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Oct 18, 2021

# DRAGEN COVID Lineage App SARS-CoV-2 Strain Characterization on the Illumina BaseSpace Platform V.1



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**Technical Outreach and Assistance for States Team<sup>1</sup>**<sup>1</sup>Centers for Disease Control and Prevention

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This protocol provides instructions on how to run the DRAGEN COVID Lineage app on the Illumina BaseSpace Sequence Hub. The DRAGEN COVID Lineage app reads in fastq sequence files and produces Lineage, Clade, and Kmer Detection reports for each run. The information provided in these reports include Pangolin lineage classification, variant calls, and the fraction of human and SARS-CoV-2 kmers detected. The DRAGEN COVID Lineage app also assembles a consensus sequence fasta file for each sample. This document applies to all whole-genome sequencing runs on the Illumina platform and downstream bioinformatics for public health laboratories.

For technical assistance, please contact: **TOAST@cdc.gov**

Technical Outreach and Assistance for States Team 2021. DRAGEN COVID Lineage App SARS-CoV-2 Strain Characterization on the Illumina BaseSpace Platform.

**protocols.io**

<https://protocols.io/view/dragen-covid-lineage-app-sars-cov-2-strain-characterization-on-the-illumina-basespace-platform-v1>



## (ILLUMINA MENU) Protocols for SARS-CoV-2 Library Prep with ARTIC Primers, Bioinformatic Analysis, and Database Submission

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Feb 12, 2021

Oct 18, 2021

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Part of collection

(ILLUMINA MENU) Protocols for SARS-CoV-2 Library Prep with ARTIC Primers, Bioinformatic Analysis, and Database Submission

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### Create Basespace Project

- 1 Login to the Illumina BaseSpace Platform and create a new BaseSpace project

Login to the BaseSpace Sequence Hub at:  
<https://basespace.illumina.com/>

Click on the 'Projects' tab on the top of the Sequence Hub homepage and create a new project directory by going to FILE -> NEW -> PROJECT:

## Projects

NAME	LAST UPDATED	OWNER
COVIDSeq_Test	2021-02-11	TOAST CDC
	2021-02-10	TOAST CDC
	2021-02-09	TOAST CDC

The Projects panel in the BaseSpace Sequence Hub

### Add Fastq files to BaseSpace Project

- 2 Add Fastq files to the new project directory. This can be done by either uploading fastq files from a local directory or by importing sequences using their Sequence Read Archive (SRA) accession numbers from NCBI using the 'SRA Import' app.

The step case format does not translate well in a pdf file. Be aware the pdf will present each step case sequentially which may be confusing when following the protocol.

Step 2 includes a Step case.

**Upload Fastqs**

**Import SRA**

### Run DRAGEN COVID Lineage App

step case

## Upload Fastqs

Go to FILE --> UPLOAD --> FILES and it will bring you to the 'Upload Files' page

## Upload Files

Projects: COVIDSeq\_Test2

Select the type of files you want to upload



The Basespace Projects 'Upload Files' page

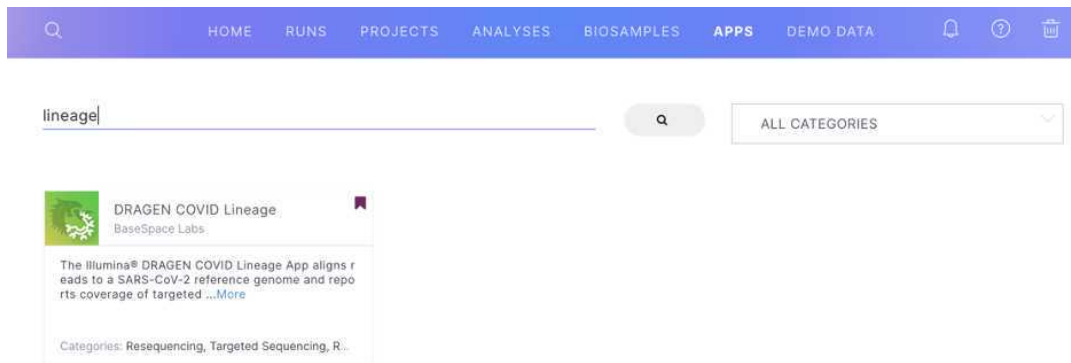
Click on the FASTQ tile and add the selected fastq files for upload. It should be noted that BaseSpace has specific upload requirements for fastq files including file naming schemes and header formats. They are described here:

[https://support.illumina.com/help/BaseSpace\\_Sequence\\_Hub/Source/Informatics/BS/UploadFastqReq\\_swBS.htm?Highlight=fastq](https://support.illumina.com/help/BaseSpace_Sequence_Hub/Source/Informatics/BS/UploadFastqReq_swBS.htm?Highlight=fastq)

### 3

Run the DRAGEN COVID Lineage App

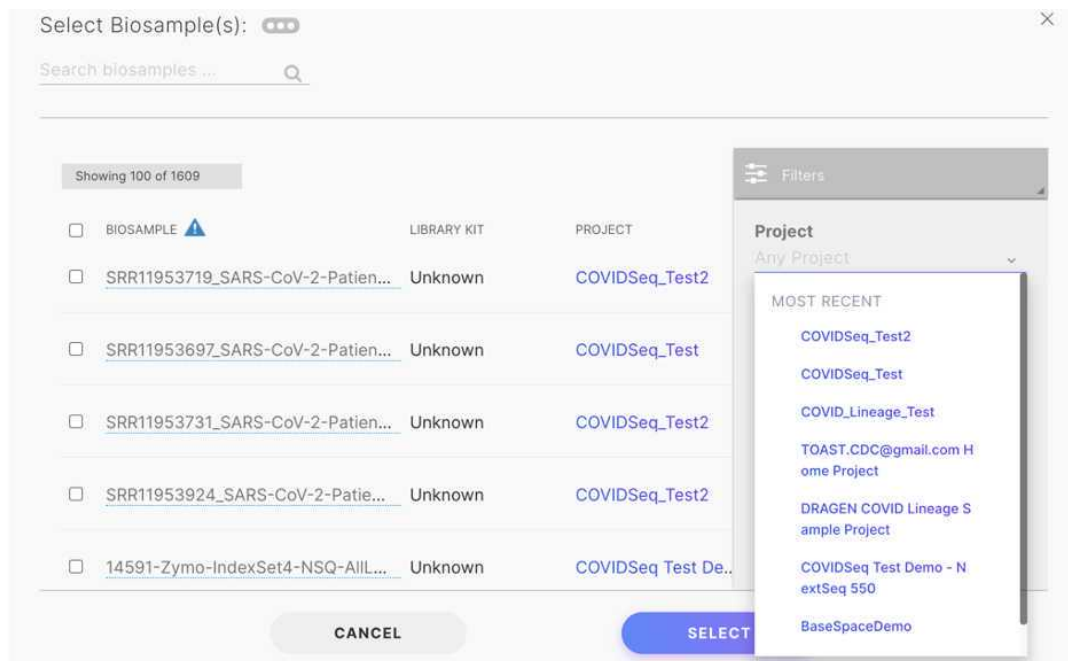
Go to the Apps panel and search 'lineage'. Click on the DRAGEN COVID Lineage app.



The 'DRAGEN COVID Lineage' BaseSpace app

Click 'LAUNCH APPLICATION' and then click 'SELECT PROJECT' and select the new project directory. Click 'SELECT BIOSAMPLE(S)'. In the 'SELECT BIOSAMPLE(S)' window click the

'Filters' button on the right. Choose the new project directory



The 'SELECT BIOSAMPLE(S)' window

Select the samples to be used in the analysis and click the 'SELECT' button. At the bottom of the application page click 'LAUNCH APPLICATION'. This will bring you to a new 'ANALYSIS' page similar to the following

## Analysis: DRAGEN COVID Lineage 02/11/2021 10:53:14

Project

SUMMARY REPORTS INPUTS FILES



### General Info

Name	DRAGEN COVID Lineage 02/11/2021 10:53:14
Application	DRAGEN COVID Lineage   Version: 3.5.0
Date Started	2021-02-11 23:01
Date Completed	N/A
Duration	N/A
Compute Charge	0.00 iCredits
Session Type	Multi-Node
Node Count	1 Node Queued for Analysis
Status	Queued for Analysis
Delivery	None

### Logs

Last che

The Analysis panel with a queued DRAGEN COVID Lineage run

## 4 Accessing the DRAGEN COVID Lineage app summary report

The DRAGEN COVID Lineage App is finished when the status line says 'Complete'. To view the 'Summary Report,' click on your analysis, navigate to the 'REPORTS' tab, and then click 'Summary' on the left side of the page.

