Introduction to Galaxy (answers to questions)

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Answer 1 : Log in

• Go to <u>http://use.galaxeast.fr</u>

🚍 Galaxy / Galaxeast	Analyze Data Workflow	Shared Data -	Visualization 🗸	Help -	User⊤		Using 0 bytes
This Galaxy instance has been configured s	uch that only users who are I	ogged in may use	e it.				
Login							
Username / Email Address: Password:		(us	Enter you erN@gala	r log axea	gin st.fr)		
Forgot password? Reset here		— En (1	ter your p NGStraini	oassy ng09	word 920)	1	

Answer 2 : History

• Create a new history



 Change the name of the new history to "RNA-seq data analysis" by clicking on "Unnamed history" on top of the history panel. Then type "RNA-seq data analysis" and [ENTER].

Histor	y 2 ‡⊡
sear	ch datasets 🛛 🕄
Unnan	ned history
0 b	Click to rename history 🔊 🗩

Answer 3 : Import data to Galaxy

• 1.

- Go to Shared data (top menu) > Data libraries.
- Go to NGS data analysis training > RNAseq > rawdata.
- Tick the box beside the sample name "siLuc3_S12040.fastq".
- Click on the button "to History".
- The history "RNAseq data analysis" is selected. Click on import.
- Click on "Analyze Data" (top menu) to go back to the main Galaxy page.



4

Answer 3 : Import data to Galaxy

- 3
 - Click on the button to display the drag and drop utility 🔔
 - Drag and drop the file sample.bed.gz into the opened window.
 - Select Type: bed
 - Select Genome: Mouse July 2007 (NCBI37/mm9) (mm9)

Download from web or upload from disk

Name	Size	Туре	Genome	Settings	Status	
□ sample.bed.gz	482.9 KB	bed 🔻 Q	Mouse July 2007 (🔻	•		Û
Type (set all):	Auto-det	ect 🔻 Q	Genome (set all):	unspec	ified (?)	
	므 Choose	local file 🕞 Choos	e FTP file 🛛 🖉 Paste/Fe	tch data Pause	e Reset Star	CI

Answer 4: remove dataset

• 1. Click on the cross of the dataset box in the history



- B) Click on "hide deleted"
- NOTE: to permanently remove from disk multiple deleted datasets, click on the History option button and select "Purge deleted datasets"

Answer 5 : Running a tool

• Search "fastqc" in the list or using the search field of the tool panel.



• Select the file to analyze and click on "Execute"

FastQC Read Quality reports (Galaxy Version 0.63)	▼ Options					
Short read data from your current history						
1: siLuc3_S12040.fastq	•					
Contaminant list						
C C Nothing selected	-					
tab delimited file with 2 columns: name and sequence. For example: Illumina Small RNA RT Primer CAAGCAGAAGACGGCATACGA						
Submodule and Limit specifing file						
C Nothing selected	-					
a file that specifies which submodules are to be executed (default=all) and also specifies the thresholds for the each submodules was	rning					
parameter						
✓ Execute						

t.