	<b>Analyzing HAI Pathogens in Terra using CDC's Phoenix Workflow Version 1</b>		
	Document TG-PX-V1, Version 1		
	Date:	Effective Date:	Workflow Version
	7/20/2023	8/2023	v1

## 1. PURPOSE/SCOPE

To standardize the process of analyzing Healthcare-Associated Infection (HAI) pathogen next generation sequencing (NGS) data using CDC's Phoenix workflow in Terra to generate assemblies, quality control (QC) metrics, and identify and characterize bacterial HAI pathogens for sequence type, antibiotic resistance and hypervirulence genes, and plasmid detection. Acceptable data types include Illumina paired end (PE) raw read files.

## 2. REQUIRED RESOURCES

- Computer
- Internet connection: at least 10 and 5Mbps for download and upload speeds, respectively
- Internet browser
  - Google Chrome, Firefox, or Edge
- Google account
- Terra account, linked to Google account
- Illumina PE raw sequencing read files uploaded to Terra workspace, see [TG-TER-03](#)
- CDC's Phoenix Workflow in Terra, see [TG-TER-03 appendix 9.2](#)

## 3. RELATED DOCUMENTS

Document Number	Document Name
TG-TER-03	Uploading Local or SRA NGS Data & Creating a Results Metadata Table in Terra

## 4. PROCEDURE

### 4.1 CREATE A SAMPLE METADATA FILE (TSV FILE) FOR RAW READS, ASSEMBLIES, AND SRA FETCH

1. In Excel, **create a list**

containing the following sample information:

#### a. For all analyses (Fig 1):

- i. Column 1 header:

**entity:HAI\_id**, where **HAI** is the data table/group of samples to be analyzed

- ii. List all **sample IDs** in column 1


#### b. For analysis from raw sequencing reads (Fig 1):

- i. Column 2 and 3 headers: **read1** and **read2**, respectively

- ii. List the **full file paths** to read1 and read2 files in the cloud

entity:HAI_id	read1	read2	run_id
03-98DDCS	gs://theiagen-public-file	gs://theiagen-public-file	SEQ137
0398KI	gs://theiagen-public-file	gs://theiagen-public-file	SEQ137
Figure 1: Raw Reads Metadata File.		ic-file:gs://theiagen-public-file	SEQ137

entity:HAI_id	assembly_fasta	run_id
03-98DDCS	gs://theiagen-pub	SEQ137
19050801924	gs://theiagen-pub	SEQ137
2022AZMC-0005	gs://theiagen-pub	SEQ137
612031-00282104	gs://theiagen-pub	SEQ137
Figure 2: Assembly Metadata file.		SEQ137

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**c. For analysis from assembly data (Fig 2):**

i. Column 2 header: `assembly_fastq`, or similar

**d. For analysis using SRA fetch to pull read data (Fig 3):**

i. Column 2 header: `sra_accession`, or similar

e. *Optional: remaining columns may be used to add metadata like additional lab results, sample collection information, demographic data, etc*

f. Do not include spaces in the headers

2. `Save as` a txt or tsv file

3. `Upload` to Terra workspace; see [TG-TER-03](#) for details

entity:HAI_id	sra_accession	run_id
03-98DDCS	gs://theiagen-pub	SEQ137
19050801924	gs://theiagen-pub	SEQ137
2022AZMC-0005	gs://theiagen-pub	SEQ137
2022AZMC-0005	gs://theiagen-pub	SEQ137
2022AZMC-0005	gs://theiagen-pub	SEQ137

Figure 3: SRA Accession Metadata File.

