# ACEMFS FUT Minna Bioinformatics Workshop

Sequence Retrieval & Quality Control using Galaxy

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#### **Workshop Outline**

By the end of this session, participants will be able to:

- Retrieve sequences of mycotoxins & fungal enzymes from databases.
- Upload & organize datasets in <u>Galaxy</u>.
- Perform QC using Galaxy tools (<u>FastQC</u>)

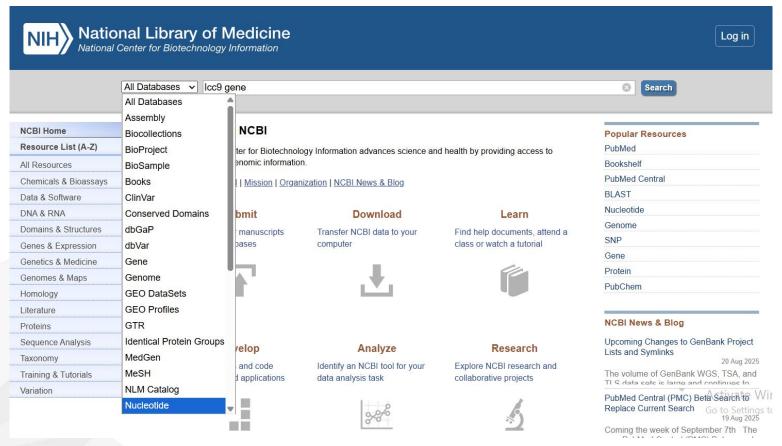
#### **Introduction to Galaxy**

- Web-based platform for bioinformaticians.
- No coding required.
- Supports reproducible research.
- Widely used for genomics, proteomics and transcriptomics analysis.



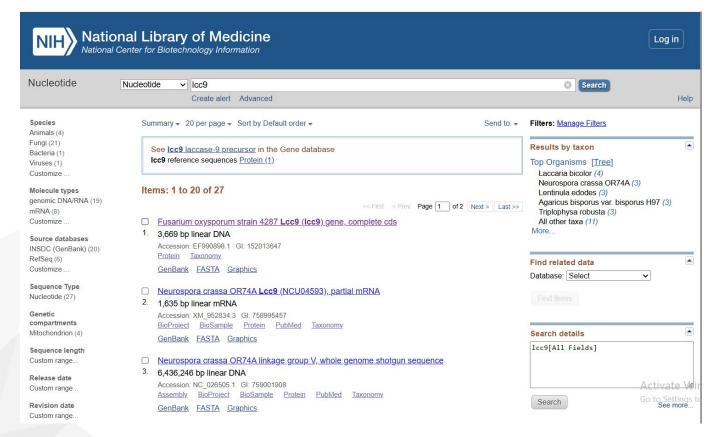
#### NCBI Exploration:

- 1. Go to NCBI
- 2. In the search bar, paste the **Icc9** gene.
- 3. Click on the dropdown arrow next to the search bar and select **Nucleotide**.
- 4. Click the search button.

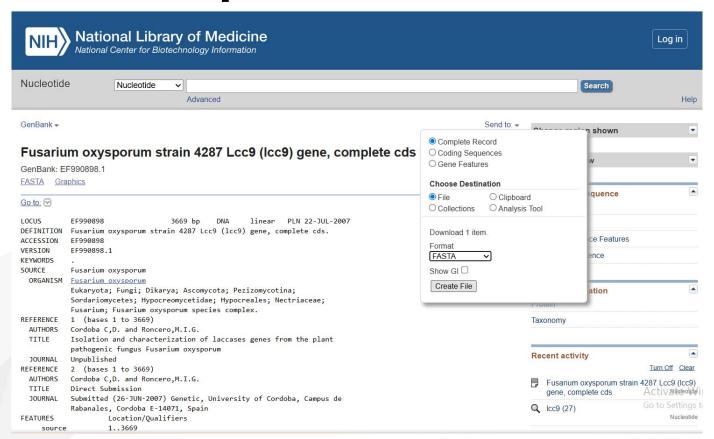


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- 5. On the results page click the **first hit**.
- 6. What is the name of the **organism**?
- 7. How many **base pairs** does it have?
- 8. What is its accession and version numbers?



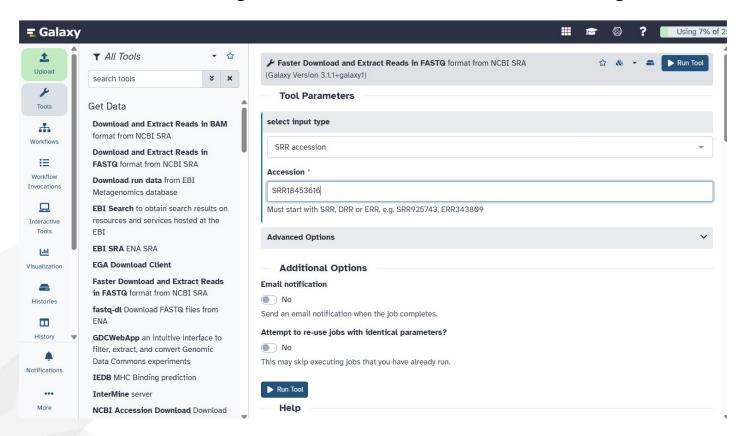
- 9. On the right, click on the dropdown arrow next to Send to:
- 10. Select file under, Choose **Destination**.
- 11. Change the format to **FASTA**
- 12. Click on Create File to download the FASTA File.
- 13. Repeat this process for a gene interesting to you.





