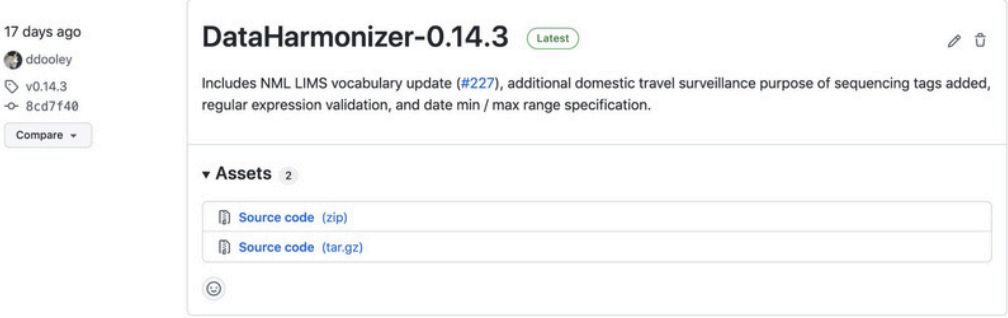
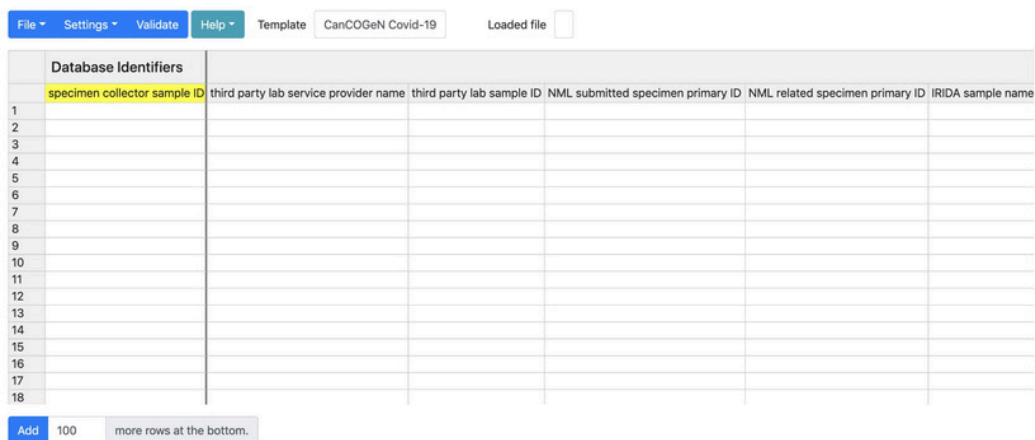


## Contextual Data (Metadata) Curation

- I. Purpose: To harmonize SARS-CoV-2 contextual data across data providers in the CanCOGeN network.
- Data providers will extract and curate lab-specific contextual data according to the steps outlined in the procedure below.
  - Laboratories will populate the harmonized template with information from their datasets using the *DataHarmonizer* application.
  - Data providers will share the harmonized data with the national database according to the agreed upon mechanism.
- II. Data: The contextual data describing sample collection and processing, host information, sequencing, and bioinformatics and QC metrics as supplied by the data provider.
- III. Procedure:

	Action	Related docs
1	<p>Download the zip file (“Source code (zip)”) containing The DataHarmonizer application from the following link: <a href="https://github.com/Public-Health-Bioinformatics/covid19ValidationGrid/releases">https://github.com/Public-Health-Bioinformatics/covid19ValidationGrid/releases</a></p>  <p>Extract the zip file’s contents, and navigate into the extracted folder. Open main.html. The validator application will open in your default browser. It should look like this:</p>	

	Action	Related docs
--	--------	--------------



Database Identifiers						
	specimen collector sample ID	third party lab service provider name	third party lab sample ID	NML submitted specimen primary ID	NML related specimen primary ID	IRIDA sample name
1						
2						
3						
4						
5						
6						
7						
8						
9						
10						
11						
12						
13						
14						
15						
16						
17						
18						

Add 100 more rows at the bottom.