## 4.2 RUNNING THE PHOENIX WORKFLOW

1. In Terra, open the `workspace` containing the data of interest and click the `workflows` tab

2. Select the `phoenix` workflow (Fig 4)

3. Choose the latest version of `version 1` in the version dropdown field or the version internally validated (Fig 5, a)

4. Select the second bullet to `run workflow(s) with inputs defined by data table` (Fig 5, b)

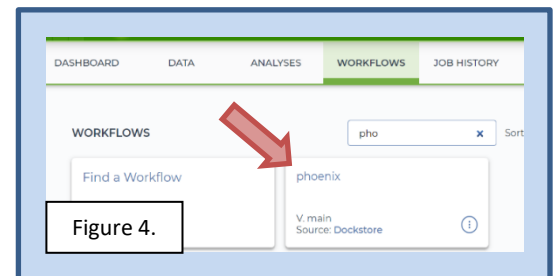
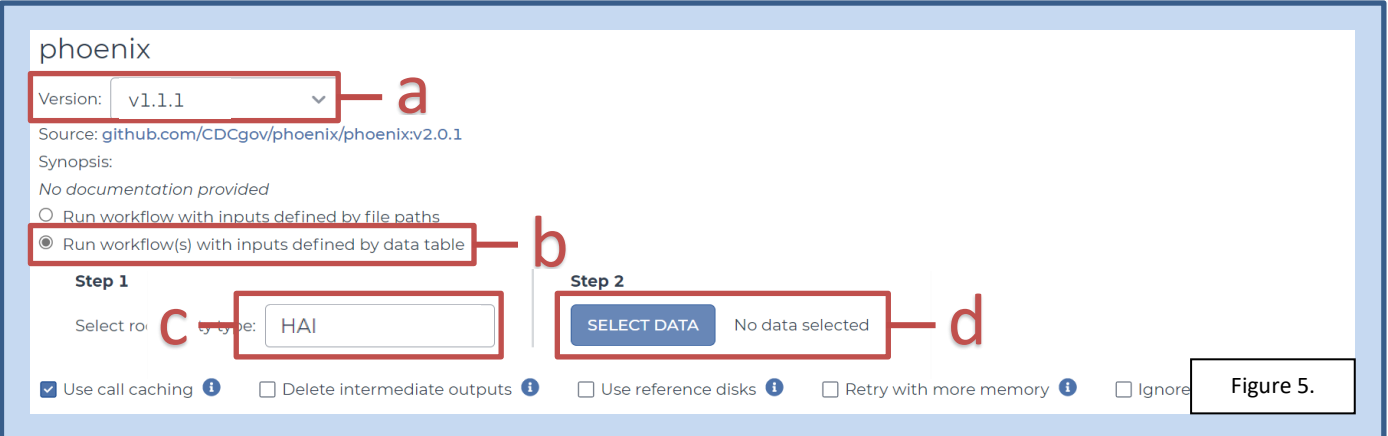


Figure 4.



phoenix

Version: v1.1.1 — a

Source: [github.com/CDCgov/phoenix/phoenix:v2.0.1](https://github.com/CDCgov/phoenix/phoenix:v2.0.1)

Synopsis:

No documentation provided

☐ Run workflow with inputs defined by file paths

☒ Run workflow(s) with inputs defined by data table — b

Step 1

Select root entity type: HAI — c

Step 2

SELECT DATA — d No data selected

☒ Use call caching ☐ Delete intermediate outputs ☐ Use reference disks ☐ Retry with more memory ☐ Ignore

Figure 5.


5. Select the relevant data table under the `select root entity type` dropdown (Fig 5, c)

6. Click `select data` (Fig 5, d)

7. In the pop-up window, `select the checkbox` for each sample to be included in the analysis (Fig 6)

a. *Click the checkbox dropdown and all to select all samples in the data table; if the checkbox at the top is checked, only the first 100 samples in the data table will be selected*

b. *A subset of samples may be chosen using the search bar to filter before selecting the checkbox dropdown and all to select only samples matching the search criteria*

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c. Optional: name the output set name to differentiate this analysis from others, e.g. *Phoenix\_YYYYMMDDn*; this populates a new row to the SET data table

d. Click **ok**

8. In the **inputs** tab, set the first 4 attributes to the following, respectively (Fig 7)

a. **workspace.kraken2\_phoenix**

i. *kraken2\_phoenix* must be uploaded as a workspace data element; see [appendix 10.1](#)

b. **this.read1\***

c. **this.read2\***

i. \*Where **read1** and **read2** are the metadata file column names containing the relevant files ([section 4.1 b](#))

