Cumulus Documentation

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Dec 04, 2019

Contents

All of our docker images are publicly available on Docker Hub and Quay. Our workflows use Docker Hub as the default Docker registry. Users can use Quay as the Docker registry by entering `quay.io/cumulus/` for the workflow input `docker_registry`, or can enter a custom registry URL of their choice.

CHAPTER 1

Version 0.11.0 *December 4, 2019*

Reorganized Cumulus documentation.

CHAPTER 2

Version 0.10.0 *October 2, 2019*

scCloud renamed to cumulus.

cumulus can accept either a sample sheet or a single file.

CHAPTER 3

Version 0.7.0 *Feburary 14, 2019*

Added support for 10x genomics scATAC assays.

scCloud runs FIt-SNE as default.

CHAPTER 4

Version 0.6.0 *January 31, 2019*

Added support for 10x genomics V3 chemistry.

Added support for extracting feature matrix for Perturb-Seq data.

Added R script to convert output_name.seurat.h5ad to Seurat object. Now the raw.data slot stores filtered raw counts.

Added min_umis and max_umis to filter cells based on UMI counts.

Added QC plots and improved filtration spreadsheet.

Added support for plotting UMAP and FLE.

Now users can upload their JSON file to annotate cell types.

Improved documentation.

Added lightGBM based marker detection.

CHAPTER 5

Version 0.5.0 *November 18, 2018*

Added support for plated-based SMART-Seq2 scRNA-Seq data.

CHAPTER 6

Version 0.4.0 *October 26, 2018*

Added CITE-Seq module for analyzing CITE-Seq data.

CHAPTER 7

Version 0.3.0 *October 24, 2018*

Added the demuxEM module for demultiplexing cell-hashing/nuclei-hashing data.

CHAPTER 8

Version 0.2.0 *October 19, 2018*

Added support for V(D)J and CITE-Seq/cell-hashing/nuclei-hashing.

CHAPTER 9

Version 0.1.0 *July 27, 2018*

KCO tools released!

9.1 First Time Running

9.1.1 Authenticate with Google

If you've done this before you can skip this step - you only need to do this once.

1. Ensure the Google Cloud SDK is installed on your computer.

   Note: Broad users do not have to install this-they can type:

   ```
   reuse Google-Cloud-SDK
   ```

   to make the Google Cloud tools available.

2. Execute the following command to login to Google Cloud.:

   ```
   gcloud auth login
   ```

3. Copy and paste the link in your unix terminal into your web browser.
4. Enter authorization code in unix terminal.

9.1.2 Create a Terra workspace

1. Create a new Terra workspace by clicking Create New Workspace in Terra

   For more detailed instructions please see this document.

9.2 Latest and stable versions on Terra

Cumulus is a fast growing project. As a result, we frequently update WDL snapshot versions on Terra. See below for latest and stable WDL versions you can use.

9.2.1 Latest version

WDL	Snapshot	Function
cumulus/cellranger_workflow	4	Run Cell Ranger tools, which include extracting sequence reads using cellranger mkfastq or cellranger-atac mkfastq, generate count matrix using cellranger count or cellranger-atac count, run cellranger vdj or feature-barcode extraction
cumulus/smartseq2	3	Run Bowtie2 and RSEM to generate gene-count matrices for SMART-Seq2 data from FASTQ files
cumulus/cumulus	8	Run cumulus analysis module for variable gene selection, batch correction, PCA, diffusion map, clustering, visualization, differential expression analysis, cell type annotation, etc.
cumulus/cumulus_subcluster	5	Run subcluster analysis using cumulus
cumulus/cumulus_hashing_cite_seq	5	Run cumulus for cell-hashing/nucleus-hashing/CITE-Seq analysis

9.2.2 Stable version - v0.11.0

WDL	Snapshot	Function
cumulus/cellranger_workflow	4	Run Cell Ranger tools, which include extracting sequence reads using cellranger mkfastq or cellranger-atac mkfastq, generate count matrix using cellranger count or cellranger-atac count, run cellranger vdj or feature-barcode extraction
cumulus/smartseq2	3	Run Bowtie2 and RSEM to generate gene-count matrices for SMART-Seq2 data from FASTQ files
cumulus/cumulus	8	Run cumulus analysis module for variable gene selection, batch correction, PCA, diffusion map, clustering, visualization, differential expression analysis, cell type annotation, etc.
cumulus/cumulus_subcluster	5	Run subcluster analysis using cumulus
cumulus/cumulus_hashing_cite_seq	5	Run cumulus for cell-hashing/nucleus-hashing/CITE-Seq analysis

9.2.3 Stable version - v0.10.0

WDL	Snapshot	Function
cumulus/cellranger_workflow	3	Run Cell Ranger tools, which include extracting sequence reads using cellranger mkfastq or cellranger-atac mkfastq, generate count matrix using cellranger count or cellranger-atac count, run cellranger vdj or feature-barcode extraction
cumulus/smartseq2	3	Run Bowtie2 and RSEM to generate gene-count matrices for SMART-Seq2 data from FASTQ files
cumulus/cumulus	7	Run cumulus analysis module for variable gene selection, batch correction, PCA, diffusion map, clustering, visualization, differential expression analysis, cell type annotation, etc.
cumulus/cumulus_subcluster	4	Run subcluster analysis using cumulus
cumulus/cumulus_hashing_cite_seq	4	Run cumulus for cell-hashing/nucleus-hashing/CITE-Seq analysis

9.2.4 Stable version - HTAPP v2

WDL	Snapshot	Function
regev/cellranger_mkfastq_count	45	Run Cell Ranger to extract FASTQ files and generate gene-count matrices for 10x genomics data
scCloud/smartseq2	5	Run Bowtie2 and RSEM to generate gene-count matrices for SMART-Seq2 data from FASTQ files
scCloud/scCloud	14	Run scCloud analysis module for variable gene selection, batch correction, PCA, diffusion map, clustering and more
scCloud/scCloud_subcluster	9	Run subcluster analysis using scCloud
scCloud/scCloud_hashing_cite_seq	9	Run scCloud for cell-hashing/nucleus-hashing/CITE-Seq analysis

9.2.5 Stable version - HTAPP v1

WDL	Snapshot	Function
regev/cellranger_mkfastq_count	39	Run Cell Ranger to extract FASTQ files and generate gene-count matrices for 10x genomics data
scCloud/scCloud	3	Run scCloud analysis module for variable gene selection, batch correction, PCA, diffusion map, clustering and more

9.3 Run Cell Ranger tools using cellranger_workflow

`cellranger_workflow` wraps Cell Ranger to process single-cell/nucleus RNA-seq, feature barcoding (cell/nucleus hashing, CITE-seq, Perturb-seq), single-cell ATAC-seq and single-cell immune profiling data.

9.3.1 A general step-by-step instruction

1. Import `cellranger_workflow`

Import *cellranger_workflow* workflow to your workspace.

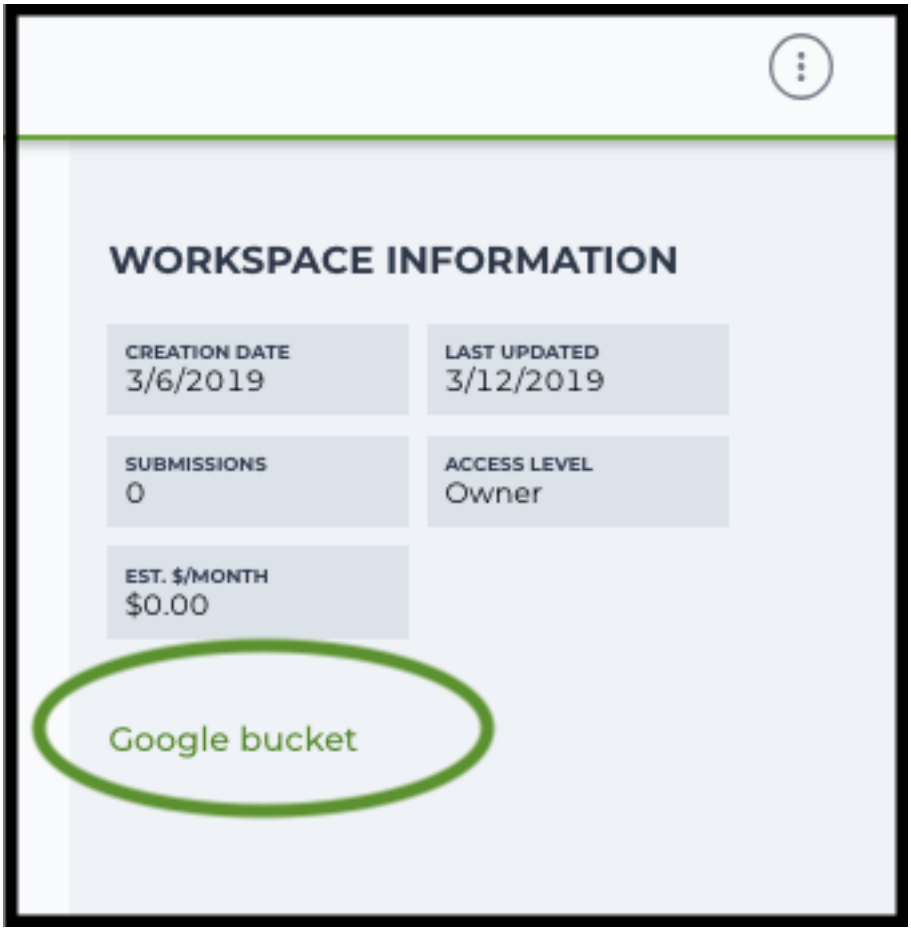
See the Terra documentation for adding a workflow. The *cellranger_workflow* workflow is under `Broad Methods Repository` with name "**cumulus/cellranger_workflow**".

Moreover, in the workflow page, click the `Export to Workspace...` button, and select the workspace to which you want to export *cellranger_workflow* workflow in the drop-down menu.

2. Upload sequencing data to Google bucket

Copy your sequencing output to your workspace bucket using gsutil (you already have it if you've installed Google cloud SDK) in your unix terminal.

You can obtain your bucket URL in the dashboard tab of your Terra workspace under the information panel.



Use `gsutil cp [OPTION]... src_url dst_url` to copy data to your workspace bucket. For example, the following command copies the directory at /foo/bar/nextseq/Data/VK18WBC6Z4 to a Google bucket:

```
gsutil -m cp -r /foo/bar/nextseq/Data/VK18WBC6Z4 gs://fc-e0000000-0000-0000-
↪0000-000000000000/VK18WBC6Z4
```

`-m` means copy in parallel, `-r` means copy the directory recursively, and `gs://fc-e0000000-0000-0000-0000-000000000000` should be replaced by your own workspace Google bucket URL.

Note: Broad users need to be on an UGER node (not a login node) in order to use the `-m` flag

Request an UGER node:

```
reuse UGER
qrsh -q interactive -l h_vmem=4g -pe smp 8 -binding linear:8 -P regevlab
```

The above command requests an interactive node with 4G memory per thread and 8 threads. Feel free to change the memory, thread, and project parameters.

Once you're connected to an UGER node, you can make gsutil available by running:

```
reuse Google-Cloud-SDK
```

3. Prepare a sample sheet

3.1 Sample sheet format:

Please note that the columns in the CSV can be in any order, but that the column names must match the recognized headings.

The sample sheet describes how to demultiplex flowcells and generate channel-specific count matrices. Note that *Sample*, *Lane*, and *Index* columns are defined exactly the same as in 10x's simple CSV layout file.

A brief description of the sample sheet format is listed below (**required column headers are shown in bold**).

Column	Description
Sample	Contains sample names. Each 10x channel should have a unique sample name.
Reference	Provides the reference genome used by Cell Ranger for each 10x channel. The elements in the *reference* column can be either Google bucket URLs to reference tarballs or keywords such as *GRCh38_v3.0.0*. A full list of available keywords is included in each of the following data type sections (e.g. sc/snRNA-seq) below.
Flowcell	Indicates the Google bucket URLs of uploaded BCL folders. If starts with FASTQ files, this should be Google bucekt URLs of uploaded FASTQ folders. The FASTQ folders should contain one subfolder for each sample in the flowcell with the sample name as the subfolder name. Each subfolder contains FASTQ files for that sample.
Lane	Tells which lanes the sample was pooled into. Can be either single lane (e.g. 8) or a range (e.g. 7-8) or all (e.g. *).
Index	Sample index (e.g. SI-GA-A12).
Chemistry	Describes the 10x chemistry used for the sample. This column is optional.
DataType	Describes the data type of the sample — *rna*, *vdj*, *adt*, or *crispr*. **rna** refers to gene expression data (*cellranger count*), **vdj** refers to V(D)J data (*cellranger vdj*), **adt** refers to antibody tag data, which can be either CITE-Seq, cell-hashing, or nucleus-hashing, **crispr** refers to Perturb-seq guide tag data, **atac** refers to scATAC-Seq data (*cellranger-atac count*). This column is optional and the default data type is *rna*.
FeatureBarcodeFile	Google bucket urls pointing to feature barcode files for *adt* and *crispr* data. Features can be either antibody for CITE-Seq, cell-hashing, nucleus-hashing or gRNA for Perburb-seq. This column is optional provided no *adt* or *crispr* data are in the sample sheet.

The sample sheet supports sequencing the same 10x channels across multiple flowcells. If a sample is sequenced across multiple flowcells, simply list it in multiple rows, with one flowcell per row. In the following example, we have 4 samples sequenced in two flowcells.