Data can be entered into the validator application manually, by typing values into the application's spreadsheet, or data can be imported from localxlsx, xls, tsvandcsvfiles.

To import local data, clickFileon the top-left toolbar, and then clickOpen. To enter data in a new file, clickFileon the top-lefttoolbar, and then clickNew. Data entered into the spreadsheet can be copied and pasted.

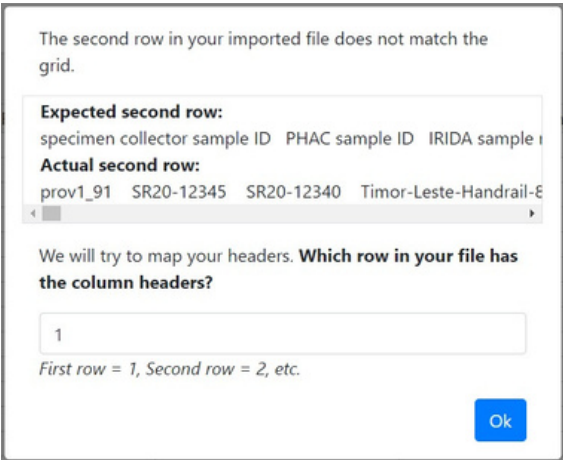
*Note: Only files containing the headers expected by the DataHarmonizer can be opened in the application. Example:*

Database Identifiers					
specimen colle	NML submitte	NML related s	IRIDA sample	umbrella biop	bio
prov1_91					
prov1_92					
prov1_93					
nrov1_94					

*If you are missing the first row, you will get the following warning:*

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	Action	Related docs
	 <p>Resolve by declaring “1” as the row in which your column headers reside.</p>	
2	<p>Before you begin to curate sample metadata:</p> <ul style="list-style-type: none"> <li>● Review your dataset</li> <li>● Review the fields in the template of the Validator application</li> <li>● Review the field descriptions in the SOP Appendix</li> </ul>	
3	<p>Familiarize yourself with DataHarmonizer functionality by reviewing the “Getting Started”. To access "Getting Started", click on the green Help button on the top-left toolbar, then click Getting Started. Definitions, examples and further guidance are available by double clicking on the field headers, or by using the “Reference Guide”. To access the “Reference Guide” click on the Help button, then click Reference Guide.</p>	
4	<p>Confirm mapping of your data fields to those in the harmonized template with the data steward (e.g. your supervisor).</p> <p><i>Note: A version of this information will be made public in GISAID and NCBI, however, another version of this data will be captured in the access controlled national database. Confirm the level of granularity of information that can be shared publicly vs in the national database, with the data steward and/or the privacy officer. The most detailed information allowable should be included here.</i></p>	
5	<p>Enter data into the validator spreadsheet.</p> <ul style="list-style-type: none"> <li>● Hide non-required fields (colour-coded purple and white/grey) by clicking Settings on the top-left toolbar, followed by clicking on Show Required Columns (colour-coded in yellow).</li> <li>● Double click in the field headers to see definitions and detailed guidance as needed (or consult Appendix A).</li> </ul>	