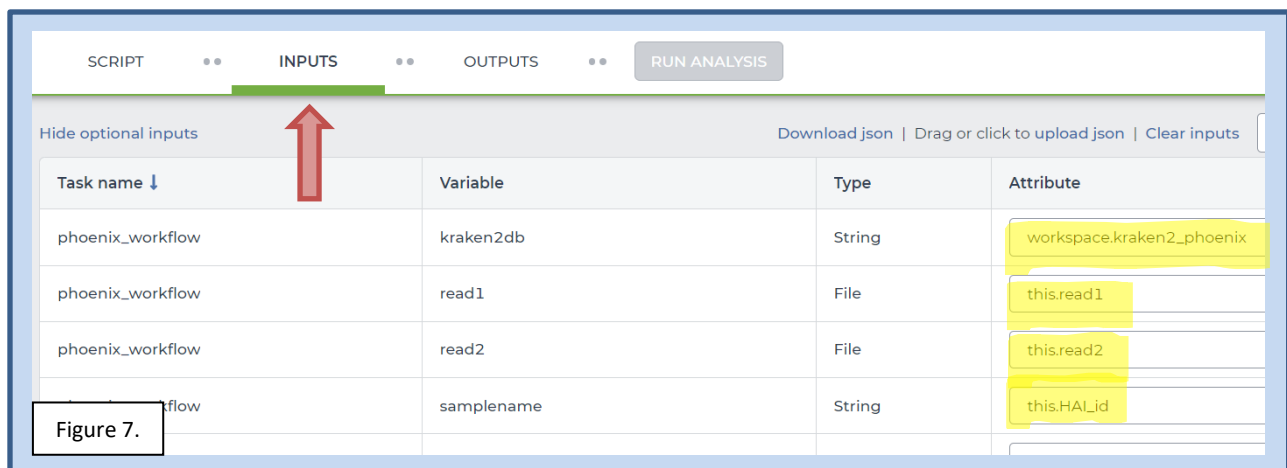
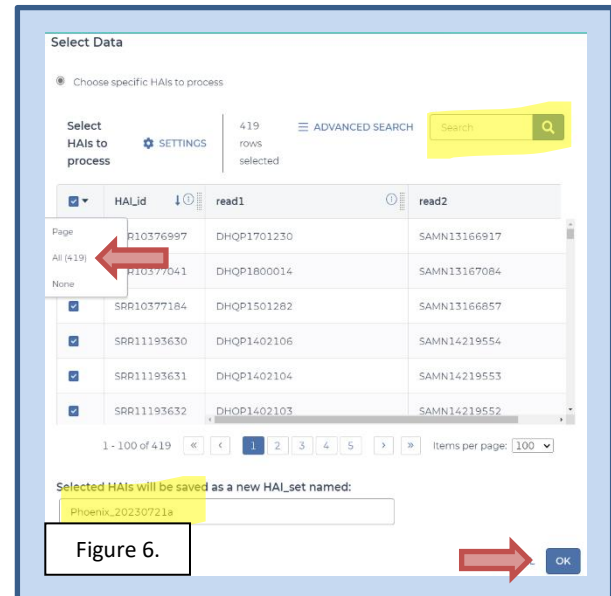
d. **this.HAI\_id**


i. Where **HAI** is the column name in the data table containing sample IDs

9. Specify outputs in the **outputs** tab by clicking **use defaults** (Fig 8)

10. Click **save**

11. Launch the workflow by clicking **run analysis**; enter desired comments and click **launch**



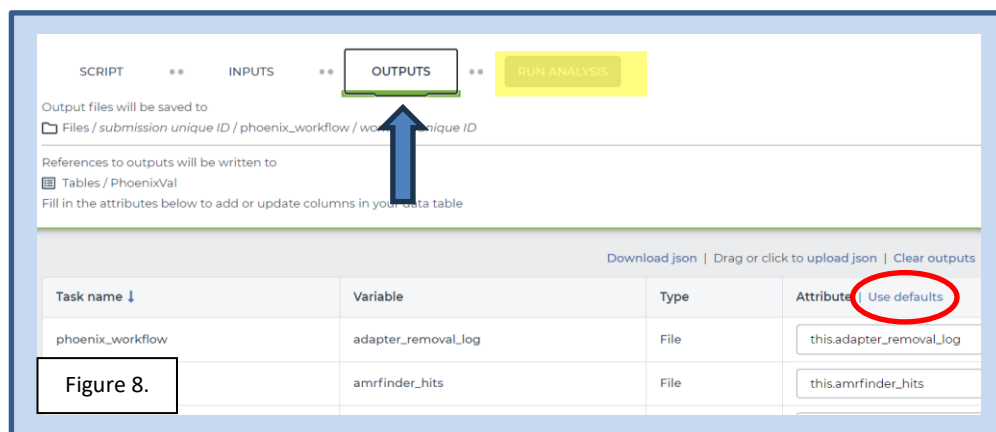
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#### 4.3 DETERMINING:

##### TAXONOMY, AMR CHARACTERIZATION, HYPERVIRULENCE, AND PLASMID MARKERS

1. In the Terra **workspace** containing Phoenix data, navigate to the **data** tab
2. **Open the data table** by clicking on the name of the data table in the left sidebar
3. View **settings** above the data table, select **none** (Fig 9)

- a. Select lab-specific QC metric columns needed to make a sample pass/fail determination




- b. Additionally, select the following result columns: (Fig 9)

- i. **amrfinder\_point\_mutations**
- ii. **beta\_lactam\_resistance\_genes**
- iii. **hypervirulence\_genes**
- iv. **mlst1**
- v. **mlst2**
- vi. **mlst1\_scheme\_1**
- vii. **mlst2\_scheme\_2**
- viii. **other\_ar\_genes**
- ix. **plasmid\_incompatability\_replicons**
- x. **species**

1. **Optional:** save this column group for future use by clicking the **save this column selection** field, **naming it** (e.g. *PhoenixResults*), and clicking **save**

- c. Click **done**
4. Determine the predicted taxonomy, sequence type, and AMR, hypervirulence, and plasmid characterization for each sample by viewing the corresponding columns
  5. Follow lab-specific QC assessment, resulting, and reporting procedures, as applicable

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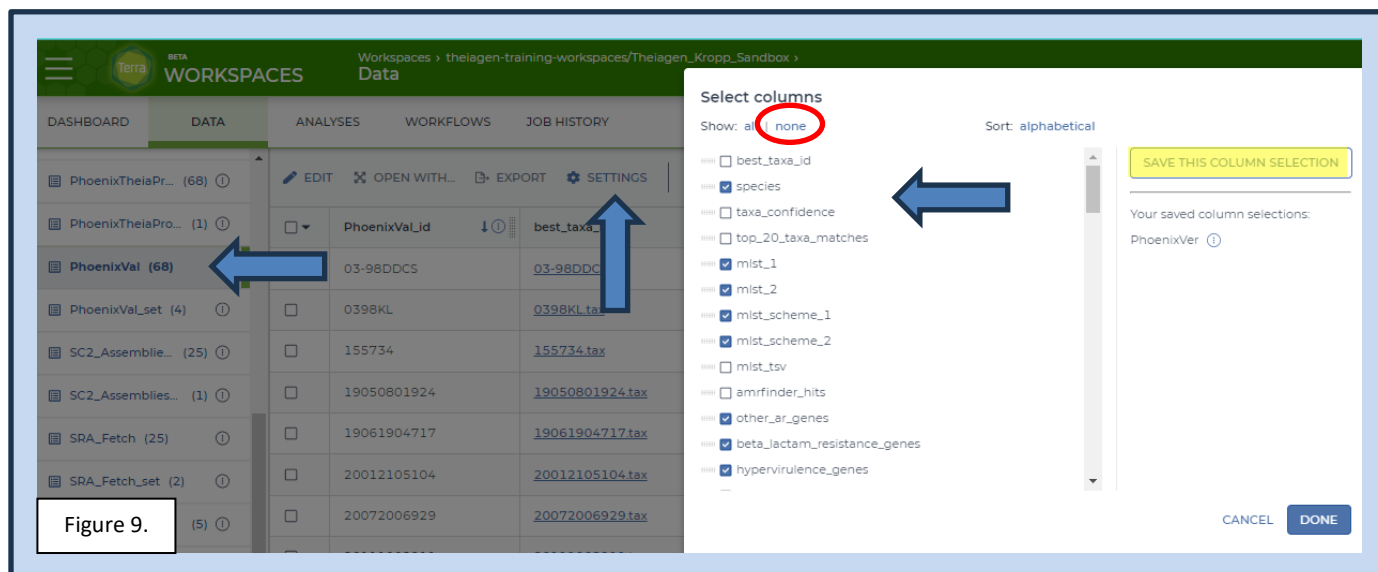


Figure 9.