Example:

```
Sample,Reference,Flowcell,Lane,Index,Chemistry,DataType,FeatureBarcodeFile
sample_1,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/
↪VK18WBC6Z4,1-2,SI-GA-A8,threeprime,rna
```

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```
sample_2,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/
↪VK18WBC6Z4,3-4,SI-GA-B8,SC3Pv3,rna
sample_3,mm10_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK18WBC6Z4,
↪5-6,SI-GA-C8,fiveprime,rna
sample_4,mm10_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK18WBC6Z4,
↪7-8,SI-GA-D8,fiveprime,rna
sample_1,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/
↪VK10WBC9Z2,1-2,SI-GA-A8,threeprime,rna
sample_2,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/
↪VK10WBC9Z2,3-4,SI-GA-B8,SC3Pv3,rna
sample_3,mm10_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK10WBC9Z2,
↪5-6,SI-GA-C8,fiveprime,rna
sample_4,mm10_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK10WBC9Z2,
↪7-8,SI-GA-D8,fiveprime,rna
```

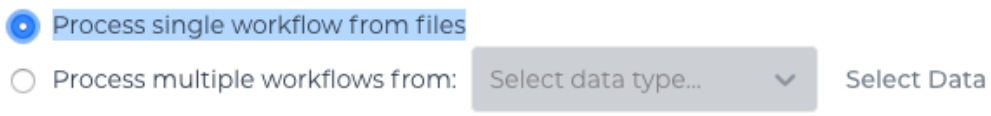
3.2 Upload your sample sheet to the workspace bucket:

Example:

```
gsutil cp /foo/bar/projects/sample_sheet.csv gs://fc-e0000000-0000-
↪0000-0000-000000000000/
```

4. Launch analysis

In your workspace, open `cellranger_workflow` in `WORKFLOWS` tab. Select the desired snapshot version (e.g. latest). Select `Process single workflow from files` as below



and click `SAVE` button. Select `Use call caching` and click `INPUTS`. Then fill in appropriate values in the `Attribute` column. Alternative, you can upload a JSON file to configure input by clicking `Drag or click to upload json`.

Once INPUTS are appropriated filled, click `RUN ANALYSIS` and then click `LAUNCH`.

5. Notice: run `cellranger mkfastq` if you are non Broad Institute users

Non Broad Institute users that wish to run `cellranger mkfastq` must create a custom docker image that contains `bcl2fastq`.

See *bcl2fastq* instructions.

6. Do not run `cellranger mkfastq`

Sometimes, users might want to perform demultiplexing locally and only run the count part on the cloud. This section describes how to only run the count part via `cellranger_workflow`.

1. Copy your FASTQ files to the workspace using gsutil in your unix terminal.

   You should upload folders of FASTQ files. The uploaded folder (for one flowcell) should contain one subfolder for each sample belong to the this flowcell. In addition, the subfolder name should be the sample name. Each subfolder contains FASTQ files for that sample.

Example:

```
gsutil -m cp -r /foo/bar/fastq_path/K18WBC6Z4 gs://fc-e0000000-0000-0000-
↪0000-000000000000/K18WBC6Z4_fastq
```

2. Create a sample sheet.

   Flowcell column should list Google bucket URLs of the FASTQ folders for flowcells.

   Example:

```
Sample,Reference,Flowcell
sample_1,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/
↪K18WBC6Z4_fastq
```

3. Set optional input `run_mkfastq` to `false`.

9.3.2 Single-cell and single-nucleus RNA-seq

To process sc/snRNA-seq data, follow the specific instructions below.

Sample sheet

1. **Reference** column.

   Pre-built scRNA-seq references are summarized below.

Keyword	Description
GRCh38_v3.0.0	Human GRCh38, cellranger reference 3.0.0, Ensembl v93 gene annotation
hg19_v3.0.0	Human hg19, cellranger reference 3.0.0, Ensembl v87 gene annotation
mm10_v3.0.0	Mouse mm10, cellranger reference 3.0.0, Ensembl v93 gene annotation
GRCh38_and_mm10_v3.1.0	Human (GRCh38) and mouse (mm10), cellranger references 3.1.0, Ensembl v93 gene annotations for both human and mouse
GRCh38_v1.2.0 or **GRCh38**	Human GRCh38, cellranger reference 1.2.0, Ensembl v84 gene annotation
hg19_v1.2.0 or **hg19**	Human hg19, cellranger reference 1.2.0, Ensembl v82 gene annotation
mm10_v1.2.0 or **mm10**	Mouse mm10, cellranger reference 1.2.0, Ensembl v84 gene annotation
GRCh38_and_mm10_v1.2.0 or **GRCh38_and_mm10**	Human and mouse, built from GRCh38 and mm10 cellranger references, Ensembl v84 gene annotations are used

   Pre-built snRNA-seq references are summarized below.

Keyword	Description
GRCh38_premrna_v1.2.0 or **GRCh38_premrna**	Human, introns included, built from GRCh38 cellranger reference 1.2.0, Ensembl v84 gene annotation, treating annotated transcripts as exons
mm10_premrna_v1.2.0 or **mm10_premrna**	Mouse, introns included, built from mm10 cellranger reference 1.2.0, Ensembl v84 gene annotation, treating annotated transcripts as exons
GRCh38_premrna_and_mm10_premrna_v1.2.0 or **GRCh38_premrna_and_mm10_premrna**	Human and mouse, introns included, built from GRCh38_premrna_v1.2.0 and mm10_premrna_v1.2.0

2. **Index** column.

   Put 10x single cell 3' sample index set names (e.g. SI-GA-A12) here.

3. *Chemistry* column.

   According to *cellranger count*'s documentation, chemistry can be

Chemistry	Explanation
auto	autodetection (default). If the index read has extra bases besides cell barcode and UMI, autodetection might fail. In this case, please specify the chemistry
threeprime	Single Cell 3
fiveprime	Single Cell 5
SC3Pv1	Single Cell 3 v1
SC3Pv2	Single Cell 3 v2
SC3Pv3	Single Cell 3 v3. You should set cellranger version input parameter to >= 3.0.2
SC5P-PE	Single Cell 5 paired-end (both R1 and R2 are used for alignment)
SC5P-R2	Single Cell 5 R2-only (where only R2 is used for alignment)

4. *DataType* column.

   This column is optional with a default **rna**. If you want to put a value, put **rna** here.

5. *FetureBarcodeFile* column.

   Leave it blank for scRNA-seq and snRNA-seq.

6. Example:

```
Sample,Reference,Flowcell,Lane,Index,Chemistry,DataType,FeatureBarcodeFile
sample_1,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK18WBC6Z4,1-
↪2,SI-GA-A8,threeprime,rna
sample_2,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK18WBC6Z4,3-
↪4,SI-GA-B8,SC3Pv3,rna
sample_3,mm10_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK18WBC6Z4,5-6,
↪SI-GA-C8,fiveprime,rna
sample_4,mm10_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK18WBC6Z4,7-8,
↪SI-GA-D8,fiveprime,rna
sample_1,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK10WBC9Z2,1-
↪2,SI-GA-A8,threeprime,rna
sample_2,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK10WBC9Z2,3-
↪4,SI-GA-B8,SC3Pv3,rna
sample_3,mm10_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK10WBC9Z2,5-6,
↪SI-GA-C8,fiveprime,rna
```

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```
sample_4,mm10_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/VK10WBC9Z2,7-8,
↪SI-GA-D8,fiveprime,rna
```

Workflow input

For sc/snRNA-seq data, `cellranger_workflow` takes Illumina outputs as input and runs `cellranger mkfastq` and `cellranger count`. Revalant workflow inputs are described below, with required inputs highlighted in bold.

Name	Description	Example	Default
input_samplesheet	Sample Sheet (contains Sample, Reference, Flowcell, Lane, Index as required and Chemistry, DataType, FeatureBarcodeFile as optional)	"gs://fc-e0000000-0000-0000-0000-000000000000/sample_sheet.csv"	
output_directory	Output directory	"gs://fc-e0000000-0000-0000-0000-000000000000/cellranger_output"	
run_mkfastq	If you want to run `cellranger mkfastq`	true	true
run_count	If you want to run `cellranger count`	true	true
delete_input_bcl_dir	If delete input BCL directories after demux. If false, you should delete this folder yourself so as to not incur storage charges	false	false
force_cells	Force pipeline to use this number of cells, bypassing the cell detection algorithm, mutually exclusive with expect_cells	6000	
expect_cells	Expected number of recovered cells. Mutually exclusive with force_cells	3000	
secondary	Perform Cell Ranger secondary analysis (dimensionality reduction, clustering, etc.)	false	false
cellranger_version	cellranger version, could be 3.1.0, 3.0.2, or 2.2.0	"3.1.0"	"3.1.0"
docker_registry	Docker registry to use for cellranger_workflow. Options: • "cumulusprod/" for Docker Hub images; • "quay.io/cumulus/" for backup images on Red Hat registry.	"cumulusprod/"	"cumulusprod/"
cellranger_mkfastq_docker_registry	Docker registry for `cellranger mkfastq`. Default is the registry to which only Broad users have access. See *bcl2fastq* for making your own registry.	"gcr.io/broad-cumulus"	"gcr.io/broad-cumulus"
zones	Google cloud zones	"us-central1-a us-west1-a"	"us-central1-a us-central1-b us-central1-c us-central1-f us-east1-b us-east1-c us-east1-d us-west1-a us-west1-b us-west1-c"
num_cpu	Number of cpus to request for one node for cellranger mkfastq and cellranger count	32	32
memory	Memory size string for cellranger mkfastq and cellranger count	"120G"	"120G"
mkfastq_disk_space	Disk space in GB for mkfastq	1500	1500
count_disk_space	Disk space in GB needed for cellranger count	500	500

Workflow output

See the table below for important sc/snRNA-seq outputs.

Name	Type	Description
output_fastqs_directory	Array[String]	A list of google bucket urls containing FASTQ files, one url per flowcell.
output_count_directory	Array[String]	A list of google bucket urls containing count matrices, one url per sample.
metrics_summaries	File	A excel spreadsheet containing QCs for each sample.
output_web_summary	Array[File]	A list of htmls visualizing QCs for each sample (cellranger count output).
count_matrix	String	gs url for a template count_matrix.csv to run Cumulus.

9.3.3 Feature barcoding assays (cell & nucleus hashing, CITE-seq and Perturb-seq)

`cellranger_workflow` can extract feature-barcode count matrices in CSV format for feature barcoding assays such as *cell and nucleus hashing*, *CITE-seq*, and *Perturb-seq*. For cell and nucleus hashing as well as CITE-seq, the feature refers to antibody. For Perturb-seq, the feature refers to guide RNA. Please follow the instructions below to configure `cellranger_workflow`.

Prepare feature barcode files

Prepare a CSV file with the following format: feature_barcode,feature_name. See below for an example:

```
TTCCTGCCATTACTA,sample_1
CCGTACCTCATTGTT,sample_2
GGTAGATGTCCTCAG,sample_3
TGGTGTCATTCTTGA,sample_4
```

The above file describes a cell hashing application with 4 samples.

Then upload it to your google bucket:

```
gsutil antibody_index.csv gs://fc-e0000000-0000-0000-0000-000000000000/
↪antibody_index.csv
```

Sample sheet

1. **Reference** column.

   This column is not used for extracting feature-barcode count matrix. To be consistent, please put the reference for the associated scRNA-seq assay here.

2. **Index** column.

   The index can be either Illumina index primer sequence (e.g. `ATTACTCG`, also known as `D701`), or 10x single cell 3' sample index set names (e.g. SI-GA-A12).

   Note 1: All index sequences (including 10x's) should have the same length (8 bases). If one index sequence is shorter (e.g. ATCACG), pad it with P7 sequence (e.g. ATCACGAT).

Note 2: It is users' responsibility to avoid index collision between 10x genomics' RNA indexes (e.g. SI-GA-A8) and Illumina index sequences for used here (e.g. `ATTACTCG`).

3. *Chemistry* column.

   The following keywords are accepted for *Chemistry* column:

Chemistry	Explanation
SC3Pv3	Single Cell 3 v3 (default).
SC3Pv2	Single Cell 3 v2
fiveprime	Single Cell 5
SC5P-PE	Single Cell 5 paired-end (both R1 and R2 are used for alignment)
SC5P-R2	Single Cell 5 R2-only (where only R2 is used for alignment)

4. *DataType* column.

   Put **adt** here if the assay is CITE-seq, cell or nucleus hashing. Put **crispr** here if Perturb-seq.

5. *FetureBarcodeFile* column.

   Put Google Bucket URL of the feature barcode file here.

6. Example:

```
Sample,Reference,Flowcell,Lane,Index,Chemistry,DataType,FeatureBarcodeFile
sample_1_rna,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/
↪VK18WBC6Z4,1-2,SI-GA-A8,threeprime,rna
sample_1_adt,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/
↪VK18WBC6Z4,1-2,ATTACTCG,threeprime,adt,gs://fc-e0000000-0000-0000-0000-
↪000000000000/antibody_index.csv
sample_2_adt,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/
↪VK18WBC6Z4,3-4,TCCGGAGA,SC3Pv3,adt,gs://fc-e0000000-0000-0000-0000-000000000000/
↪antibody_index.csv
sample_3_crispr,GRCh38_v3.0.0,gs://fc-e0000000-0000-0000-0000-000000000000/
↪VK18WBC6Z4,5-6,CGCTCATT,SC3Pv3,crispr,gs://fc-e0000000-0000-0000-0000-
↪000000000000/crispr_index.csv
```

In the sample sheet above, despite the header row,

- First row describes the normal 3' RNA assay;
- Second row describes its associated antibody tag data, which can from either a CITE-seq, cell hashing, or nucleus hashing experiment.
- Third row describes another tag data, which is in 10x genomics' V3 chemistry. For tag and crispr data, it is important to explicitly state the chemistry (e.g. `SC3Pv3`).
- Last row describes one gRNA guide data for Perturb-seq (see `crispr` in *DataType* field).