	Action	Related docs						
	<div><ul style="list-style-type: none"><li>● Jump to a specific field header by clicking <b>Settings</b> on the top-left toolbar, followed by clicking on <b>Jump to</b>, then select the field header of the column you would like to view from the drop down list.</li><li>● Populate the validator template with the information from your dataset.</li><li>● Use picklists when provided.</li><li>● A value must be entered for every <u>required field</u> in each row. If data is missing or not collected, choose a null value from the picklist.<ul style="list-style-type: none"><li><input type="radio"/> Not Applicable</li><li><input type="radio"/> Missing</li><li><input type="radio"/> Not Collected</li><li><input type="radio"/> Not Provided</li><li><input type="radio"/> Restricted Access</li></ul></li><li>● Free text can be provided when picklists are not available.</li><li>● For filling an entire column with the same data, use the <b>Fill Column</b> function. Click <b>Settings</b>, followed by <b>Fill Column</b>. Type in the name of the desired field, followed by the value that should be used to fill every row in that column. Then click <b>OK</b>.</li></ul><p><b>If a desired term is not present in a picklist, contact Emma Griffiths at <a href="mailto:ega12@sfu.ca">ega12@sfu.ca</a>.</b></p><p><i>Note: Sometimes there will be constraints on what information can be shared, other times a field may not be applicable to your sample. Use the null values (controlled vocabulary indicating the reason why information is not provided) in the picklist to report missing data.</i></p><p>Required fields are organized into subsections (see Appendix A for required field definitions and guidance, and Appendix B for examples of how to structure sample descriptions):</p><table><tr><th>Subsection</th><th>Required Fields</th></tr><tr><td><b>Database Identifiers Sample</b></td><td>specimen collector sample ID</td></tr><tr><td><b>Collection and Processing</b> <i>Note: Evaluate with your supervisor whether the specimen collector sample ID is considered identifiable by your institutional policies. If not, copy the sample ID into the sample ID field in the validator spreadsheet.</i></td><td>sample collected by sequence submitted by sample collection date sample collection date precision geo_loc (country) geo_loc (province/territory) organism isolate</td></tr></table></div>	Subsection	Required Fields	<b>Database Identifiers Sample</b>	specimen collector sample ID	<b>Collection and Processing</b> <i>Note: Evaluate with your supervisor whether the specimen collector sample ID is considered identifiable by your institutional policies. If not, copy the sample ID into the sample ID field in the validator spreadsheet.</i>	sample collected by sequence submitted by sample collection date sample collection date precision geo_loc (country) geo_loc (province/territory) organism isolate	
Subsection	Required Fields							
<b>Database Identifiers Sample</b>	specimen collector sample ID							
<b>Collection and Processing</b> <i>Note: Evaluate with your supervisor whether the specimen collector sample ID is considered identifiable by your institutional policies. If not, copy the sample ID into the sample ID field in the validator spreadsheet.</i>	sample collected by sequence submitted by sample collection date sample collection date precision geo_loc (country) geo_loc (province/territory) organism isolate							

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	Action	Related docs
	<p><i>If yes, provide the alternative sample ID as specified by the lab. Be sure to keep a copy of the key.</i></p>	<p>purpose of sampling purpose of sampling details</p>
	<p>Describing the material and/or site sampled.</p> <p><i>Note: Seven fields have been introduced to capture different kinds of anatomical and environmental samples, as well as collection methods. Populate only the fields that pertain to your sample - provide null values for the fields that are not applicable. Provide the most granular information allowable according to your organization's data sharing policies. NML submitted specimen type is required for upload to CNPHI. Select the appropriate value from the available pick list (consult the reference guide for further support).</i></p>	<p>anatomical material anatomical part body product environmental material environmental site collection device collection_method NML submitted specimen type</p>
	<p><b>Host Information</b></p>	<p>host (scientific name) host disease host age host age unit host age bin host gender</p>
	<p><b>Sequencing</b></p>	<p>sequencing instrument sequencing date purpose of sequencing purpose of sequencing details</p>
	<p><b>Bioinformatics and QC Metrics</b></p>	<p>raw sequencing data processing method hosting method consensus sequence software name consensus sequence software version bioinformatics protocol</p>