#### Obtain **FASTQ** files:

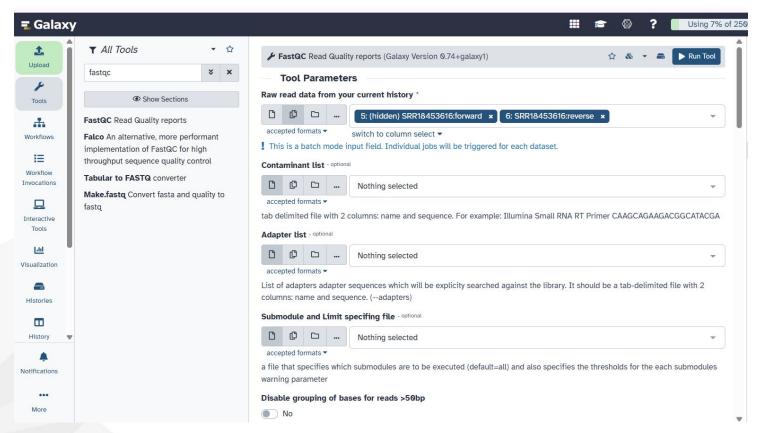
- 1. Go to Galaxy
- 2. In the tool Bar, click on Get Data
- 3. Choose "Faster Download and Extract Reads in FASTA/Q format from NCBI SRA".
- 4. In the Accession tab, write the accession number of the fastq file: SRR18453616.





#### Perform **QC**:

- 1. On the search bar, type **fastqc**.
- 2. **Choose** the desired fastq file (paired end) in the raw read tab.
- 3. Leave all other tabs unchanged.





- 4. Once it runs, two files are generated, a raw data file and a Web Page file.
- 5. **View** the result by clicking on the web page file produced.

#### **Quality Control Result**

#### Summary

- Basic Statistics
- Per base sequence quality
- Per sequence quality scores
- Per base sequence content
- Per sequence GC content
- Per base N content
- Sequence Length Distribution
- Sequence Duplication Levels



# **Quality Control (MultiQC)**

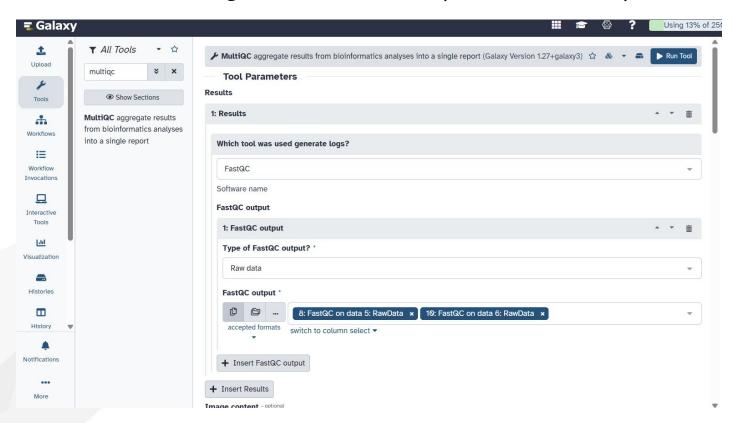
**Why:** It helps us to obtain a more intuitive comparison

- 1. On the search bar, type multiqc
- 2. On the "Which tool was used generate logs?" tab, choose **Fastqc**
- 3. Then click on "Insert FastQC output"

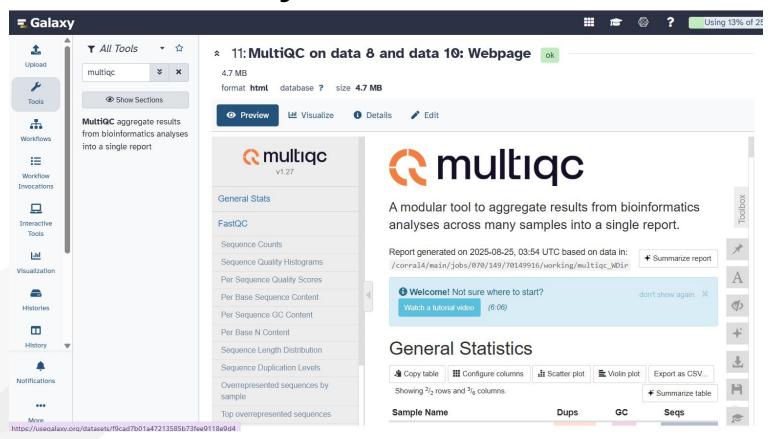
# **Quality Control (MultiQC)**

- 4. Type of output is raw data
- 5. Add the raw data files generated earlier
- 6. **Leave** all other parameters at default
- 7. Run tool
- 8. **View** the result by clicking on the web page file produced

## **Quality Control (MultiQC)**



#### **Quality Control Result**



#### Why QC is Important

- Ensures data integrity before downstream analysis.
- Detects contamination, errors, or poor-quality sequences.
- Prevents misleading results.