Workflow input

For feature barcoding data, `cellranger_workflow` takes Illumina outputs as input and runs `cellranger mkfastq` and `cumulus adt`. Revalant workflow inputs are described below, with required inputs highlighted in bold.

Name	Description	Example	Default
input_csv_file	Sample Sheet (contains Sample, Reference, Flowcell, Lane, Index as required and Chemistry, DataType, FeatureBarcodeFile as optional)	"gs://fc-e0000000-0000-0000-0000-000000000000/sample_sheet.csv"	
output_directory	Output directory	"gs://fc-e0000000-0000-0000-0000-000000000000/cellranger_output"	
run_mkfastq	If you want to run `cellranger mkfastq`	true	true
delete_input_bcl_dir	If delete BCL directories after demux. If false, you should delete this folder yourself so as to not incur storage charges	false	false
scaffold_sequence	Scaffold sequence in sgRNA for Purturb-seq, only used for crispr data type	"GTTTAAGAGCTAAGCTGGAA"	
max_mismatch	Maximum hamming distance in feature barcodes for the adt task	3	3
min_read_ratio	Minimum read count ratio (non-inclusive) to justify a feature given a cell barcode and feature combination, only used for the adt task and crispr data type	0.1	0.1
cellranger_version	cellranger version, could be 3.1.0, 3.0.2, 2.2.0	"3.1.0"	"3.1.0"
cumulus_version	Cumulus version for extracting feature barcode matrix, currently only 0.11.0	"0.11.0"	"0.11.0"
docker_registry	Docker registry to use for cellranger_workflow. Options: • "cumulusprod/" for Docker Hub images; • "quay.io/cumulus/" for backup images on Red Hat registry.	"cumulusprod/"	"cumulusprod/"
cellranger_mkfastq_docker_registry	Docker registry for `cellranger mkfastq`. Default is the registry to which only Broad users have access. See *bcl2fastq* for making your own registry.	"gcr.io/broad-cumulus"	"gcr.io/broad-cumulus"
zones	Google cloud zones	"us-central1-a us-west1-a"	"us-central1-a us-central1-b us-central1-c us-central1-f us-east1-b us-east1-c us-east1-d us-west1-a us-west1-b us-west1-c"
num_cpu	Number of cpus to request for one node for cellranger mkfastq	32	32
memory	Memory size string for cellranger mkfastq	"120G"	"120G"
feature_ref_mem	Optional memory string for extracting feature count matrix	"32G"	32G
mkfastq_disk_space	Optional disk space in GB for mkfastq	1500	1500

Parameters used for feature count matrix extraction

If the chemistry is V2, 10x genomics v2 cell barcode white list will be used, a hamming distance of 1 is allowed for matching cell barcodes, and the UMI length is 10. If the chemistry is V3, 10x genomics v3 cell barcode white list will be used, a hamming distance of 0 is allowed for matching cell barcodes, and the UMI length is 12.

For Perturb-seq data, a small number of sgRNA protospace sequences will be sequenced ultra-deeply and we may have PCR chimeric reads. Therefore, we generate filtered feature count matrices as well in a data driven manner:

1. First, plot the histogram of UMIs with certain number of read counts. The number of UMIs with `x` supporting reads decreases when `x` increases. We start from `x = 1`, and a valley between two peaks is detected if we find `count[x] < count[x + 1] < count[x + 2]`. We filter out all UMIs with `< x` supporting reads since they are likely formed due to chimeric reads.
2. In addition, we also filter out barcode-feature-UMI combinations that have their read count ratio, which is defined as total reads supporting barcode-feature-UMI over total reads supporting barcode-UMI, no larger than `min_read_ratio` parameter set above.

Workflow outputs

See the table below for important outputs.

Name	Type	Description
output_fastqs_directory	Array[String]	A list of google bucket urls containing FASTQ files, one url per flowcell.
output_count_directory	Array[String]	A list of google bucket urls containing feature-barcode count matrices, one url per sample.
count_matrix	String	gs url for a template count_matrix.csv to run cumulus.

In addition, For each antibody tag or crispr tag sample, a folder with the sample ID is generated under `output_directory`. In the folder, two files — `sample_id.csv` and `sample_id.stat.csv.gz` — are generated.

`sample_id.csv` is the feature count matrix. It has the following format. The first line describes the column names: `Antibody/CRISPR,cell_barcode_1,cell_barcode_2,...,cell_barcode_n`. The following lines describe UMI counts for each feature barcode, with the following format: `feature_name,umi_count_1,umi_count_2,...,umi_count_n`.

`sample_id.stat.csv.gz` stores the gzipped sufficient statistics. It has the following format. The first line describes the column names: `Barcode,UMI,Feature,Count`. The following lines describe the read counts for every barcode-umi-feature combination.

If data type is `crispr`, three additional files, `sample_id.umi_count.pdf`, `sample_id.filt.csv` and `sample_id.filt.stat.csv.gz`, are generated.

`sample_id.umi_count.pdf` plots number of UMIs against UMI with certain number of reads and colors UMIs with high likelihood of being chimeric in blue and other UMIs in red. This plot is generated purely based on number of reads each UMI has.

`sample_id.filt.csv` is the filtered feature count matrix. It has the same format as `sample_id.csv`.

`sample_id.filt.stat.csv.gz` is the filtered sufficient statistics. It has the same format as `sample_id.stat.csv.gz`.

9.3.4 Single-cell ATAC-seq

To process scATAC-seq data, follow the specific instructions below.

Sample sheet

1. **Reference** column.

   Pre-built scATAC-seq references are summarized below.

Keyword	Description
GRCh38_atac_v1.1.0	Human GRCh38, cellranger-atac reference 1.1.0
mm10_atac_v1.1.0	Mouse mm10, cellranger-atac reference 1.1.0
hg19_atac_v1.1.0	Human hg19, cellranger-atac reference 1.1.0
b37_atac_v1.1.0	Human b37 build, cellranger-atac reference 1.1.0
GRCh38_and_mm10_atac_v1.1.0	Human GRCh38 and mouse mm10, cellranger-atac reference 1.1.0
hg19_and_mm10_atac_v1.1.0	Human hg19 and mouse mm10, cellranger-atac reference 1.1.0

2. **Index** column.

   Put 10x single cell ATAC sample index set names (e.g. SI-NA-B1) here.

3. *Chemistry* column.

   This column is not used for scATAC-seq data. Put **auto** here as a placeholder if you decide to include the Chemistry column.

4. *DataType* column.

   Set it to **atac**.

5. *FetureBarcodeFile* column.

   Leave it blank for scATAC-seq.

6. Example:

```
Sample,Reference,Flowcell,Lane,Index,Chemistry,DataType
sample_atac,GRCh38_atac_v1.1.0,gs://fc-e0000000-0000-0000-0000-000000000000/
↪VK10WBC9YB,*,SI-NA-A1,auto,atac
```

Workflow input

`cellranger_workflow` takes Illumina outputs as input and runs `cellranger-atac mkfastq` and `cellranger-atac count`. Please see the description of inputs below. Note that required inputs are shown in bold.

Name	Description	Example	Default
input_csv_file	Sample Sheet (contains Sample, Reference, Flowcell, Lane, Index as required and Chemistry, DataType, FeatureBarcodeFile as optional)	"gs://fc-e0000000-0000-0000-0000-000000000000/sample_sheet.csv"	
output_directory	Output directory	"gs://fc-e0000000-0000-0000-0000-000000000000/cellranger_output"	
run_mkfastq	If you want to run `cellranger-atac mkfastq`	true	true
run_count	If you want to run `cellranger-atac count`	true	true
delete_input_bcl_directory	If true, delete BCL directories after demux. If false, you should delete this folder yourself so as to not incur storage charges	false	false
force_cells	Force pipeline to use this number of cells, bypassing the cell detection algorithm	6000	
cellranger_atac_version	cellranger-atac version, currently only 1.1.0	"1.1.0"	"1.1.0"
docker_registry	Docker registry to use for cellranger_workflow. Options: • "cumulusprod/" for Docker Hub images; • "quay.io/cumulus/" for backup images on Red Hat registry.	"cumulusprod/"	"cumulusprod/"
zones	Google cloud zones	"us-central1-a us-west1-a"	"us-central1-a us-central1-b us-central1-c us-central1-f us-east1-b us-east1-c us-east1-d us-west1-a us-west1-b us-west1-c"
atac_num_core	Number of cpus for cellranger-atac count	64	64
atac_memory	Memory string for cellranger-atac count	"57.6G"	"57.6G"
mkfastq_disk_space	Optional disk space in GB for cellranger-atac mkfastq	1500	1500
atac_disk_space	Disk space in GB needed for cellranger-atac count	500	500
preemptible	Number of preemptible tries	2	2

Workflow output

See the table below for important scATAC-seq outputs.

Name	Type	Description
output_fastqs_directory	Array[String]	A list of google bucket urls containing FASTQ files, one url per flowcell.
output_count_directory	Array[String]	A list of google bucket urls containing cellranger-atac count outputs, one url per sample.
metrics_summaries	File	A excel spreadsheet containing QCs for each sample.
output_web_summary	Array[File]	A list of htmls visualizing QCs for each sample (cellranger count output).
count_matrix	String	gs url for a template count_matrix.csv to run cumulus.

9.3.5 Single-cell immune profiling

To process single-cell immune profiling (scIR-seq) data, follow the specific instructions below.

Sample sheet

1. **Reference** column.

   Pre-built scIR-seq references are summarized below.

Keyword	Description
GRCh38_vdj_v3.1.0	Human GRCh38 V(D)J sequences, cellranger reference 3.1.0, annotation built from Ensembl *Homo_sapiens.GRCh38.94.chr_patch_hapl_scaff.gtf*
GRCm38_vdj_v3.1.0	Mouse GRCm38 V(D)J sequences, cellranger reference 3.1.0, annotation built from Ensembl *Mus_musculus.GRCm38.94.gtf*
GRCh38_vdj_v2.0.0 or **GRCh38_vdj**	Human GRCh38 V(D)J sequences, cellranger reference 2.0.0, annotation built from Ensembl *Homo_sapiens.GRCh38.87.chr_patch_hapl_scaff.gtf* and *vdj_GRCh38_alts_ensembl_10x_genes-2.0.0.gtf*
GRCm38_vdj_v2.2.0 or **GRCm38_vdj**	Mouse GRCm38 V(D)J sequences, cellranger reference 2.2.0, annotation built from Ensembl *Mus_musculus.GRCm38.90.chr_patch_hapl_scaff.gtf*

2. **Index** column.

   Put 10x single cell V(D)J sample index set names (e.g. SI-GA-A3) here.

3. *Chemistry* column.

   This column is not used for scIR-seq data. Put **fiveprime** here as a placeholder if you decide to include the Chemistry column.

4. *DataType* column.

   Set it to **vdj**.

5. *FetureBarcodeFile* column.

   Leave it blank for scIR-seq.

6. Example:

```
Sample,Reference,Flowcell,Lane,Index,Chemistry,DataType
sample_vdj,GRCh38_vdj_v3.1.0,gs://fc-e0000000-0000-0000-0000-000000000000/
↪VK10WBC9ZZ,1,SI-GA-A1,fiveprime,vdj
```

Workflow input

For scIR-seq data, `cellranger_workflow` takes Illumina outputs as input and runs `cellranger mkfastq` and `cellranger vdj`. Revalant workflow inputs are described below, with required inputs highlighted in bold.

Name	Description	Example	Default
input_csv_file	Sample Sheet (contains Sample, Reference, Flowcell, Lane, Index as required and Chemistry, DataType, FeatureBarcodeFile as optional)	"gs://fc-e0000000-0000-0000-0000-000000000000/sample_sheet.csv"	
output_directory	Output directory	"gs://fc-e0000000-0000-0000-0000-000000000000/cellranger_output"	
run_mkfastq	If you want to run `cellranger mkfastq`	true	true
delete_input_bcl_directory	Delete input BCL directories after demux. If false, you should delete this folder yourself so as to not incur storage charges	false	false
force_cells	Force pipeline to use this number of cells, bypassing the cell detection algorithm	6000	
vdj_denovo	Do not align reads to reference V(D)J sequences before de novo assembly	false	false
vdj_chain	Force the web summary HTML and metrics summary CSV to only report on a particular chain type. The accepted values are: auto for autodetection based on TR vs IG representation, TR for T cell receptors, IG for B cell receptors, all for all chain types	TR	
cellranger_version	Cellranger version, could be 3.1.0, 3.0.2, 2.2.0	"3.1.0"	"3.1.0"
docker_registry	Docker registry to use for cellranger_workflow. Options: • "cumulusprod/" for Docker Hub images; • "quay.io/cumulus/" for backup images on Red Hat registry.	"cumulusprod/"	"cumulusprod/"
cellranger_mkfastq_docker_registry	Docker registry to use for `cellranger mkfastq`. Default is the registry to which only Broad users have access. See *bcl2fastq* for making your own registry.	"gcr.io/broad-cumulus"	"gcr.io/broad-cumulus"
zones	Google cloud zones	"us-central1-a us-west1-a"	"us-central1-a us-central1-b us-central1-c us-central1-f us-east1-b us-east1-c us-east1-d us-west1-a us-west1-b us-west1-c"
num_cpu	Number of cpus to request for one node for cellranger mkfastq and cellranger vdj	32	32
memory	Memory size string for cellranger mkfastq and cellranger vdj	"120G"	"120G"
mkfastq_disk_space	Optional disk space in GB for mkfastq	1500	1500
vdj_disk_space	Disk space in GB needed for cellranger vdj	500	500
preemptible	Number of preemptible tries	2	2

Workflow output

See the table below for important scIR-seq outputs.

Name	Type	Description
output_fastqs_directory	Array[String]	A list of google bucket urls containing FASTQ files, one url per flowcell.
output_vdj_directory	Array[String]	A list of google bucket urls containing vdj results, one url per sample.
metrics_summaries	File	A excel spreadsheet containing QCs for each sample.
output_web_summary	Array[File]	A list of htmls visualizing QCs for each sample (cellranger count output).
count_matrix	String	gs url for a template count_matrix.csv to run cumulus.

9.4 bcl2fastq

9.4.1 License

bcl2fastq license

9.4.2 Docker

Read this tutorial if you are new to Docker and don't know how to write your Dockerfile.

For a Debian based docker, add the lines below into its Dockerfile to install `bcl2fastq`:

```
RUN apt-get install --no-install-recommends -y alien unzip
ADD https://support.illumina.com/content/dam/illumina-support/documents/downloads/
↪software/bcl2fastq/bcl2fastq2-v2-20-0-linux-x86-64.zip /software
RUN unzip -d /software/ /software/bcl2fastq2-v2-20-0-linux-x86-64.zip && alien -i /
↪software/bcl2fastq2-v2.20.0.422-Linux-x86_64.rpm && rm /software/bcl2fastq2-v2*
```

You can host your private docker images in the Google Container Registry.

9.4.3 Workflows

Workflows such as **cellranger_workflow** and **dropseq_workflow** provide the option of running `bcl2fastq`. We provide dockers containing `bcl2fastq` that are accessible only by members of the Broad Institute. Non-Broad Institute members will have to provide their own docker images.

9.4.4 Example

In this example we create a docker image for running `cellranger mkfastq` version 3.0.2.

1. Create a GCP project or reuse an existing project.
2. Enable the Google Container Registry
3. Clone the cumulus repository:

```
git clone https://github.com/klarman-cell-observatory/cumulus.git
```

4. Add the lines to cumulus/docker/cellranger/3.0.2/Dockerfile to include bcl2fastq (see *Docker*).
5. Ensure you have Docker installed
6. Download cellranger from https://support.10xgenomics.com/single-cell-gene-expression/software/downloads/3.0
7. Build, tag, and push the docker. Remember to replace PROJECT_ID with your GCP project id:

```
cd cumulus/docker/cellranger/3.0.2/
docker build -t cellranger-3.0.2 .
docker tag cellranger-3.0.2 gcr.io/PROJECT_ID/cellranger:3.0.2
gcr.io/PROJECT_ID/cellranger:3.0.2
```

8. Import **cellranger_workflow** workflow to your workspace (see cellranger_workflow steps), and enter your docker registry URL (in this example, `"gcr.io/PROJECT_ID/"`) in `cellranger_mkfastq_docker_registry` field of cellranger_workflow inputs.

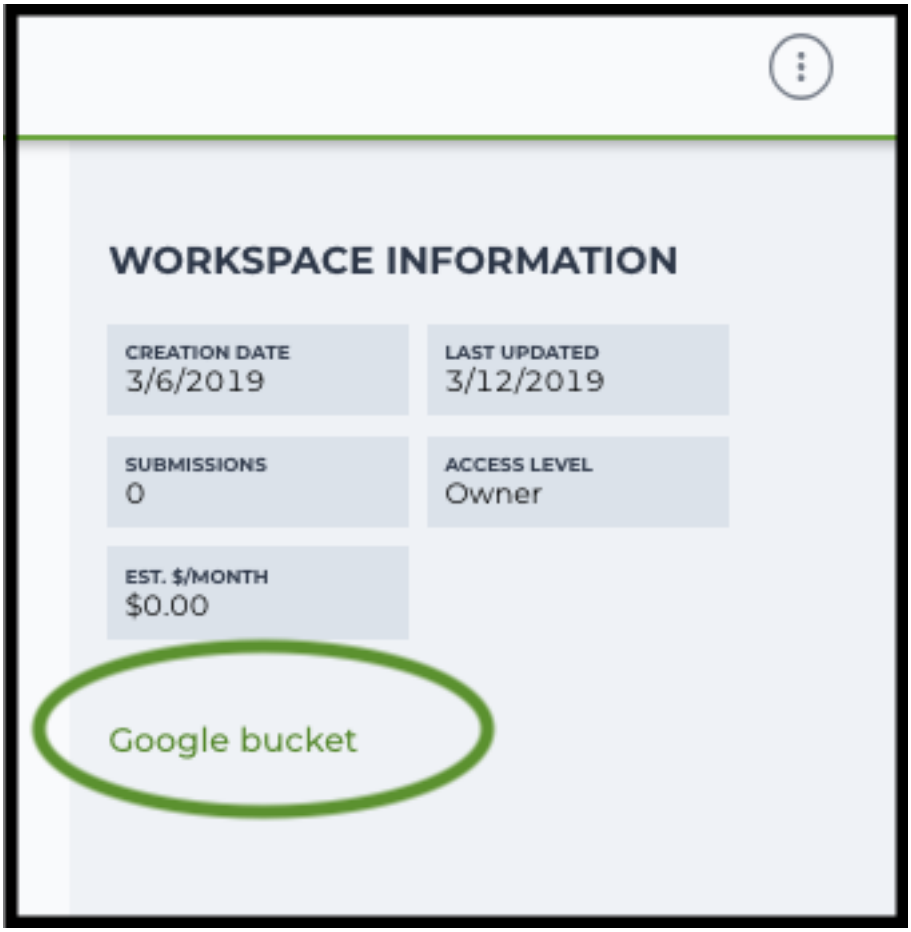
9.5 Extract gene-count matrices from plated-based SMART-Seq2 data

9.5.1 Run SMART-Seq2 Workflow

Follow the steps below to extract gene-count matrices from SMART-Seq2 data on Terra. This WDL aligns reads using *Bowtie 2* and estimates expression levels using *RSEM*.

1. Copy your sequencing output to your workspace bucket using gsutil in your unix terminal.

   You can obtain your bucket URL in the dashboard tab of your Terra workspace under the information panel.



Note: Broad users need to be on an UGER node (not a login node) in order to use the `-m` flag

Request an UGER node:

```
reuse UGER
qrsh -q interactive -l h_vmem=4g -pe smp 8 -binding linear:8 -P regevlab
```

The above command requests an interactive node with 4G memory per thread and 8 threads. Feel free to change the memory, thread, and project parameters.

Once you're connected to an UGER node, you can make gsutil available by running:

```
reuse Google-Cloud-SDK
```

Use `gsutil cp [OPTION]... src_url dst_url` to copy data to your workspace bucket. For example, the following command copies the directory at /foo/bar/nextseq/Data/VK18WBC6Z4 to a Google bucket:

```
gsutil -m cp -r /foo/bar/nextseq/Data/VK18WBC6Z4 gs://fc-e0000000-0000-
↪0000-0000-000000000000/VK18WBC6Z4
```

`-m` means copy in parallel, `-r` means copy the directory recursively.

2. Create a sample sheet.

   Please note that the columns in the CSV can be in any order, but that the column names must match the recognized headings.

The sample sheet provides metadata for each cell:

Column	Description
Cell	Cell name.
Plate	Plate name. Cells with the same plate name are from the same plate.
Read1	Location of the FASTQ file for read1 in the cloud (gsurl).
Read2	Location of the FASTQ file for read1 in the cloud (gsurl).

Example:

```
Cell,Plate,Read1,Read2
cell-1,plate-1,gs://fc-e0000000-0000-0000-0000-000000000000/smartseq2/
↪cell-1_L001_R1_001.fastq.gz,gs://fc-e0000000-0000-0000-0000-
↪000000000000/smartseq2/cell-1_L001_R2_001.fastq.gz
cell-2,plate-1,gs://fc-e0000000-0000-0000-0000-000000000000/smartseq2/
↪cell-2_L001_R1_001.fastq.gz,gs://fc-e0000000-0000-0000-0000-
↪000000000000/smartseq2/cell-2_L001_R2_001.fastq.gz
cell-3,plate-2,gs://fc-e0000000-0000-0000-0000-000000000000/smartseq2/
↪cell-3_L001_R1_001.fastq.gz,gs://fc-e0000000-0000-0000-0000-
↪000000000000/smartseq2/cell-3_L001_R2_001.fastq.gz
cell-4,plate-2,gs://fc-e0000000-0000-0000-0000-000000000000/smartseq2/
↪cell-4_L001_R1_001.fastq.gz,gs://fc-e0000000-0000-0000-0000-
↪000000000000/smartseq2/cell-4_L001_R2_001.fastq.gz
```

3. Upload your sample sheet to the workspace bucket.

   Example:

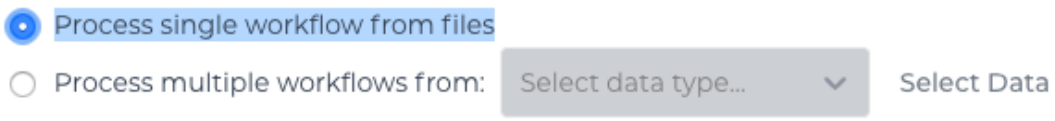
   ```
   gsutil cp /foo/bar/projects/sample_sheet.csv gs://fc-e0000000-0000-0000-
   ↪0000-000000000000/
   ```

4. Import *smartseq2* workflow to your workspace.

   See the Terra documentation for adding a workflow. The *smartseq2* workflow is under `Broad Methods Repository` with name "**cumulus/smartseq2**".

   Moreover, in the workflow page, click `Export to Workspace...` button, and select the workspace to which you want to export *smartseq2* workflow in the drop-down menu.

5. In your workspace, open `smartseq2` in `WORKFLOWS` tab. Select `Process single workflow from files` as below

   and click `SAVE` button.

Inputs:

Please see the description of inputs below. Note that required inputs are shown in bold.

Name	Description	Example	Default
input_csv_file	Sample Sheet (contains Cell, Plate, Read1, Read2)	"gs://fc-e0000000-0000-0000-0000-000000000000/sample_sheet.csv"	
output_directory	Output directory	"gs://fc-e0000000-0000-0000-0000-000000000000/smartseq2_output"	
reference	Reference transcriptome to align reads to. Acceptable values: • Pre-created genome references: "GRCh38" for human; "GRCm38" and "mm10" for mouse. • Create a custom genome reference using smartseq2_create_reference workflow, and specify its Google bucket URL here.	"GRCh38", or "gs://fc-e0000000-0000-0000-0000-000000000000/rsem_ref.tar.gz"	
smartseq2_version	SMART-Seq2 version to use. Versions available: 1.0.0.	"1.0.0"	"1.0.0"
docker_registry	Docker registry to use. Options: • "cumulusprod/" for Docker Hub images; • "quay.io/cumulus/" for backup images on Red Hat registry.	"cumulusprod/"	"cumulusprod/"
zones	Google cloud zones	"us-east1-d us-west1-a us-west1-b"	"us-east1-d us-west1-a us-west1-b"
num_cpu	Number of cpus to request for one node	4	4
memory	Memory size string	"3.60G"	"3.60G"
disk_space	Disk space in GB	10	10
preemptible	Number of preemptible tries	2	2

Outputs:

See the table below for important outputs.

Name	Type	Description
output_count_matrix	Array[String]	A list of google bucket urls containing gene-count matrices, one per plate. Each gene-count matrix file has the suffix `.dge.txt.gz`.

This WDL generates one gene-count matrix per SMART-Seq2 plate. The gene-count matrix uses Drop-Seq format:

- The first line starts with `"Gene"` and then gives cell barcodes separated by tabs.
- Starting from the second line, each line describes one gene. The first item in the line is the gene name and the rest items are TPM-normalized count values of this gene for each cell.

The gene-count matrices can be fed directly into **cumulus** for downstream analysis.

TPM-normalized counts are calculated as follows:

1. Estimate the gene expression levels in TPM using *RSEM*.
2. Suppose `c` reads are achieved for one cell, then calculate TPM-normalized count for gene `i` as `TPM_i / 1e6 * c`.

TPM-normalized counts reflect both the relative expression levels and the cell sequencing depth.

9.5.2 Custom Genome

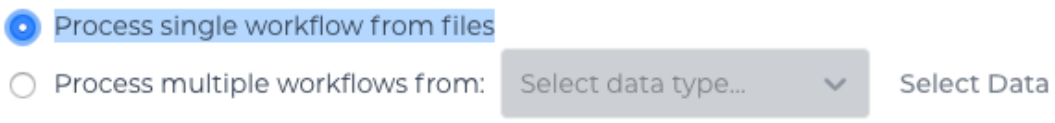
We also provide a way of generating user-customized Genome references for SMART-Seq2 workflow.

1. Import smartseq2_create_reference workflow to your workspace.

   See the Terra documentation for adding a workflow. The smartseq2_create_reference workflow is under `Broad Methods Repository` with name "**cumulus/smartseq2_create_reference**".

   Moreover, in the workflow page, click `Export to Workflow...` button, and select the workspace to which you want to export `smartseq2_create_reference` in the drop-down menu.

2. In your workspace, open `smartseq2_create_reference` in `WORKFLOWS` tab. Select `Process single workflow from files` as below

   and click `SAVE` button.

Inputs:

Please see the description of inputs below. Note that required inputs are shown in bold.

Name	Description	Type or Example	Default
fasta	Genome fasta file	File. For example, "gs://fc-e0000000-0000-0000-0000-000000000000/Homo_sapiens.GRCh38.dna.primary_assembly.fa"	
gtf	GTF gene annotation file (e.g. Homo_sapiens.GRCh38.83.gtf)	File. For example, "gs://fc-e0000000-0000-0000-0000-000000000000/Homo_sapiens.GRCh38.83.gtf"	
smartseq2_version	SMART-Seq2 version to use. Versions available: 1.0.0.	String	"1.0.0"
docker_registry	Docker registry to use. Options: • "cumulusprod/" for Docker Hub images; • "quay.io/cumulus/" for backup images on Red Hat registry.	String	"cumulusprod/"
zones	Google cloud zones	String	"us-east1-b us-east1-c us-east1-d"
cpu	Number of CPUs	Integer	8
memory	Memory size string	String	"7.2G"
extra_disk_space	Extra disk space in GB	Integer	15
preemptible	Number of preemptible tries	Integer	2

Outputs

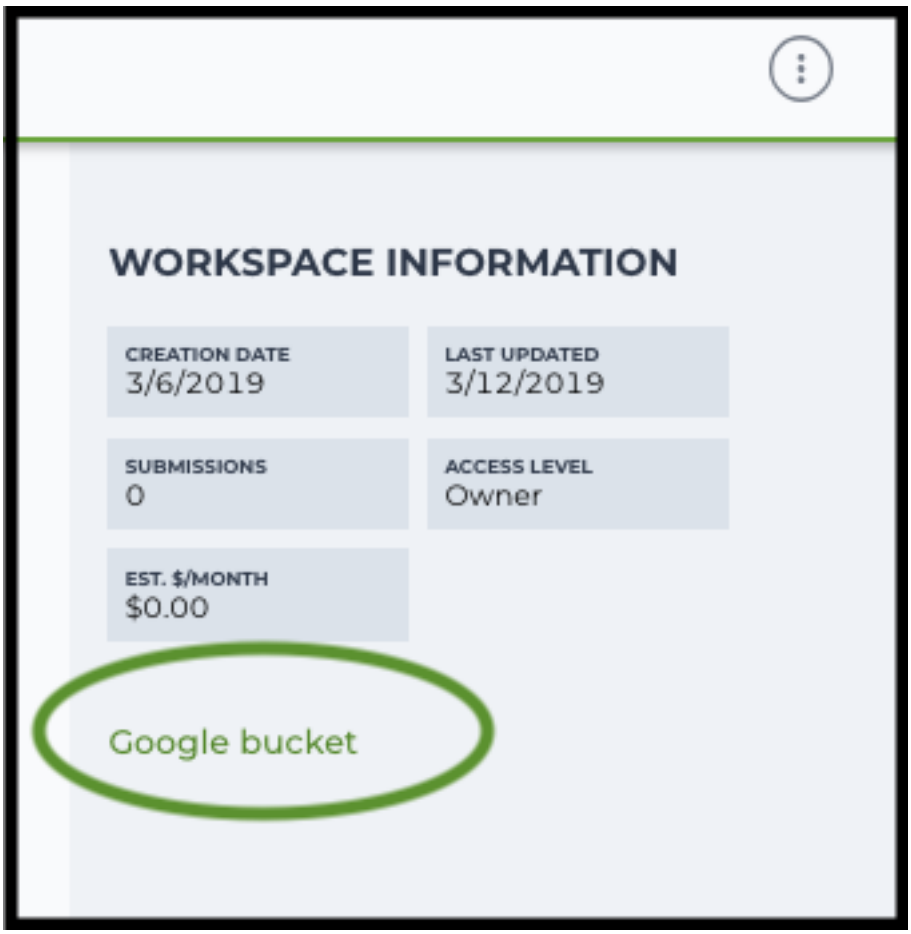
Name	Type	Description
reference	File	The custom Genome reference generated. Its default file name is `rsem_ref.tar.gz`.

9.6 Drop-seq pipeline

This workflow follows the steps outlined in the Drop-seq alignment cookbook from the McCarroll lab , except the default STAR aligner flags are *–limitOutSJcollapsed 1000000 –twopassMode Basic*. Additionally the pipeline provides the option to generate count matrices using dropEst.

1. Copy your sequencing output to your workspace bucket using gsutil in your unix terminal.

   You can obtain your bucket URL in the dashboard tab of your Terra workspace under the information panel.



   Note: Broad users need to be on an UGER node (not a login node) in order to use the `-m` flag

   Request an UGER node:

```
reuse UGER
qrsh -q interactive -l h_vmem=4g -pe smp 8 -binding linear:8 -P regevlab
```

   The above command requests an interactive node with 4G memory per thread and 8 threads. Feel free to change the memory, thread, and project parameters.

   Once you're connected to an UGER node, you can make gsutil available by running:

```
reuse Google-Cloud-SDK
```

   Use `gsutil cp [OPTION]... src_url dst_url` to copy data to your workspace bucket. For example, the following command copies the directory at /foo/bar/nextseq/Data/VK18WBC6Z4 to a Google bucket:

```
gsutil -m cp -r /foo/bar/nextseq/Data/VK18WBC6Z4 gs://fc-e0000000-0000-
↪0000-0000-000000000000/VK18WBC6Z4
```

   `-m` means copy in parallel, `-r` means copy the directory recursively.

2. Non Broad Institute users that wish to run bcl2fastq must create a custom docker image.

   See *bcl2fastq* instructions.

3. Create a sample sheet.

   Please note that the columns in the CSV must be in the order shown below and does not contain a header line. The sample sheet provides either the FASTQ files for each sample if you've already run bcl2fastq or a list of BCL directories if you're starting from BCL directories. Please note that BCL directories must contain a valid bcl2fastq sample sheet (SampleSheet.csv):

Column	Description
Name	Sample name.
Read1	Location of the FASTQ file for read1 in the cloud (gsurl).
Read2	Location of the FASTQ file for read2 in the cloud (gsurl).

   Example using FASTQ input files:

```
sample-1,gs://fc-e0000000-0000-0000-0000-000000000000/dropseq-1/sample1-1_
↪L001_R1_001.fastq.gz,gs://fc-e0000000-0000-0000-0000-000000000000/
↪dropseq-1/sample-1_L001_R2_001.fastq.gz
sample-2,gs://fc-e0000000-0000-0000-0000-000000000000/dropseq-1/sample-2_
↪L001_R1_001.fastq.gz,gs://fc-e0000000-0000-0000-0000-000000000000/
↪dropseq-1/sample-2_L001_R2_001.fastq.gz
sample-1,gs://fc-e0000000-0000-0000-0000-000000000000/dropseq-2/sample1-1_
↪L001_R1_001.fastq.gz,gs://fc-e0000000-0000-0000-0000-000000000000/
↪dropseq-2/sample-1_L001_R2_001.fastq.gz
```

   Note that in this example, sample-1 was sequenced across two flowcells.

   Example using BCL input directories:

```
gs://fc-e0000000-0000-0000-0000-000000000000/flowcell-1
gs://fc-e0000000-0000-0000-0000-000000000000/flowcell-2
```

   Note that the flow cell directory must contain a bcl2fastq sample sheet named SampleSheet.csv.

4. Upload your sample sheet to the workspace bucket.

   Example:

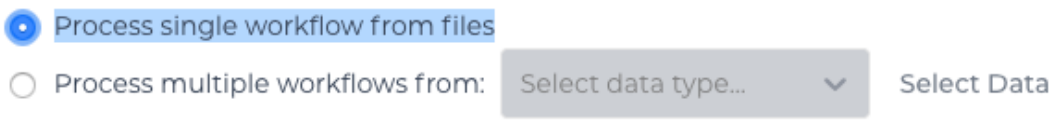
```
gsutil cp /foo/bar/projects/sample_sheet.csv gs://fc-e0000000-0000-0000-
↪0000-000000000000/
```

5. Import *dropseq_workflow* workflow to your workspace.

   See the Terra documentation for adding a workflow. The *dropseq_workflow* is under `Broad Methods Repository` with name "**cumulus/dropseq_workflow**".

   Moreover, in the workflow page, click the `Export to Workspace...` button, and select the workspace you want to export *dropseq_workflow* workflow in the drop-down menu.

6. In your workspace, open `dropseq_workflow` in `WORKFLOWS` tab. Select `Process single workflow from files` as below



and click the SAVE button.

9.6.1 Inputs

Please see the description of important inputs below.

Name	Description
input_csv_file	CSV file containing sample name, read1, and read2 or a list of BCL directories.
output_directory	Pipeline output directory (gs URL e.g. "gs://fc-e0000000-0000-0000-0000-000000000000/dropseq_output")
reference	hg19, GRCh38, mm10, hg19_mm10, mmul_8.0.1 or a path to a custom reference JSON file
run_bcl2fastq	Whether your sample sheet contains one BCL directory per line or one sample per line (default false)
run_dropseq_tools	Whether to generate count matrixes using Drop-Seq tools from the McCarroll lab (default true)
run_dropest	Whether to generate count matrixes using dropEst (default false)
cellular_barcodes_whitelist	Optional whitelist of known cellular barcodes
drop_seq_tools_force_cells	If supplied, bypass the cell detection algorithm (the elbow method) and use this number of cells.
dropest_cells_max	Maximal number of output cells
dropest_genes_min	Minimal number of genes for cells after the merge procedure (default 100)
dropest_min_merge_threshold	The threshold for the merge procedure (default 0.2)
dropest_max_cb_merge_edit_distance	Max edit distance between barcodes (default 2)
dropest_max_umi_merge_edit_distance	Max edit distance between UMIs (default 1)
dropest_min_genes_before_merge	Minimal number of genes for cells before the merge procedure. Used mostly for optimization. (default 10)
dropest_merge_barcodes_precise	Use precise merge strategy (can be slow), recommended to use when the list of real barcodes is not available (default true)
dropest_velocyto	Save separate count matrices for exons, introns and exon/intron spanning reads (default true)
trim_sequence	The sequence to look for at the start of reads for trimming (default "AAGCAGTGGTATCAACGCAGAGTGAATGGG")
trim_num_bases	How many bases at the beginning of the sequence must match before trimming occur (default 5)
umi_base_range	The base location of the molecular barcode (default 13-20)
cellular_barcode_base_range	The base location of the cell barcode (default 1-12)
star_flags	Additional options to pass to STAR aligner

Please note that run_bcl2fastq must be set to true if you're starting from BCL files instead of FASTQs.

Custom Genome JSON

If you're reference is not one of the predefined choices, you can create a custom JSON file. Example:

```
{
        "refflat":        "gs://fc-e0000000-0000-0000-0000-000000000000/human_mouse/
↪hg19_mm10_transgenes.refFlat",
        "genome_fasta":    "gs://fc-e0000000-0000-0000-0000-000000000000/human_mouse/
↪hg19_mm10_transgenes.fasta",
        "star_genome":    "gs://fc-e0000000-0000-0000-0000-000000000000/human_mouse/
↪STAR2_5_index_hg19_mm10.tar.gz",
        "gene_intervals":        "gs://fc-e0000000-0000-0000-0000-000000000000/human_
↪mouse/hg19_mm10_transgenes.genes.intervals",
        "genome_dict":    "gs://fc-e0000000-0000-0000-0000-000000000000/human_mouse/
↪hg19_mm10_transgenes.dict",
```