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	Action	Related docs
6	<p>Validate the entered data by clicking on the <b>Validate</b> button on the top-left toolbar.</p> <p>Missing information and invalid entries in required fields will be highlighted in red.</p> <ul style="list-style-type: none"> <li>● Observe invalid rows by clicking <b>Settings</b> in the top-left toolbar, and then clicking on <b>Show invalid rows</b>.</li> <li>● Address errors systematically by clicking the <b>Next Error</b> button. When all errors have been corrected, the <b>Next Error</b> button will disappear.</li> <li>● Observe valid rows by clicking <b>Settings</b> in the top-left toolbar, and then clicking on <b>Show valid rows</b>.</li> <li>● Return view to all rows by clicking <b>Settings</b> in the top-left toolbar, and then clicking on <b>Show all rows</b>.</li> </ul> <p><i>Note: Row viewing options only appear after a validation attempt has been made.</i></p>	
7	<p>Address any invalid data that was flagged in red in the template.</p> <ul style="list-style-type: none"> <li>●  Pale Red = Incorrect data format</li> <li>●  Dark Red = Required data missing</li> </ul> <p><i>Note: It is possible to export incomplete or invalid data. Make sure to review any errors prior to exporting.</i></p>	
8	<p>Export validated data by clicking <b>File</b> on the top-left toolbar, and then clicking on <b>Save as</b>. Enter the file name and press <b>Save</b>. Export to IRIDA, GISAID, or NML-LIMS formats by clicking <b>File</b> on the top-left toolbar, and then clicking <b>Export to</b>.</p> <ul style="list-style-type: none"> <li>● Have the validated data reviewed by the data steward (i.e. your supervisor)</li> </ul>	
9	<p>Submit validated data to the national database. You can submit either by i) emailing the validated data to your NML contact, or</p> <p>ii) uploading the validated data directly through the CNPHI Collaboration Centre interface.</p> <ul style="list-style-type: none"> <li>● Before uploading to CNPHI, export your data in “NML-LIMS” format by clicking <b>File</b> on the top-left toolbar, then clicking <b>Export To</b>. Type in the file name, and select “<b>NML-LIMS</b>” from the Format picklist. Then click <b>Export</b>.</li> <li>● See CNPHI documentation for more information regarding Metadata Upload.</li> </ul>	<p>Submission to CNPHI LaSER SOP:  <a href="https://docs.google.com/document/d/e/2PACX-1vRUAWGPP44_Dlbmpx8Fi_mS-iweA2d3KIBGs_h2JgUiWzKIXT_O4gz44fCbqqH1JA_nprWsRaFHkjZMKj/pub">https://docs.google.com/document/d/e/2PACX-1vRUAWGPP44_Dlbmpx8Fi_mS-iweA2d3KIBGs_h2JgUiWzKIXT_O4gz44fCbqqH1JA_nprWsRaFHkjZMKj/pub</a></p>

## CanCOGeN\_2.5 Contextual Data Curation

	Action	Related docs
10	<p>Optional: Format validated data for GISAID submission. The DataHarmonizer will automate the preparation of a GISAID submission form from the entered data by exporting the data in GISAID format.</p> <ul style="list-style-type: none"> <li>● Export your data in “GISAID” format by clicking <b>File</b> on the top-left toolbar, then clicking <b>Export To</b>. Type in the filename, and select “GISAID” from the Format picklist. Then click <b>Export</b>.</li> </ul>	
11	<p>Optional: Format validated data for IRIDA submission. The DataHarmonizer will automate the preparation of an IRIDA submission form from the entered data by exporting the data in IRIDA format.</p> <ul style="list-style-type: none"> <li>● Export your data in “IRIDA” format by clicking <b>File</b> on the top-left toolbar, then clicking <b>Export To</b>. Type in the file name, and select “IRIDA” from the Format picklist. Then click <b>Export</b>.</li> </ul> <p><i>Note: If the top row containing the broad headings (Database identifiers, Sample collection and processing, Host information, Sequencing, Bioinformatics and QC, Authors) is not removed, the IRIDA metadata upload will fail. By exporting the sheet in IRIDA format, the DataHarmonizer completes this formatting for you.</i></p>	<p>Upload to IRIDA SOP:</p> <p><a href="https://irida.cerifacility.ca/documentation/user/user/samples/#adding-a-new-sample">https://irida.cerifacility.ca/documentation/user/user/samples/#adding-a-new-sample</a></p>
12	<p>Additional Information:</p> <p>A local copy of the <b>Standard Operating Procedure (SOP)</b> is included in every download of the DataHarmonizer. To access it, click on the green <b>Help</b> button on the top-left toolbar, then click <b>SOP</b>.</p> <p>The latest version of the SOP is <a href="#">published online</a> and accessible via a web browser at all times.</p> <p>Datasets that can be used for testing, training, and quality control purposes are also available.</p>	<p>Testing/training datasets:</p> <p><a href="https://github.com/cidgoh/DataHarmonizer/wiki/CanCOGeN-Contextual-Data-Datasets-for-Harmonization-Training-and-Quality-Control">https://github.com/cidgoh/DataHarmonizer/wiki/CanCOGeN-Contextual-Data-Datasets-for-Harmonization-Training-and-Quality-Control</a></p>