## 5. QUALITY RECORDS

1. Raw reads
2. Metadata (tsv)
3. All Phoenix workflow outputs relevant to results, including tool & database versions

## 6. TROUBLESHOOTING

- Consult with internal staff familiar with this procedure or contact [support@theiagen.com](mailto:support@theiagen.com) for troubleshooting inquiries
- For document edit requests, contact [support@theiagen.com](mailto:support@theiagen.com)

## 7. INTERFERENCES


N/A

## 8. REFERENCES

None

## 9. REVISION HISTORY

Revision	Version	Release Date
Document creation	1	7/2023

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## 10. APPENDICES

### 10.1 ADD A WORKSPACE DATA ELEMENT

1. Navigate to the **Terra workspace** where Phoenix will be run
2. To upload local files, open the **Files** tab in the bottom left of the workspace (Fig 8)
  - a. Click **upload**
  - b. Once the upload is complete, **right click** on the file name and click **copy link**

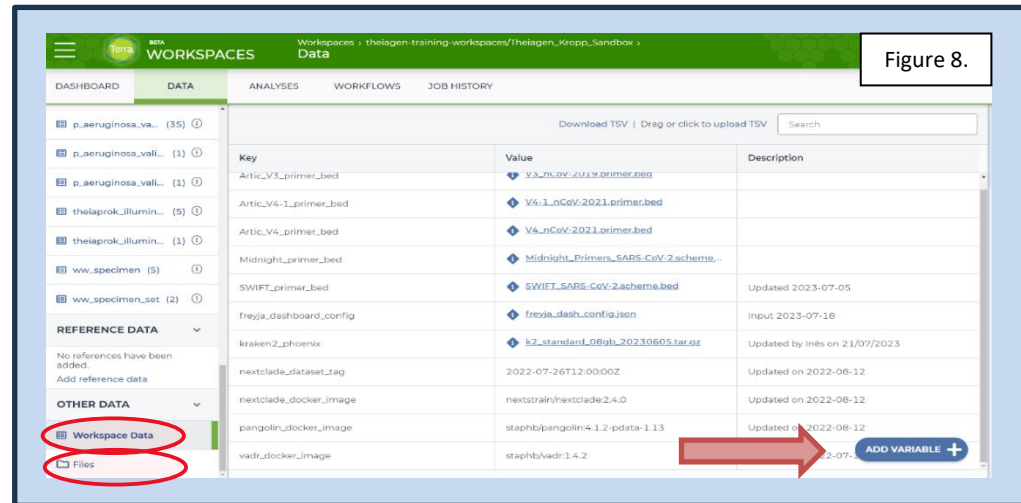


Figure 8.

3. Open the **workspace data** tab and click the **blue plus symbol** in the bottom right (Fig 8)
4. Click in the **key field** and **name the element** being added (Fig 9)
  - a. E.g. to add the Kraken2 database, the key **kraken2\_phoenix** may be used to specify its use with the Phoenix workflow
5. In the value field, choose **string** as the value type
  - a. **Paste the file path**
    - i.E.g. for the kraken2 database, paste **gs://theiagen-public-files-rp/terra/theiaprok-files/k2\_standard\_08gb\_20230605.tar.gz**
    - b. For other string elements like docker images and dataset tags, **paste the ID value**
      - i.E.g. for the nextclade docker image, add **nextstrain/nextclade:2.13.0**
      - ii.Always ensure the docker images and dataset tags are aligned with versions used for internal validation procedures


Key	Value	Description
Artic_V3_primer_bed	 <a href="#">V3_nCoV-2019.primer.bed</a>	
nextclade_dataset_tag	2022-07-26T12:00:00Z	Updated on 2022-08-12
nextclade_docker_image	nextstrain/nextclade:2.4.0	Updated on 2022-08-12
pangolin_docker_image	staphb/pangolin:4.1.2-pdata-1.13	Updated on 2022-08-12
vadr_docker_image	staphb/vadr:1.4.2	Updated on 2022-07-15
kraken2_phoenix	gs://theiagen-publi	String Updated on 7/24/2023

Figure 9.