Analysis: DRAGEN COVID Lineage 02/11/2021 6:02:03

Project COVIDSeq\_Test2

SUMMARY REPORTS INPUTS FILES

Summary

SRR11953697...  
Report  
Metrics

SRR11953731...  
Report

### Summary Report

DRAGEN Host Software Version 05.021.510.3.5.88-84-g76be5329 and Bio-IT Processor Version 0x04261818  
DRAGEN COVID Lineage 3.5.0

Kmer Config: kmer\_config.json

The Analysis panel showing the DRAGEN COVID Lineage app Summary Report

The 'Summary Report' compiles together three individual reports: 'Lineage Report', 'Clade Report' and 'Kmer Detection Report'. The 'Lineage Report', which provides Pangolin lineage assignments is shown below

## Summary Report







DRAGEN Host Software Version 05.021.510.3.5.88-84-g76be5329 and Bio-IT Processor Version 0x04261818  
DRAGEN COVID Lineage 3.5.0

Kmer Config: [kmer\\_config.json](#)

[Download Combined FASTA](#) <https://clades.nextstrain.org/> <https://pangolin.cog-uk.io/>

Note: Some tables in this report are very wide. Use mouse-wheel click or drag to scroll right to see more columns.

## Lineage Report

Copy	CSV	Excel	Search: <input type="text"/>			
Taxon 	Lineage 	Probability 	Version 	Status 	Note 	% of non-N bases 
SRR11953697_SARS-CoV-2-Patient-Sequencing-From-Partners-MGH	B.1	1	02/11/2021 00:00:00	passed_qc		77.66%
SRR11953731_SARS-CoV-2-Patient-Sequencing-From-Partners-MGH	B.1.268	1	02/11/2021 00:00:00	passed_qc		96.41%
SRR11953924_SARS-CoV-2-Patient-Sequencing-From-Partners-MGH	B.1	1	02/11/2021 00:00:00	passed_qc		99.87%

Showing 1 to 3 of 3 entries

The lineage section of the DRAGEN COVID Lineage Summary Report showing Pangolin lineage assignments

## Clade Report

Copy CSV Excel

Search:

seqName ↓↑	clade ↓↑	substitutions ↓↑	deletions ↓↑	insertions ↓↑	nonACGTNs ↓↑	pcrPri
SRR11953697_SARS-CoV-2-Patient-Sequencing-From-Partners-MGH	20C	C241T, C1059T, C14408T, A23403G, G25563T, C25710T, G28960T				China
SRR11953731_SARS-CoV-2-Patient-Sequencing-From-Partners-MGH	20C	C241T, C1059T, C3037T, G3892T, C14408T, A23403G, G25563T				
SRR11953924_SARS-CoV-2-Patient-Sequencing-From-Partners-MGH	20C	C84T, C241T, C1059T, C3037T, C6730T, C14177T, C14408T,				China Pasteur

The Clade section of the DRAGEN COVID Lineage Summary Report showing NextClade clade assignments

## 5 Downloading consensus sequences generated by the DRAGEN COVID Lineage app

To download the SARS-CoV-2 consensus sequences produced by the DRAGEN COVID Lineage program, click on the 'FILES' tab and then click on one of the samples. This will bring you to a new 'FILES' page similar to the following:



Analysis: DRAGEN COVID Lineage 02/11/2021 6:02:03

Project COVIDSeq\_Test2

SUMMARY REPORTS INPUTS FILES

NAME CREATED TYPE PATH SIZE

< Back

NAME	CREATED	TYPE	PATH	SIZE
.basespace		Folder	.basespace	--
consensus		Folder	consensus	--

The FILES tab of the Analysis panel for this DRAGEN COVID Lineage app run

Click on the 'consensus' directory and scroll down to the first file that has 'TYPE' 'fa'. Clicking on this sequence will pull up a new window the looks similar to the following:

SRR11953924\_SARS-CoV-2-Patient-Sequencing-From-Partners-MGH.consensus\_hard\_masked\_sequence.fa