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IV. Appendix A: Required Field Definitions and Guidance

Field definitions for required fields, as well as guidance and examples, are provided below. This information has been sourced from the DataHarmonizer reference guide. Guidance for strongly recommended and optional fields can be found in the reference guide. For access to information on non-required fields, refer to “Procedure - Action 3”.

Database Identifiers specimen collector sample ID

*The user-defined name for the sample.*

Store the collector sample ID. If this number is considered identifiable information, provide an alternative ID. Make sure to store the key between this alternative ID and the original ID for traceability. Every collector sample ID from a single submitter must be unique. It can have any format, but we suggest that you make it concise, unique and consistent within your lab.

e.g. prov\_rona\_99

Sample Collection and Processing

sample collected by

*The name of the agency that collected the original sample.*

The name of the sample collector should be written out in full, (with minor exceptions) and be consistent across multiple submissions e.g. Public Health Agency of Canada, Public Health Ontario, BC Centre for Disease Control. The sample collector specified is at the discretion of the data provider (i.e. may be hospital, provincial public health lab, or other).

e.g. BC Centre for Disease Control

sequence submitted by

*The name of the agency that generated the sequence.*

The name of the agency should be written out in full, (with minor exceptions) and be consistent across multiple submissions. If submitting specimens rather than sequencing data, please put the "National Microbiology Laboratory (NML)". e.g. Public Health Ontario (PHO)

sample collection date

*The date on which the sample was collected.*

Sample collection date is critical for surveillance and many types of analyses. Required granularity includes year, month and day. Record the collection date accurately in the template. Before sharing this data, ensure you have consulted the data steward and/or your privacy officer regarding whether they consider this date to be identifiable information. If this date is considered identifiable, it is acceptable to add "jitter" to the collection date you share by adding or subtracting a calendar day (acceptable by GISAID). Do not change the collection date in your original records. Alternatively, "received date" may be used as a substitute in the data you share. The date should be provided in ISO 8601 standard format "YYYY-MM-DD". e.g. 2020-03-16



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sample collection date precision

*The precision to which the "sample collection date" was provided.*

Provide the precision of granularity to the "day", "month", or "year" for the date provided in the "sample collection date" field. The "sample collection date" will be truncated to the precision specified upon export; "day" for "YYYY-MM-DD", "month" for "YYYY-MM", or "year" for "YYYY".  
e.g. year

geo\_loc\_name (country)

*The country where the sample was collected.*

Provide the country name from the controlled vocabulary provided.

e.g. Canada

geo\_loc\_name (province/territory)

*The province/territory where the sample was collected.*

Provide the province/territory name from the controlled vocabulary provided.

e.g. Saskatchewan

organism

*Taxonomic name of the organism.*

Use Severe acute respiratory syndrome coronavirus 2. This value is provided in the template.

e.g. Severe acute respiratory coronavirus 2

isolate

*Identifier of the specific isolate.*

Provide the isolate name. The isolate name should be identical to the GISAID virus name in the format "hCov-19/CANADA/xxxxx/2020", where xxxxx represents the sample ID.

e.g. hCov-19/CANADA/prov\_rona\_99/2020

purpose of sampling

*The reason that the sample was collected.*

The reason a sample was collected may provide information about potential biases in sampling strategy. Provide the purpose of sampling from the picklist in the template. Most likely, the sample was collected for Diagnostic testing. The reason why a sample was originally collected may differ from the reason why it was selected for sequencing, which should be indicated in the "purpose of sequencing" field.

e.g. Diagnostic testing

purpose of sampling details

*The description of why the sample was collected providing specific details.*

Provide an expanded description of why the sample was collected using free text. The description may include the importance of the sample for a particular public health investigation/surveillance activity/research question. If details are not available, provide a null value.

e.g. The sample was collected to investigate the prevalence of variants associated with mink-to-human transmission in Canada.

Describing the material and/or site sampled.

anatomical material

*A substance obtained from an anatomical part of an organism e.g. tissue, blood.*

Provide a descriptor if an anatomical material was sampled. Use the picklist provided in the template. If a desired term is missing from the picklist, contact [emma.griffiths@bccdc.ca](mailto:emma.griffiths@bccdc.ca). If not applicable, do not leave blank. Choose a null value.

e.g. Blood

anatomical part

*An anatomical part/location of an organism e.g. oropharynx.*

Provide a descriptor if an anatomical part was sampled. Use the picklist provided in the template. If a desired term is missing from the picklist, contact [emma.griffiths@bccdc.ca](mailto:emma.griffiths@bccdc.ca). If not applicable, do not leave blank. Choose a null value.

e.g. Nasopharynx (NP)

body product

*A substance excreted/secreted from an organism e.g. feces, urine, sweat.*

Provide a descriptor if a body product was sampled. Use the picklist provided in the template. If a desired term is missing from the picklist, contact [emma.griffiths@bccdc.ca](mailto:emma.griffiths@bccdc.ca). If not applicable, do not leave blank. Choose a null value.

e.g. Feces

environmental material

*A substance or object obtained from the natural or man-made environment e.g. soil, water, sewage.*

Provide a descriptor if an environmental material was sampled. Use the picklist provided in the template. If a desired term is missing from the picklist, contact [emma.griffiths@bccdc.ca](mailto:emma.griffiths@bccdc.ca). If not applicable, do not leave blank. Choose a null value.

e.g. Face Mask

environmental site

*An environmental location may describe a site in the natural or built environment e.g. contact surface, metal can, hospital, wet market, bat cave.*

Provide a descriptor if an environmental site was sampled. Use the picklist provided in the template. If a desired term is missing from the picklist, contact [emma.griffiths@bccdc.ca](mailto:emma.griffiths@bccdc.ca). If not applicable, do not leave blank. Choose a null value.

e.g. Building floor

collection device

*The instrument or container used to collect the sample e.g. swab.*

Provide a descriptor if a device was used for sampling. Use the picklist provided in the template. If a desired term is missing from the picklist, contact [emma.griffiths@bccdc.ca](mailto:emma.griffiths@bccdc.ca). If not applicable, do not leave blank. Choose a null value.

e.g. Swab

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collection method

*The process used to collect the sample e.g. phlebotomy, necropsy.*

Provide a descriptor if a collection method was used for sampling. Use the picklist provided in the template. If a desired term is missing from the picklist, contact emma.griffiths@bccdc.ca. If not applicable, do not leave blank. Choose a null value.

e.g. Bronchoalveolar Lavage (BAL)

NML submitted specimen type

*The type of specimen submitted to the NML for testing.*

This information is required for upload through the CNPHI LaSER system. Select the specimen type from the pick list provided. If sequence data is being submitted rather than a specimen for testing, select “Not Applicable”.

