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Submitting Genomes

DOI

dx.doi.org/10.17504/protocols.io.8epv5jdkdl1b/v1

Emma Pearce¹

¹UCSF



Emma Pearce

UCSF

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DOI: https://dx.doi.org/10.17504/protocols.io.8epv5jdkdl1b/v1

External link: https://www.ncbi.nlm.nih.gov/WebSub/?form=history&tool=genbank

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Protocol status: In development

We are still developing and optimizing this protocol



Created: February 14, 2023

Last Modified: March 10, 2023

Protocol Integer ID: 77005

Keywords: bankit, submitting, ncbi, genome to ncbi, submitting genomes step, genomes step, ncbi, genome, bankit

Abstract

Steps to submit a genome to NCBI through Bankit

Troubleshooting



Submitting to Bankit

- 1 Log into Bankit with UCSF username and password:
 - Follow this link: https://www.ncbi.nlm.nih.gov/WebSub/?form=history&tool=genbank
 - Choose more login options and select University of California, San Francisco
- 2 Select "Start Bankit Submission"
- 3 Fill out the contact information

4 Sequence Authors:

List all sequence authors (use the Add button for more than one)

Reference Information #1:

Select publication status (unpublished, in-press, or Published)

- If in-press add reference title, journal title, year, volume, issue, and page numbers
- If published add reference title, journal title, year, volume, issue, page numbers, and PubMed ID
 - Optional: if unpublished add reference title

Select reference authors as "Same as Sequence Authors" or "Specify New Authors"

- If specify new authors, list authors

(Optional) Add Another Reference:

Follow Reference Information #1 steps

5 Sequencing Technology:

Select the sequencing technology (or other and specify)

Select unassembled sequence reads or assembled sequences (consisting of two or more sequence reads)

List assembly program and version or date

Example: Illumina, Geneious v. Geneious Prime 2023.0.1

6 **Submission Release Date:**

Select "Immediately after processing" or "Release date"

- If Release Date is selected, specify date in DD-Mon-YYYY format

Sequence(s) and Definition Line(s):

Choose molecule type, linear or circular, if it is a complete sequence, and fasta or alignment format

Either download sequence file or copy and paste it in the box

- Note: make sure the first line follows this format; make everything in bold specific to your submission
 - >Seq#[organism=genus species] species straininfo



Example: >Seq1[organism=Enterovirus rhinovirus] Rhinovirus C17 strain RvC17/USA/2019

7 **Submission Category:**

Select "Original" or "Third Party Annotation"

8 **Source Modifiers:**

Choose organelle/location if applicable

Choose source modifiers and list information in value column, add as many as desired

Example: Strain = RvC17/USA/2019, host = homo sapiens, tissue_type = Lung, country = USA, collection_date = 2019

9 **Features (Overview):**

Select "Add features by completing input forms" Select "Coding Region (CDS) / Gene / mRNA" Select "providing intervals" Click Add

10 Features (Detail):

Choose if the CDS is on the + or - strand

If the CDS is not compete, select 5' or 3' for which end it is incomplete at Select "Entire Sequence" or "Specific Spans" for Nucleotide Interval Spans

- If Specific Spans, specify start and stop and which strand it is on List protein name (optional: also list protein description and EC number) Click accept

Example: specific spans: start = 19 stop = 6498 strand = -minus; protein name = polyprotein

11 **Review Submission:**

Make sure everything looks good

If resubmitting check the box under the resubmission section