>SRR11953924\_SARS-CoV-2-Patient-Sequencing-From-Partners-MGH|NC\_045512.2 Consensus sequence derived from variants in SRR11953924\_SARS-CoV-2-Patient-Sequencing-From-Partners-MGH applied to NC\_045512.2

```

NNNNNNGGTTTATACCTTCCAGGTAACAAACCAACCACTTTCGATCTCTGTAGATCTGTTCTCTAAA
CGAACTTTAAATTTGTGTGGCTGCTACTCGGCTGCATGCTTAGTGCACTCACGAGTATAATTAATAAC
TAATTACTGTGTTGACAGGACACAGTAAGTCTGCTATCTTCTGAGGCTGCTTACGGTTTCGTCGGTG
TTGCAGCCGATCATCAGCACATCTAGGTTTTGTCCGGGTGTGACCGAAAGGTAAGATGGAGAGCCTTGTC
CCTGGTTTCAACGAGAAAAACACAGTCCAACTCAGTTTGGCTGTTTTACAGGTTCCGACGTGCTGTAC
GTGGCTTTGGAGACTCCGTGGAGAGGCTTATCAGAGGCAGCTCAACATCTTAAGATGGCACTTGTGG
CTTAGTAGAAGTTGAAAAAGGCGTTTGCCTCAACTGAAACAGCCCTATGTTTCATCAACGTTCCGGAT
GCTCGAACTGCACCTCATGGTCATGTTATGGTTGAGCTGGTAGCAGAACTCGAAGGCATTTCAGTACGGTC
GTAGTGGTGAGACACTTGGTGTCTTGTCCCTCATGTGGGCGAAATACAGTGGCTTACCGCAAGGTTCT
TCTTCGTAAGAACGGTAATAAAGGAGCTGGTGGCCATAGTTACGGCGCCGATCTAAAGTCATTGACTTA
GGCGACGAGCTTGGCACTGATCCTTATGAAGATTTTCAAGAAAACTGGAACACTAAACATAGCAGTGGTG
TTACCCGTGAACCTCATGCGTGAGCTTAACGGAGGGGCATACACTCGCTATGTCGATAACAACTTCTGTGG
CCCTGATGGCTACCCTCTTGAAGTCAATTAAGACCTTCTAGCAGCTGCTGGTAAAGCTTTCATGCACCTTG
TCCGAACAACTGGACTTTATTGACACTAAGAGGGGTGTATCTGCTGCGTGAACATGAGCATGAAATTG
CTTGTACACGGAACGTTCTGAAAAGAGCTATGAATTGCAGACACCTTTTGAATTAATTTGGCAAGAA
ATTTGACATCTCAATGGGGAATGTCAAATTTGTATTTCCCTTAAATCCATAATCAAGACTATTCAA

```

CLOSE DOWNLOAD (29.82 KB)

SRR11953924\_SARS-CoV-2-Patient-Sequencing-From-Partners-MGH.consensus\_hard\_masked\_sequence.fa 32 B

Windows displaying the hard masked SARS-CoV-2 consensus sequence

This filename should end with 'consensus\_hard\_masked\_sequence.fa'. Click 'DOWNLOAD'

to save the consensus sequence to a local directory. Note there is an additional TYPE 'fa' file that ends with a 'consensus\_soft\_masked\_sequence.fa'. This consensus sequence should not be uploaded to a public database like Genbank or Gisaidd.

Additional documentation for the DRAGEN COVID Lineage app is available here:

[DRAGEN COVID Lineage \(illumina.com\)](https://illumina.com/DRAGEN-COVID-Lineage)

Before submitting the resulting SARS-CoV-2 consensus sequence assemblies to public repositories, such as NCBI GenBank or GISAID, refer to the following documentation describing submission criteria and minimum quality control thresholds:

GenBank Submission Criteria: [About GenBank Submission \(nih.gov\)](https://www.ncbi.nlm.nih.gov/genbank/submission/)

Gisaidd Submission Criteria: [Gisaidd inclusion criteria.pdf](#)

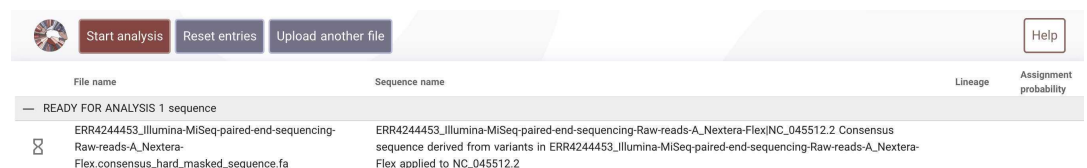
## Alternative Lineage Assignment

- The SARS-CoV-2 consensus sequence assembly generated by the CDRAGEN COVID Lineage app can also be uploaded to other lineage assignment software.