e.g. Swab

Host Information

host (scientific name)

*The taxonomic, or scientific name of the host.*

Common name or scientific name are required if there was a host. Both can be provided, if known. Use terms from the pick lists in the template. Scientific name e.g. Homo sapiens, If the sample was environmental, put “Not Applicable”.

e.g. Homo sapiens

host disease

*The name of the disease experienced by the host.*

This field is only required if there was a host. If the host was a human select COVID-19 from the pick list. If the host was asymptomatic, this can be recorded under “host health state details”. If the host is not human, and the disease state is not known or the host appears healthy, put “Not Applicable”.

e.g. COVID-19

host age

*Age of host at the time of sampling.*

Enter the age of the host in years. If not available, provide a null value. If there is not host, put “Not Applicable”.

e.g. 79

host age bin

*Age of host at the time of sampling, expressed as an age group.*

Select the corresponding host age bin from the pick list provided in the template. If not available, provide a null value. The “host age bin” field will automatically propagate with the bin that corresponds to the input in “host age”. If not available or you are not permitted to share, put a null value.

Age Bins:

0 - 9

10 -19

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20 -29  
30 -39  
40 -49  
50 -59  
60 -69  
70 -79  
80 -89  
90 -99  
100+

host age unit

*The unit used to measure the host age, in either months or years.*

Indicate whether the host age is in months or years. Age indicated in months will be binned to the 0 - 9 year age bin.

host gender

*The gender of the host at the time of sample collection.*

Select the corresponding host gender from the pick list provided in the template. If not available, choose a null value. If there is no host, put "Not Applicable".

e.g. Male

### Sequencing

sequencing instrument

*The model of the sequencing instrument used.*

Select a sequencing instrument from the picklist provided in the template.

e.g. MinIon

sequencing date

*The date the sample was sequenced.*

Provide the date that the sample was sequenced in ISO 8601 standard "YYYY-MM-DD" format.

If the exact sequencing date is unknown, proxy dates may be used instead (e.g. library preparation date)

e.g. 2020-06-22

purpose of sequencing

*The reason that the isolate was sequenced.*

The reason an isolate was sequenced may provide information about potential biases in sequencing strategy. Provide the purpose of sequencing from the picklist in the template. Most likely, the sample was collected for Surveillance or Research. The reason why a sample was originally collected may differ from the reason why it was selected for sequencing. The reason for sample collection should be indicated in the "purpose of sampling" field.

purpose of sequencing details

*The description of why the sample was sequenced providing specific details.*

Provide an expanded description of why the sample was sequenced using free text. The description may include the importance of the sequences for a particular public health

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investigation/surveillance activity/research question. If details are not available, provide a null value.

e.g. The sample was sequenced to investigate the differences in lineages circulating in Canada during the spring and fall waves of the pandemic.

Bioinformatics and QC Metrics

raw sequencing data processing method

*The names of the software and version number used for raw data processing such as removing barcodes, adapter trimming, filtering etc.*

Provide the software name followed by the version.

e.g. Trimmomatic v. 0.38, Porechop v. 0.2.3

dehosting method

*The method used to remove host reads from the pathogen sequence.*

Provide the name and version number of the software used to remove host reads.

e.g. BWA 0.7.17

consensus sequence software name

*The name of software used to generate the consensus sequence.*

Provide the name of the software used to generate the consensus sequence.

e.g. iVar

consensus sequence software version

*The version of the software used to generate the consensus sequence.*

Provide the version of the software used to generate the consensus sequence.

e.g. 1.3

bioinformatics protocol

*A description of the overall bioinformatics strategy used.*

Further details regarding the methods used to process raw data, and/or generate assemblies, and/or generate consensus sequences can. This information can be provided in an SOP or protocol or pipeline/workflow. Provide the name and version number of the protocol, or a GitHub link to a pipeline or workflow.

e.g. <https://github.com/phac-nml/ncov2019-artic-nf>

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## V. Appendix B: Structuring Sample Descriptions (Examples)

Several examples are provided below which illustrate how to structure common sample descriptions.

