### 4- filters: nb_reads > 10 reads + length > 50 nt
for i in "NC_011744" "NC_011753" ; do
    selectByTag.py -i ${i}_cis_e10_cluster20InSeed2515.gff3 -f gff -o ${i}_cis_e10_cluster20InSeed2515_nbEUp10 -g "nbElements" -m 10 -d 0 ; # nb_reads > 10 filter
done ;
...
```

Wrap tool, share (test/toolshed)  
Import on my local Galaxy instance  
Manual launch, follow the readme file



Extract Workflow from History  
Edit workflow: close or open parameter  
(« set at run time »), add comments,  
change name of the steps  
Share Workflow (test/toolshed)

# Technical issues

- Use « virtual\_env »
  - Dissociate « your » directories from the Galaxy distribution (configure the universe\_wsgi.ini file)
  - « underlying » database :  
SQLite => postgres  
allow concurrential accesses, count DB in the ressources needed
  - Install a Galaxy instance on a stratuslab virtual machine during revision process of the publication
- 
- ```
graph TD; . --- galaxy-dist; . --- galaxy_env; . --- repository_dependencies; . --- toolshed; . --- upload_libraries;
```



# Feedbacks

## User's point of view: local instance vs. Galaxy server ?

User : import workflows, not the tools (admin account).

Ok in a local instance, contact the admin in a server => not so easy !

## Open questions (perhaps have you the answers?):

- How expliciting the need of specific tools? **dependencies?**
- « **Global** » **variables?** Launching a tool many times implies to re-enter the same values for open parameters. Environment variables?
- **Meta-workflow?** Begin = bam file (user chooses the mapper). How add the mapping step? another workflow and combine the two?

# DETR'PROK



## Methods

Volume 63, Issue 1, 1 September 2013, Pages 60–65

Diversity of the non-coding transcriptomes revealed by RNA-seq technologies



## Detection of non-coding RNA in bacteria and archaea using the DETR'PROK Galaxy pipeline

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2279 valid tools on Oct 24, 2013

Repositories with matching workflows

workflow name: **detrprok**  
exact matches only: False

| <input type="checkbox"/> Repository name | Synopsis                                      | Revision     | Owner    |
|------------------------------------------|-----------------------------------------------|--------------|----------|
| <input type="checkbox"/> detrprok_wf     | ncRNA detection in prokaryote oriented RNAseq | f0ca0981eb86 | clairetn |
| <input type="checkbox"/> detrprok_wf     | ncRNA detection in prokaryote oriented RNaseq | 8f7eb127baf3 | clairetn |
| <input type="checkbox"/> detrprok_wf     | ncRNA detection in prokaryote oriented RNAseq | bb71a378053a | clairetn |

For 0 selected items:

