Introduction to Galaxy (answers to questions)

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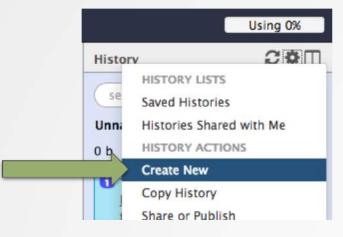
Answer 1 : Log in

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

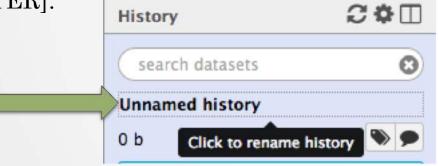
= Galaxy / Galaxeast	Analyze Data	Workflow	Shared Data -	Visualization -	Help 🗸	User≁	Using 0 bytes
This Galaxy instance has been configured s	such that only use	ers who are l	ogged in may us	e it.			
Login							
Username / Email Address: Password:			— E	nter you (user		gin	
Forgot password? Reset here				ter your GStrain	-		

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].



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.



Answer 3 : Import data to Galaxy

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- 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

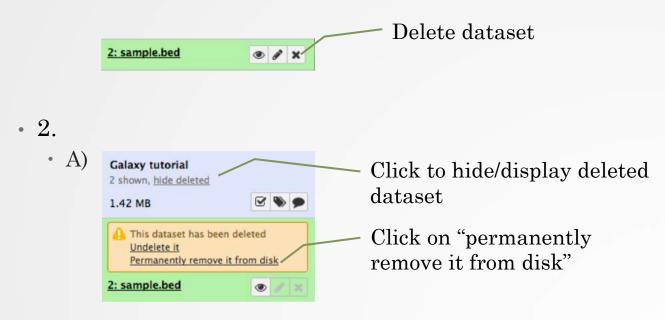
Download from web or upload from disk

• Select Genome: Mouse July 2007 (NCBI37/mm9) (mm9)

		You add	ed 1 file(s) to the queu	e. Add more fil	es or click 'Start'	to proceed.		
Name		Size	Туре	e Genome		Settings	Status	
	sample.bed.gz	482.9 KB	bed 💌 Q	Mouse Jul	y 2007 (💌	٥		ê
Type (set all):	Auto-det	ect 💌 Q	Gen	ome (set all):		unspecified (?)		
		Choose	local file 🖻 Cho	ose FTP file	Paste/Feto	ch data	Pause Reset Start	

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. Tools

Enter: "fastqc" Click on the tool name FastQC Read Quality reports

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

read data	from your current history	
20	1: siLuc3_\$12040.fastq	
minant lis		
20	Nothing selected	
limited file	with 2 columns: name and sequence. For example: Illumina Small RNA RT	Primer CAAGCAGAAGACGGCATACGA
odule and	Limit specifing file	
20	Nothing selected	
hat specifi	s which submodules are to be executed (default=all) and also specifies the	e thresholds for the each submodules warning
	Iminant list Iminant list Imited file Imited file Imited file Imited file	minant list C Nothing selected limited file with 2 columns: name and sequence. For example: Illumina Small RNA RT odule and Limit specifing file

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