*e.g. nasal swab* should be recorded:

host (scientific name)	host (common name)	host disease	anatomical part	collection device
Homo sapiens	Human	COVID-19	Nasopharynx (NP)	Swab

*e.g. throat swab* should be recorded:

host (scientific name)	host (common name)	host disease	anatomical part	collection device
Homo sapiens	Human	COVID-19	Oropharynx (OP)	Swab

*e.g. saliva* should be recorded:

host (scientific name)	host (common name)	host disease	anatomical material
Homo sapiens	Human	COVID-19	Saliva

*e.g. salt water gargle* should be recorded:

host (scientific name)	host (common name)	host disease	collection method
Homo sapiens	Human	COVID-19	Saline gargle (mouth rinse and gargle)

*e.g. human feces* should be recorded:

host (scientific name)	host (common name)	host disease	body product
Homo sapiens	Human	COVID-19	Feces

*e.g. swab of a hospital bed rail* should be recorded:

environmental site	environmental material	collection device
Hospital	Bed Rail	Swab

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*e.g. tissue from a bat (Platyrrhinus lineatus) in a cave* should be recorded:

Host (common name)	Host (scientific name)	host disease	anatomical_part	environmental_site
Bat	Platyrrhinus lineatus	Not applicable	Tissue	Cave

*e.g. particulates from air filter* should be recorded:

environmental material	collection method
Particulate Matter	Air Filtration

## VI. Appendix C: Null Value Definitions

### Not Applicable

Information is inappropriate to report, can indicate that the standard itself fails to model or represent the information appropriately.

### Missing

Information was known to be recorded in the past, but the observed value cannot be located or retrieved for some reason.

### Not Collected

Information of an expected format was not given because it has not been collected.

### Not Provided

Information of an expected format was not given, a value may be given at the later stage.

### Restricted Access

Information exists but can not be released openly because of privacy concerns.

Source:

International Nucleotide Sequence Database Collaboration (INSDC) Missing Value Reporting Terms (2017-2018).  
ENA Training Modules : <https://ena-docs.readthedocs.io/en/latest/submit/samples/missing-values.html>

CanCOGeN – SARS-CoV-2  
CanCOGeN\_2.5 Contextual Data Curation

Revision History

Version	Date	Writer	Description of Change
0.0	May 25, 2020	Lauren Tindale, Emma Griffiths	Created protocol
1.0	June 8, 2020	Emma Griffiths	Protocol edited
1.1	June 16, 2020	Emma Griffiths	Protocol edited
1.2	October 05, 2020	Rhiannon Cameron	Protocol edited
1.3	October 06, 2020	Rhiannon Cameron	Protocol edited
1.4	October 23, 2020	Rhiannon Cameron	Protocol & Appendix A edited
1.5	November 04, 2020	Rhiannon Cameron, Emma Griffiths	Protocol & Appendix A edited
1.6	November 12, 2020	Emma Griffiths, Rhiannon Cameron	Protocol & Appendix A edited
1.7	November 13, 2020	Rhiannon Cameron	Protocol edited
1.8	December 11, 2020	Emma Griffiths	Protocol & Appendix A edited
1.9	December 18, 2020	Emma Griffiths	Appendix A edited
2.0	March 5, 2021	Emma Griffiths	Protocol & Appendices edited
2.1	April 26, 2021	Rhiannon Cameron	Appendix A edited
2.2	November 8, 2021	Emma Griffiths	Required fields updated, extra instructions describing new functionality added
2.3	February 2 2022	Emma Griffiths	Link to testing/training datasets added
2.5	December 2024	Charlie Barclay	Version compatibility update