### 6.1 Upload the consensus sequence for each sample to the **Pangolin COVID-19 Lineage Assigner** at:

<https://pangolin.cog-uk.io/>

Click the 'Start analysis' button:



File name	Sequence name	Lineage	Assignment probability
READY FOR ANALYSIS 1 sequence			
ERR4244453_Illumina-MiSeq-paired-end-sequencing-Raw-reads-A_Nextera-Flex-consensus_hard_masked_sequence.fa	ERR4244453_Illumina-MiSeq-paired-end-sequencing-Raw-reads-A_Nextera-Flex(NC_045512.2 Consensus sequence derived from variants in ERR4244453_Illumina-MiSeq-paired-end-sequencing-Raw-reads-A_Nextera-Flex applied to NC_045512.2		

Pangolin COVID-19 Lineage Assigner example

The Pangolin COVID-19 Lineage Assigner returns the lineage classification and assignment probability:

Reset entries          Upload another file		Help	
File name	Sequence name	Lineage	Assignment probability
ANALYSED (Click tick icon for more info) 1 sequence			
✓ ERR4244453_Illumina-MiSeq-paired-end-sequencing-Raw-reads-A_Nextera-Flex.consensus_hard_masked_sequence.fa	ERR4244453_Illumina-MiSeq-paired-end-sequencing-Raw-reads-A_Nextera-Flex(NC_045512.2 Consensus sequence derived from variants in ERR4244453_Illumina-MiSeq-paired-end-sequencing-Raw-reads-A_Nextera-Flex applied to NC_045512.2	B.1.1	1.0

Pangolin COVID-19 Lineage Assigner output

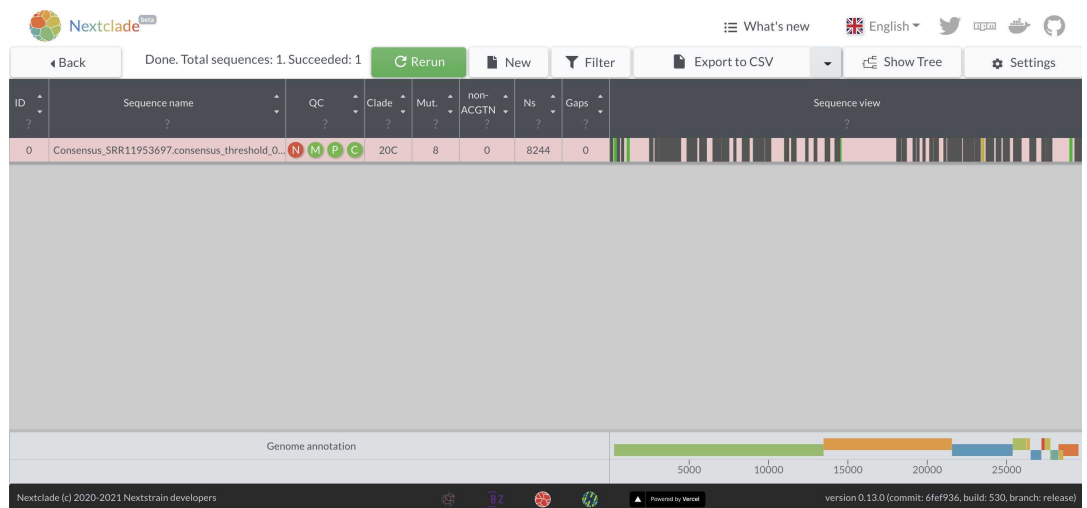
6.2 Or upload the consensus sequence for each sample to the **Nextclade** clade assignment web portal at:

<https://clades.nextstrain.org/>

The image shows the Nextclade web portal interface. At the top, there's a navigation bar with the Nextclade logo, a 'To Results' button, and links for 'What's new', 'English', and social media. The main heading is 'Nextclade beta v0.13.0' with the subtitle 'Clade assignment, mutation calling, and sequence quality checks'. Below this, there are four main sections: 'Simple' (No installation or setup - drop a file and see the results), 'Private' (No remote processing - sequence data never leaves your computer), 'Mutation Calling' (Find differences of your sequences relative to the reference in standard numbering), 'Clade Assignment' (Find out which Nextstrain clades your samples are from), 'Phylogenetic Placement' (See where on the SARS-CoV-2 tree your sequences fall), and 'Quality Control' (Check your data against multiple QC metrics). On the right, there's a 'Sequences' section with a dropdown for 'SARS-CoV-2', a toggle for 'Simple mode' and 'Advanced mode', and buttons for 'From file', 'From URL', and 'Paste'. A large area for file upload is shown with the text 'Drag & Drop a file here' and a 'Select a file' button. A 'FASTA' icon is also present.

NextClade assignment web portal

The Nextclade server provides clade classification as well as QC metrics and a list of amino acid substitutions. A summary output file can be downloaded with the 'Export to CSV' button.



Nextclade clade assignment output