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```
        "star_cpus": 32,
        "star_memory": "120G"
}
```

The fields star_cpus and star_memory are optional and are used as the default cpus and memory for running STAR with your genome.

9.6.2 Outputs

The pipeline outputs a list of google bucket urls containing one gene-count matrix per sample. Each gene-count matrix file produced by Drop-seq tools has the suffix 'dge.txt.gz', matrices produced by dropEst have the extension .rds.

Building a Custom Genome

The tool **dropseq_bundle** can be used to build a custom genome. Please see the description of important inputs below.

Name	Description
fasta_file	Array of fasta files. If more than one species, fasta and gtf files must be in the same order.
gtf_file	Array of gtf files. If more than one species, fasta and gtf files must be in the same order.
genomeSAindexNbases	Length (bases) of the SA pre-indexing string. Typically between 10 and 15. Longer strings will use much more memory, but allow faster searches. For small genomes, must be scaled down to min(14, log2(GenomeLength)/2 - 1)

dropseq_workflow Terra Release Notes

Version 5

- Split preprocessing steps into separate tasks (FastqToSam, TagBam, FilterBam, and TrimBam).

Version 4

- Handle uncompressed fastq files as workflow input.
- Added optional prepare_fastq_disk_space_multiplier input.

Version 3

- Set default value for docker_registry input.

Version 2

- Added docker_registry input.

Version 1

- Renamed cumulus to cumulus
- Added use_bases_mask option when running bcl2fastq

Version 18

- Created a separate docker image for running bcl2fastq

Version 17

- Fixed bug that ignored WDL input star_flags (thanks to Carly Ziegler for reporting)

- Changed default value of star_flags to the empty string (Prior versions of the WDL incorrectly indicated that basic 2-pass mapping was done)

Version 16

- Use cumulus dockerhub organization
- Changed default dropEst version to 0.8.6

Version 15

- Added drop_deq_tools_prep_bam_memory and drop_deq_tools_dge_memory options

Version 14

- Fix for downloading files from user pays buckets

Version 13

- Set GCLOUD_PROJECT_ID for user pays buckets

Version 12

- Changed default dropEst memory from 52G to 104G

Version 11

- Updated formula for computing disk size for dropseq_count

Version 10

- Added option to specify merge_bam_alignment_memory and sort_bam_max_records_in_ram

Version 9

- Updated default drop_seq_tools_version from 2.2.0 to 2.3.0

Version 8

- Made additional options available for running dropEst

Version 7

- Changed default dropEst memory from 104G to 52G

Version 6

- Added option to run dropEst

Version 5

- Specify full version for bcl2fastq (2.20.0.422-2 instead of 2.20.0.422)

Version 4

- Fixed issue that prevented bcl2fastq from running

Version 3

- Set default run_bcl2fastq to false
- Create shortcuts for commonly used genomes

Version 2

- Updated QC report

Version 1

- Initial release

9.7 Demultiplex cell-hashing/nuclei-hashing data using demuxEM or prepare for CITE-Seq analysis

Follow the steps below to run **cumulus** for cell-hashing/nuclei-hashing/CITE-Seq data on Terra.

1. Run Cell Ranger tool to generate raw gene count matrices and antibody hashtag data.

   Please refer to the cellranger_workflow tutorial for details.

   When finished, you should be able to find the raw gene count matrix (e.g. `raw_gene_be_matrices_h5.h5`) and ADT count matrix (e.g. `sample_1_ADT.csv`) for each sample.

2. Create a sample sheet, **sample_sheet_hashing.csv**, which describes the metadata for each pair of RNA and antibody hashtag data. The sample sheet should contain 4 columns — *OUTNAME*, *RNA*, *ADT*, and *TYPE*. *OUTNAME* is the output name for one pair of RNA and ADT data. *RNA* and *ADT* are the raw gene count matrix and the ADT count matrix generated in Step 1, respectively. *TYPE* is the assay type, which can be `cell-hashing`, `nuclei-hashing`, or `cite-seq`.

   Example:

   ```
   OUTNAME,RNA,ADT,TYPE
   sample_1,gs://fc-e0000000-0000-0000-0000-000000000000/my_dir/sample_1/raw_
   ↪gene_bc_matrices_h5.h5,gs://fc-e0000000-0000-0000-0000-000000000000/my_
   ↪dir/sample_1_ADT/sample_1_ADT.csv,cell-hashing
   sample_2,gs://fc-e0000000-0000-0000-0000-000000000000/my_dir/sample_2/raw_
   ↪feature_bc_matrices.h5,gs://fc-e0000000-0000-0000-0000-000000000000/my_
   ↪dir/sample_2_ADT/sample_2_ADT.csv,nuclei-hashing
   ```

   Note that in the example above, sample_2 is 10x genomics' v3 chemistry. Cumulus can automatically detect v2/v3 chemistry when loading hdf5 files.

3. Create an additional antibody-control sheet **antibody_control.csv** if you have CITE-Seq data. This sheet contains 2 columns — *Antibody* and *Control*.

   Example:

   ```
   Antibody,Control
   CD8,Mouse-IgG1
   HLA-ABC,Mouse-IgG2a
   CD45RA,Mouse-IgG2b
   ```

4. Upload your sample sheets to the Google bucket of your workspace.

   Example:

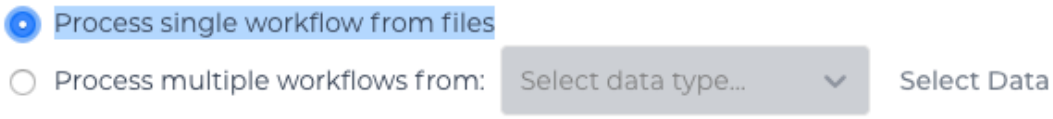
   ```
   gsutil cp /foo/bar/projects/my_sample_sheet_hashing.csv gs://fc-e0000000-
   ↪0000-0000-0000-000000000000/
   gsutil cp /foo/bar/projects/antibody_control.csv gs://fc-e0000000-0000-
   ↪0000-0000-000000000000/
   ```

5. Import *cumulus_hashing_cite_seq* to your workspace.

   See the Terra documentation for adding a workflow. The *cumulus_hashing_cite_seq* workflow is under `Broad Methods Repository` with name "**cumulus/cumulus_hashing_cite_seq**".

   Moreover, in the workflow page, click the `Export to Workspace...` button, and select the workspace to which you want to export *cumulus_hashing_cite_seq* workflow in the drop-down menu.

6. In your workspace, open `cumulus_hashing_cite_seq` in `WORKFLOWS` tab. Select `Process single workflow from files` as below



   and click the `SAVE` button.

9.7.1 cumulus_hashing_cite_seq inputs:

Name	Description	Example	Default
input_sample_sheet	Input CSV file describing metadata of RNA and ADT data pairing	"gs://fc-e0000000-0000-0000-0000-000000000000/sample_sheet_hashing.csv"	
output_directory	This is the output directory (gs url + path) for all results. There will be one folder per RNA-ADT data pair under this directory	"gs://fc-e0000000-0000-0000-0000-000000000000/my_demux_dir"	
genome	Reference genome name. If not provided, cumulus will infer the genome name from data	"GRCh38"	
demuxEM_min_num_genes	demuxEM parameter. Only demultiplex cells/nuclei with at least <demuxEM_min_num_genes> expressed genes	200	100
demuxEM_alpha_on_samples	demuxEM parameter. The Dirichlet prior concentration parameter (alpha) on samples. An alpha value < 1.0 will make the prior sparse.	2.0	0.0
demuxEM_min_num_umis	demuxEM parameter. Only demultiplex cells/nuclei with at least <demuxEM_min_num_umis> of UMIs.	200	100
demuxEM_min_signal_hashtag	demuxEM parameter. Any cell/nucleus with less than <count> hashtags from the signal will be marked as unknown. [default: 10.0]	10.0	10.0
demuxEM_random_state	demuxEM parameter. The random seed used in the KMeans algorithm to separate empty ADT droplets from others	0	0
demuxEM_generate_diagnostic_plots	demuxEM parameter. If generate a series of diagnostic plots, including the background/signal between HTO counts, estimated background probabilities, HTO distributions of cells and non-cells etc	true	true
demuxEM_generate_gender_plot	demuxEM parameter. If generate violin plots using gender-specific genes (e.g. Xist). <demuxEM_generate_gender_plot> is a comma-separated list of gene names	"XIST"	
antibody_control_csv	cellranger rna_adt parameter. This is a CSV file containing the IgG control information for each antibody.	"gs://fc-e0000000-0000-0000-0000-000000000000/antibody_control.csv"	
cumulus_version	cumulus version to use. Versions available: 0.11.0, 0.10.0.	"0.11.0"	"0.11.0"
docker_registry	Docker registry to use. Options: • "cumulusprod/" for Docker Hub images; • "quay.io/cumulus/" for backup images on Red Hat registry.	"cumulusprod/"	"cumulusprod/"
zones	Google cloud zones	"us-east1-d us-west1-a us-west1-b"	"us-central1-a us-central1-b us-central1-c us-central1-f us-east1-b us-east1-c us-east1-d us-west1-a us-west1-b us-west1-c"
num_cpu	Number of CPUs per cumulus_hashing_cite_seq job	8	8

9.7.2 cumulus_hashing_cite_seq outputs

See the table below for important *cumulus_hashing_cite_seq* outputs:

Name	Type	Description
output_folder	Array[String]	A list of google bucket urls containing results for every RNA-ADT data pairs.

In the output folder of each cell-hashing/nuclei-hashing RNA-ADT data pair, you can find the following files:

Name	Description
output_name_demux.h5ad	Demultiplexed RNA count matrix in h5ad format.
output_name_demux.h5sc	RNA expression matrix with demultiplexed sample identities in cumulus hdf5 (h5sc) format.
output_name_ADTs.h5ad	Antibody tag matrix in h5ad format.
output_name.ambient_hashtag.hist.png	Optional output. A histogram plot depicting hashtag distributions of empty droplets and non-empty droplets.
output_name.background_probabilities.png	Optional output. A bar plot visualizing the estimated hashtag background probability distribution.
output_name.real_content.hist.png	Optional output. A histogram plot depicting hashtag distributions of not-real-cells and real-cells as defined by total number of expressed genes in the RNA assay.
output_name.rna_demux.hist.png	Optional output. A histogram plot depicting RNA UMI distribution for singlets, doublets and unknown cells.
output_name.gene_name.violin.png	Optional outputs. Violin plots depicting gender-specific gene expression across samples. We can have multiple plots if a gene list is provided in `demuxEM_generate_gender_plot` field of cumulus_hashing_cite_seq inputs.

In the output folder of each CITE-Seq RNA-ADT data pair, you can find the following file:

Name	Description
output_name.h5sc	A Cumulus hdf5 format (h5sc) file containing both RNA and ADT count matrices.

9.7.3 Load demultiplexing results into Python and R

To load demultiplexing results into Python, you need to install Python package anndata first. Then follow the codes below:

```
import anndata
data = anndata.read_h5ad('output_name_demux.h5ad')
```

Once you load the data object, you can find predicted droplet types (singlet/doublet/unknown) in `data.obs['demux_type']`. You can find predicted sample assignments in `data.obs['assignment']`. You

can find estimated sample fractions (sample1, sample2, ..., samplen, background) for each droplet in `data.obsm['raw_probs']`.

To load the results into R, you need to install R package `reticulate` in addition to Python package `anndata`. Then follow the codes below:

```
library(reticulate)
ad <- import("anndata", convert = FALSE)
data <- ad$read_h5ad("output_name_demux.h5ad")
```

Results are in `data$obs['demux_type']`, `data$obs['assignment']`, and `data$obsm['raw_probs']`.

9.8 Run Cumulus for sc/snRNA-Seq data analysis

9.8.1 Run Cumulus analysis

Prepare Input Data

Case One: Sample Sheet

Follow the steps below to run **cumulus** on Terra.

1. Create a sample sheet, **count_matrix.csv**, which describes the metadata for each sample count matrix. The sample sheet should at least contain 2 columns — *Sample* and *Location*. *Sample* refers to sample names and *Location* refers to the location of the channel-specific count matrix in either of

- 10x format with v2 chemistry. For example, `gs://fc-e0000000-0000-0000-0000-000000000000/my_dir/sample_1/filtered_gene_bc_matrices_h5.h5`.
- 10x format with v3 chemistry. For example, `gs://fc-e0000000-0000-0000-0000-000000000000/my_dir/sample_1/filtered_feature_bc_matrices.h5`.
- Drop-seq format. For example, `gs://fc-e0000000-0000-0000-0000-000000000000/my_dir/sample_2/sample_2.umi.dge.txt.gz`.
- mtx, csv, tsv or loom format.

Additionally, an optional Reference column can be used to select samples generated from a same reference (e.g. mm10). If the count matrix is in either DGE, mtx, csv, tsv, or loom format, the value in this column will be used as the reference since the count matrix file does not contain reference name information. In addition, the Reference column can be used to aggregate count matrices generated from different genome versions or gene annotations together under a unified reference. For example, if we have one matrix generated from mm9 and the other one generated from mm10, we can write mm9_10 for these two matrices in their Reference column. Pegasus will change their references to 'mm9_10' and use the union of gene symbols from the two matrices as the gene symbols of the aggregated matrix. For HDF5 files (e.g. 10x v2/v3), the reference name contained in the file does not need to match the value in this column. In fact, we use this column to rename references in HDF5 files. For example, if we have two HDF files, one generated from mm9 and the other generated from mm10. We can set these two files' Reference column value to 'mm9_10', which will rename their reference names into mm9_10 and the aggregated matrix will contain all genes from either mm9 or mm10. This renaming feature does not work if one HDF5 file contain multiple references (e.g. mm10 and GRCh38).

You are free to add any other columns and these columns will be used in selecting channels for futher analysis. In the example below, we have *Source*, which refers to the tissue of origin, *Platform*, which refers to the sequencing platform, *Donor*, which refers to the donor ID, and *Reference*, which refers to the reference genome.

Example:

```
Sample,Source,Platform,Donor,Reference,Location
sample_1,bone_marrow,NextSeq,1,GRCh38,gs://fc-e0000000-0000-0000-0000-
↪000000000000/my_dir/sample_1/filtered_gene_bc_matrices_h5.h5
sample_2,bone_marrow,NextSeq,2,GRCh38,gs://fc-e0000000-0000-0000-0000-
↪000000000000/my_dir/sample_2/filtered_gene_bc_matrices_h5.h5
sample_3,pbmc,NextSeq,1,GRCh38,gs://fc-e0000000-0000-0000-0000-000000000000/
↪my_dir/sample_3/filtered_feature_bc_matrices.h5
sample_4,pbmc,NextSeq,2,GRCh38,gs://fc-e0000000-0000-0000-0000-000000000000/
↪my_dir/sample_4/filtered_feature_bc_matrices.h5
```

If you ran **cellranger_workflow** ahead, you should already obtain a template **count_matrix.csv** file that you can modify from **generate_count_config**'s outputs.

1. Upload your sample sheet to the workspace.

   Example:

   ```
   gsutil cp /foo/bar/projects/my_count_matrix.csv gs://fc-e0000000-0000-
   ↪0000-0000-000000000000/
   ```

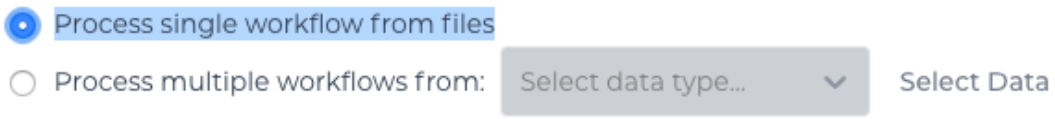
   where `/foo/bar/projects/my_count_matrix.csv` is the path to your sample sheet in local machine, and `gs://fc-e0000000-0000-0000-0000-000000000000/` is the location on Google bucket to hold it.

2. Import *cumulus* workflow to your workspace.

   See the Terra documentation for adding a workflow. The *cumulus* workflow is under `Broad Methods Repository` with name "**cumulus/cumulus**".

   Moreover, in the workflow page, click the `Export to Workspace...` button, and select the workspace to which you want to export *cumulus* workflow in the drop-down menu.

3. In your workspace, open `cumulus` in `WORKFLOWS` tab. Select `Process single workflow from files` as below

   and click the `SAVE` button.

Case Two: Single File

Alternatively, if you only have one single count matrix for analysis, you can go without sample sheets. **Cumulus** currently supports the following formats:

- 10x genomics v2/v3 formats (hdf5 or mtx);
- HCA DCP mtx and loom formats;
- Drop-seq dge formats.

Simply upload your data to the Google Bucket of your workspace, and specify its URL in `input_file` field of Cumulus' global inputs (see below). Notice that for dge and loom files, the `genome` field in global inputs is required.

In this case, the **aggregate_matrices** step will be skipped.

Cumulus steps:

Cumulus processes single cell data in the following steps:

1. **aggregate_matrices** (optional). When given a CSV format sample sheet, this step aggregates channel-specific count matrices into one big count matrix. Users can specify which channels they want to analyze and which sample attributes they want to import to the count matrix in this step. Otherwise, if a single count matrix file is given, skip this step.
2. **cluster**. This is the main analysis step. In this step, **Cumulus** performs low quality cell filtration, highly variable gene selection, batch correction, dimension reduction, diffusion map calculation, graph-based clustering and 2D visualization calculation (e.g. t-SNE/UMAP/FLE).
3. **de_analysis**. This step is optional. In this step, **Cumulus** can calculate potential markers for each cluster by performing a variety of differential expression (DE) analysis. The available DE tests include Welch's t test, Fisher's exact test, and Mann-Whitney U test. **Cumulus** can also calculate the area under ROC (AUROC) curve values for putative markers. If `find_markers_lightgbm` is on, **Cumulus** will try to identify cluster-specific markers by training a LightGBM classifier. If the samples are human or mouse immune cells, **Cumulus** can also optionally annotate putative cell types for each cluster based on known markers.
4. **plot**. This step is optional. In this step, **Cumulus** can generate 6 types of figures based on the **cluster** step results:
    - **composition** plots which are bar plots showing the cell compositions (from different conditions) for each cluster. This type of plots is useful to fast assess library quality and batch effects.
    - **tsne**, **fitsne**, and **net_tsne**: t-SNE like plots based on different algorithms, respectively. Users can specify cell attributes (e.g. cluster labels, conditions) for coloring side-by-side.
    - **umap** and **net_umap**: UMAP like plots based on different algorithms, respectively. Users can specify cell attributes (e.g. cluster labels, conditions) for coloring side-by-side.
    - **fle** and **net_fle**: FLE (Force-directed Layout Embedding) like plots based on different algorithms, respectively. Users can specify cell attributes (e.g. cluster labels, conditions) for coloring side-by-side.
    - **diffmap** plots which are 3D interactive plots showing the diffusion maps. The 3 coordinates are the first 3 PCs of all diffusion components.
    - If input is CITE-Seq data, there will be **citeseq_fitsne** plots which are FIt-SNE plots based on epitope expression.
5. **organize_results**. Copy analysis results from execution environment to destination location on Google bucket.

In the following sections, we will first introduce global inputs and then introduce the WDL inputs and outputs for each step separately. But please note that you need to set inputs from all steps simultaneously in the Terra WDL.

Note that we will make the required inputs/outputs bold and all other inputs/outputs are optional.

---

global inputs

Name	Description	Example	Default
input_file	Input CSV sample sheet describing metadata of each 10x channel, or a single input count matrix file	"gs://fc-e0000000-0000-0000-0000-000000000000/my_count_matrix.csv"	
output_name	This is the prefix for all output files. It should contain the google bucket url, subdirectory name and output name prefix	"gs://fc-e0000000-0000-0000-0000-000000000000/my_results_dir/my_results"	
cumulus_version	cumulus version to use. Versions available: 0.11.0, 0.10.0.	"0.11.0"	"0.11.0"
docker_registry	Docker registry to use. Options: • "cumulusprod/" for Docker Hub images; • "quay.io/cumulus/" for backup images on Red Hat registry.	"cumulusprod/"	"cumulusprod/"
zones	Google cloud zones to consider for execution.	"us-east1-d us-west1-a us-west1-b"	"us-central1-a us-central1-b us-central1-c us-central1-f us-east1-b us-east1-c us-east1-d us-west1-a us-west1-b us-west1-c"
num_cpu	Number of CPUs per Cumulus job	32	64
memory	Memory size string	"200G"	"200G"
disk_space	Total disk space in GB	100	100
preemptible	Number of preemptible tries	2	2

aggregate_matrices

aggregate_matrices inputs

Name	Description	Example	Default
restrictions	Select channels that satisfy all restrictions. Each restriction takes the format of name:value,. . . ,value. Multiple restrictions are separated by ';'	"Source:bone_marrow;Platform:NextSeq"	
attributes	Specify a comma-separated list of outputted attributes. These attributes should be column names in the count_matrix.csv file	"Source,Platform,Donor"	
default_reference	If sample count matrix is in either DGE, mtx, csv, tsv or loom format and there is no Reference column in the csv_file, use default_reference as the reference.	"GRCh38"	
select_only_singlets	If we have demultiplexed data, turning on this option will make cumulus only include barcodes that are predicted as singlets.	true	false
minimum_number_of_genes	Only keep barcodes with at least this number of expressed genes	100	100

aggregate_matrices output

Name	Type	Description
output_h5sc	File	Aggregated count matrix in Cumulus hdf5 (h5sc) format

---

cluster

cluster inputs

Name	Description	Example	Default
considered_refs	A string contains comma-separated reference(e.g. genome) names. Cumulus will read all groups associated with reference names in the list from the input file. If considered_refs is None, all groups will be considered.	"mm10"	
channel	Specify the cell barcode attribute to represent different samples.	"Donor"	
black_list	Cell barcode attributes in black list will be poped out. Format is "attr1,attr2,. . . ,attrn".	"attr1,attr2,attr3""	
min_genes_on_raw	If input are raw 10x matrix, which include all barcodes, perform a pre-filtration step to keep the data size small. In the pre-filtration step, only keep cells with at least <min_genes_on_raw> of genes	100	100

Continued on next page

Table 1 – continued from previous page

Name	Description	Example	Default
cite_seq	Data are CITE-Seq data. cumulus will perform analyses on RNA count matrix first. Then it will attach the ADT matrix to the RNA matrix with all antibody names changing to 'AD-' + antibody_name. Lastly, it will embed the antibody expression using FIt-SNE (the basis used for plotting is 'citeseq_fitsne')	false	false
cite_seq_capping	For CITE-Seq surface protein expression, make all cells with expression > <percentile> to the value at <percentile> to smooth outlier. Set <percentile> to 100.0 to turn this option off.	10.0	99.99
select_only_singlets	If we have demultiplexed data, turning on this option will make cumulus only include barcodes that are predicted as singlets	false	false
output_filtration_results	If write cell and gene filtration results to a spreadsheet	true	true
plot_filtration_results	If plot filtration results as PDF files	true	true
plot_filtration_figsize	Figure size for filtration plots. <figsize> is a comma-separated list of two numbers, the width and height of the figure (e.g. 6,4)	6,4	
output_seurat_compatible	If generate Seurat-compatible h5ad file. Caution: File size might be large, do not turn this option on for large data sets.	false	false
output_loom	If generate loom-formatted file	false	false
output_parquet	If generate parquet-formatted file	false	false
min_genes	Only keep cells with at least <min_genes> of genes	500	500
max_genes	Only keep cells with less than <max_genes> of genes	6000	6000
min_umis	Only keep cells with at least <min_umis> of UMIs	100	100
max_umis	Only keep cells with less than <max_umis> of UMIs	600000	600000
mito_prefix	Prefix of mitochondrial gene names. This is to identify mitochondrial genes.	"mt-"	"MT-"
percent_mito	Only keep cells with mitochondrial ratio less than <percent_mito>% of total counts	50	10.0
gene_percent_cells	Only use genes that are expressed in at <gene_percent_cells>% of cells to select variable genes	50	0.05
counts_per_cell_after	Total counts per cell after normalization, before transforming the count matrix into Log space.	1e5	1e5
select_hvf_flavor	Highly variable feature selection method. Options: • "pegasus": New selection method proposed in Pegasus, the analysis module of Cumulus workflow. • "Seurat": Conventional selection method used by Seurat and SCANPY.	"pegasus"	"pegasus"
select_hvf_ngenes	Select top <select_hvf_ngenes> highly variable features. If <select_hvf_flavor> is "Seurat" and <select_hvf_ngenes> is "None", select HVGs with z-score cutoff at 0.5.	2000	2000

Continued on next page

Table 1 – continued from previous page

Name	Description	Example	Default
no_select_hvf	Do not select highly variable features.	false	false
correct_batch_effect	If correct batch effects	false	false
batch_group_by	Batch correction assumes the differences in gene expression between channels are due to batch effects. However, in many cases, we know that channels can be partitioned into several groups and each group is biologically different from others. In this case, we will only perform batch correction for channels within each group. This option defines the groups. If <expression> is None, we assume all channels are from one group. Otherwise, groups are defined according to <expression>. <expression> takes the form of either 'attr', or 'attr1+attr2+...+attrn', or 'attr=value11,...,value1n_1;value21,...,value2n_2;...;valuem1,...,valuemn_m'. In the first form, 'attr' should be an existing sample attribute, and groups are defined by 'attr'. In the second form, 'attr1',...,'attrn' are n existing sample attributes and groups are defined by the Cartesian product of these n attributes. In the last form, there will be m + 1 groups. A cell belongs to group i (i > 0) if and only if its sample attribute 'attr' has a value among value i1,...,valuein_i. A cell belongs to group 0 if it does not belong to any other groups	"Donor"	None
random_state	Random number generator seed	0	0
nPC	Number of principal components	50	50
knn_K	Number of nearest neighbors used for constructing affinity matrix.	50	100
knn_full_speed	For the sake of reproducibility, we only run one thread for building kNN indices. Turn on this option will allow multiple threads to be used for index building. However, it will also reduce reproducibility due to the racing between multiple threads.	false	false
run_diffmap	Whether to calculate diffusion map or not. It will be automatically set to `true` when input **run_fle** or **run_net_fle** is set.	false	false
diffmap_ndc	Number of diffusion components	100	100
diffmap_maxt	Maximum time stamp in diffusion map computation to search for the knee point.	5000	5000
run_louvain	Run Louvain clustering algorithm	true	true
louvain_resolution	Resolution parameter for the Louvain clustering algorithm	1.3	1.3
louvain_class_label	Louvain cluster label name in analysis result.	"louvain_labels"	"louvain_labels"
run_leiden	Run Leiden clustering algorithm.	false	false

Continued on next page

Table 1 – continued from previous page

Name	Description	Example	Default
leiden_resolution	Resolution parameter for the Leiden clustering algorithm.	1.3	1.3
leiden_niter	Number of iterations of running the Leiden algorithm. If negative, run Leiden iteratively until no improvement.	2	-1
leiden_class_label	Leiden cluster label name in analysis result.	“leiden_labels”	“leiden_labels”
run_spectral_louvain	Run Spectral Louvain clustering algorithm	false	false
spectral_louvain_basis	Basis used for KMeans clustering. Use diffusion map by default. If diffusion map is not calculated, use PCA coordinates. Users can also specify “pca” to directly use PCA coordinates.	“diffmap”	“diffmap”
spectral_louvain_resolution	Resolution parameter for louvain.	1.3	1.3
spectral_louvain_class_label	Spectral louvain label name in analysis result.	“spectral_louvain_labels”	“spectral_louvain_labels”
run_spectral_leiden	Run Spectral Leiden clustering algorithm.	false	false
spectral_leiden_basis	Basis used for KMeans clustering. Use diffusion map by default. If diffusion map is not calculated, use PCA coordinates. Users can also specify “pca” to directly use PCA coordinates.	“diffmap”	“diffmap”
spectral_leiden_resolution	Resolution parameter for leiden.	1.3	1.3
spectral_leiden_class_label	Spectral leiden label name in analysis result.	“spectral_leiden_labels”	“spectral_leiden_labels”
run_tsne	Run multi-core t-SNE for visualization	false	false
tsne_perplexity	t-SNE’s perplexity parameter, also used by FIt-SNE.	30	30
run_fitsne	Run FIt-SNE for visualization	true	true
run_umap	Run UMAP for visualization	false	false
umap_K	K neighbors for UMAP.	15	15
umap_min_dist	UMAP parameter.	0.5	0.5
umap_spread	UMAP parameter.	1.0	1.0
run_fle	Run force-directed layout embedding (FLE) for visualization	false	false
fle_K	Number of neighbors for building graph for FLE	50	50
fle_target_change_per_node	Target change per node to stop FLE.	2.0	2.0
fle_target_steps	Maximum number of iterations before stopping the algoritm	5000	5000
net_down_sample_fraction	Down sampling fraction for net-related visualization	0.1	0.1
run_net_tsne	Run Net tSNE for visualization	false	false
net_tsne_out_basis	Basis name for Net t-SNE coordinates in analysis result	“net_tsne”	“net_tsne”
run_net_umap	Run Net UMAP for visualization	false	false
net_umap_out_basis	Basis name for Net UMAP coordinates in analysis result	“net_umap”	“net_umap”
run_net_fle	Run Net FLE for visualization	false	false
net_fle_out_basis	Basis name for Net FLE coordinates in analysis result.	“net_fle”	“net_fle”

cluster outputs

Name	Type	Description
output_h5ad	File	Output file in h5ad format (output_name.h5ad). To load this file in Python, you need to first install Pegasus on your local machine. Then use `import pegasus as pg; data = pg.read_input('output_name.h5ad')` in Python interpreter. The log-normalized expression matrix is stored in `data.X` as a CSR-format sparse matrix. The `obs` field contains cell related attributes, including clustering results. For example, `data.obs_names` records cell barcodes; `data.obs['Channel']` records the channel each cell comes from; `data.obs['n_genes']`, `data.obs['n_counts']`, and `data.obs['percent_mito']` record the number of expressed genes, total UMI count, and mitochondrial rate for each cell respectively; `data.obs['louvain_labels']`, `data.obs['leiden_labels']`, `data.obs['spectral_louvain_labels']`, and `data.obs['spectral_leiden_labels']` record each cell's cluster labels using different clustring algorithms; The `var` field contains gene related attributes. For example, `data.var_names` records gene symbols, `data.var['gene_ids']` records Ensembl gene IDs, and `data.var['highly_variable_features']` records selected variable genes. The `obsm` field records embedding coordinates. For example, `data.obsm['X_pca']` records PCA coordinates, `data.obsm['X_tsne']` records t-SNE coordinates, `data.obsm['X_umap']` records UMAP coordinates, `data.obsm['X_diffmap']` records diffusion map coordinates, `data.obsm['X_diffmap_pca']` records the first 3 PCs by projecting the diffusion components using PCA, and `data.obsm['X_fle']` records the force-directed layout coordinates from the diffusion components. The `varm` field records DE analysis results if performed: `data.varm['de_res']`. The `uns` field stores other related information, such as reference genome (`data.uns['genome']`), kNN on PCA coordinates (`data.uns['pca_knn_indices']` and `data.uns['pca_knn_distances']`), etc.
output_seurat_h5ad	File	h5ad file in seurat-compatible manner. This file can be loaded into R and converted into a Seurat object (see here for instructions)
output_filt_xlsx	File	Spreadsheet containing filtration results (output_name.filt.xlsx). This file has two sheets — Cell filtration stats and Gene filtration stats. The first sheet records cell filtering results and it has 10 columns: Channel, channel name; kept, number of cells kept; median_n_genes, median number of expressed genes in kept cells; median_n_umis, median number of UMIs in kept cells; median_percent_mito, median mitochondrial rate as UMIs between mitochondrial genes and all genes in kept cells; filt, number of cells filtered out; total, total number of cells before filtration, if the input contain all barcodes, this number is the cells left after

de_analysis

de_analysis inputs

Name	Description	Example	Default
perform_de_analysis	If to perform differential expression (DE) analysis	true	true
cluster_labels	Specify the cluster label used for DE analysis	"louvain_labels"	"louvain_labels"
alpha	Control false discovery rate at <alpha>	0.05	0.05
auc	Calculate area under ROC (AUROC)	true	true
fisher	Calculate Fisher's exact test	true	true
t_test	Calculate Welch's t-test.	true	true
mwu	Calculate Mann-Whitney U test	false	false
find_markers_lightgbm	If also to detect markers using LightGBM	false	false
remove_ribo	Remove ribosomal genes with either RPL or RPS as prefixes. Currently only works for human data	false	false
min_gain	Only report genes with a feature importance score (in gain) of at least <gain>	1.0	1.0
annotate_cluster	If also annotate cell types for clusters based on DE results	false	false
annotate_de_test	Differential Expression test to use for inference on cell types. Options: "t", "fisher", or "mwu"	"t"	"t"
organism	Organism, could either be "human_immune", "mouse_immune", "human_brain", "mouse_brain" or a Google bucket link to a JSON file describing the markers	"mouse_brain"	"human_immune"
minimum_report_score	Minimum cell type score to report a potential cell type	0.5	0.5

de_analysis outputs

Name	Type	Description
output_de_h5ad	File	h5ad-formatted results with DE results updated (output_name.h5ad)
output_de_xlsx	File	Spreadsheet reporting DE results (output_name.de.xlsx)
output_markers_xlsx	File	An excel spreadsheet containing detected markers. Each cluster has one tab in the spreadsheet and each tab has three columns, listing markers that are strongly up-regulated, weakly up-regulated and down-regulated (output_name.markers.xlsx)
output_anno_file	File	Annotation file (output_name.anno.txt)

How cell type annotation works

In this subsection, we will describe the format of input JSON cell type marker file, the *ad hoc* cell type inference algorithm, and the format of the output putative cell type file.

JSON file

The top level of the JSON file is an object with two name/value pairs:

- **title**: A string to describe what this JSON file is for (e.g. "Mouse brain cell markers").
- **cell_types**: List of all cell types this JSON file defines. In this list, each cell type is described using a separate object with 2 to 3 name/value pairs:
  - **name**: Cell type name (e.g. "GABAergic neuron").
  - **markers**: List of gene-marker describing objects, each of which has 2 name/value pairs:
    - **genes**: List of positive and negative gene markers (e.g. `["Rbfox3+", "Flt1-"]`).
    - **weight**: A real number between `0.0` and `1.0` to describe how much we trust the markers in **genes**.

    All markers in **genes** share the weight evenly. For instance, if we have 4 markers and the weight is 0.1, each marker has a weight of `0.1 / 4 = 0.025`.

    The weights from all gene-marker describing objects of the same cell type should sum up to 1.0.
  - **subtypes**: Description on cell subtypes for the cell type. It has the same structure as the top level JSON object.

See below for an example JSON snippet:

```
{
  "title" : "Mouse brain cell markers",
    "cell_types" : [
      {
        "name" : "Glutamatergic neuron",
        "markers" : [
          {
            "genes" : ["Rbfox3+", "Reln+", "Slc17a6+", "Slc17a7+"],
            "weight" : 1.0
          }
        ],
        "subtypes" : {
          "title" : "Glutamatergic neuron subtype markers",
            "cell_types" : [
              {
                "name" : "Glutamatergic layer 4",
                "markers" : [
                  {
                    "genes" : ["Rorb+", "Paqr8+"],
                    "weight" : 1.0
                  }
                ]
              }
            ]
        }
      }
    ]
}
```

Inference Algorithm

We have already calculated the up-regulated and down-regulated genes for each cluster in the differential expression analysis step.

First, load gene markers for each cell type from the JSON file specified, and exclude marker genes, along with their associated weights, that are not expressed in the data.

Then scan each cluster to determine its putative cell types. For each cluster and putative cell type, we calculate a score between `0` and `1`, which describes how likely cells from the cluster are of this cell type. The higher the score is, the more likely cells are from the cell type.

To calculate the score, each marker is initialized with a maximum impact value (which is `2`). Then do case analysis as follows:

- For a positive marker:
  - If it is not up-regulated, its impact value is set to `0`.
  - Otherwise, if it is up-regulated:
    - If it additionally has a fold change in percentage of cells expressing this marker (within cluster vs. out of cluster) no less than `1.5`, it has an impact value of `2` and is recorded as a **strong supporting marker**.
    - If its fold change (`fc`) is less than `1.5`, this marker has an impact value of `1 + (fc - 1) / 0.5` and is recorded as a **weak supporting marker**.
- For a negative marker:
  - If it is up-regulated, its impact value is set to `0`.
  - If it is neither up-regulated nor down-regulated, its impact value is set to `1`.
  - Otherwise, if it is down-regulated:
    - If it additionally has `1 / fc` (where `fc` is its fold change) no less than `1.5`, it has an impact value of `2` and is recorded as a **strong supporting marker**.
    - If `1 / fc` is less than `1.5`, it has an impact value of `1 + (1 / fc - 1) / 0.5` and is recorded as a **weak supporting marker**.

The score is calculated as the weighted sum of impact values weighted over the sum of weights multiplied by 2 from all expressed markers. If the score is larger than 0.5 and the cell type has cell subtypes, each cell subtype will also be evaluated.

Output annotation file

For each cluster, putative cell types with scores larger than `minimum_report_score` will be reported in descending order with respect to their scores. The report of each putative cell type contains the following fields:

- **name**: Cell type name.
- **score**: Score of cell type.
- **average marker percentage**: Average percentage of cells expressing marker within the cluster between all positive supporting markers.
- **strong support**: List of strong supporting markers. Each marker is represented by a tuple of its name and percentage of cells expressing it within the cluster.
- **weak support**: List of week supporting markers. It has the same structure as **strong support**.

plot

The h5ad file contains a default cell attribute `Channel`, which records which channel each that single cell comes from. If the input is a CSV format sample sheet, `Channel` attribute matches the `Sample` column in the sample sheet. Otherwise, it's specified in `channel` field of the cluster inputs.

Other cell attributes used in plot must be added via `attributes` field in the `aggregate_matrices` inputs.

plot inputs

Name	Description	Example	Default
plot_composition	Takes the format of "label:attr,label:attr,...,label:attr". If non-empty, generate composition plot for each "label:attr" pair. "label" refers to cluster labels and "attr" refers to sample conditions	"louvain_labels:Donor"	None
plot_fitsne	Takes the format of "attr,attr,...,attr". If non-empty, plot attr colored FIt-SNEs side by side	"louvain_labels,Donor"	None
plot_tsne	Takes the format of "attr,attr,...,attr". If non-empty, plot attr colored t-SNEs side by side	"louvain_labels,Channel"	None
plot_umap	Takes the format of "attr,attr,...,attr". If non-empty, plot attr colored UMAP side by side	"louvain_labels,Donor"	None
plot_fle	Takes the format of "attr,attr,...,attr". If non-empty, plot attr colored FLE (force-directed layout embedding) side by side	"louvain_labels,Donor"	None
plot_diffmap	Takes the format of "attr,attr,...,attr". If non-empty, generate attr colored 3D interactive plot. The 3 coordinates are the first 3 PCs of all diffusion components	"louvain_labels,Donor"	None
plot_citeseq_fitsne	plot cells based on FIt-SNE coordinates estimated from antibody expressions. Takes the format of "attr,attr,...,attr". If non-empty, plot attr colored FIt-SNEs side by side	"louvain_labels,Donor"	None
plot_net_tsne	Takes the format of "attr,attr,...,attr". If non-empty, plot attr colored t-SNEs side by side based on net t-SNE result.	"leiden_labels,Channel"	None
plot_net_umap	Takes the format of "attr,attr,...,attr". If non-empty, plot attr colored UMAP side by side based on net UMAP result.	"leiden_labels,Donor"	None
plot_net_fle	Takes the format of "attr,attr,...,attr".	"leiden_labels,Donor"	None

plot outputs

Name	Type	Description
output_pdfs	Array[File]	Outputted pdf files
output_htmls	Array[File]	Outputted html files

Generate SCP Output

Generate analysis result in Single Cell Portal (SCP) compatible format.

scp_output inputs

Name	Description	Example	Default
generate_scp_output	Whether to generate SCP format output or not.	false	false
output_dense	Output dense expression matrix, instead of the default sparse matrix format.	false	false

scp_output outputs

Name	Type	Description
output_scp_files	Array[File]	Outputted SCP format files.

9.8.2 Run CITE-Seq analysis

To run CITE-Seq analysis, turn on `cite_seq` option in cluster inputs of cumulus workflow.

An embedding of epitope expressions via FIt-SNE is available at basis `X_citeseq_fitsne`.

To plot this epitope embedding, specify attributes to plot in `plot_citeseq_fitsne` field of cluster inputs.

9.8.3 Run subcluster analysis

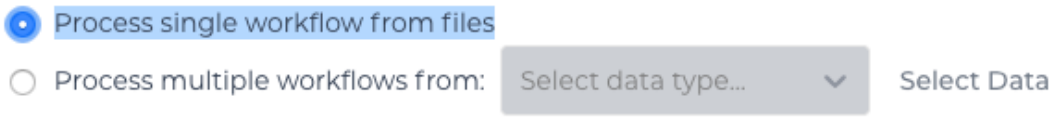
Once we have **cumulus** outputs, we could further analyze a subset of cells by running **cumulus_subcluster**. To run **cumulus_subcluster**, follow the following steps:

1. Import **cumulus_subcluster** method.

   See the Terra documentation for adding a workflow. The cumulus workflow is under `Broad Methods Repository` with name "**cumulus/cumulus_subcluster**".

   Moreover, in the workflow page, click the `Export to Workspace...` button, and select the workspace to which you want to export cumulus workflow in the drop-down menu.

2. In your workspace, open `cumulus_subcluster` in `WORKFLOWS` tab. Select `Process single workflow from files` as below



   and click the `SAVE` button.

cumulus_subcluster steps:

cumulus_subcluster processes the subset of single cells in the following steps:

1. **subcluster**. In this step, **cumulus_subcluster** first select the subset of cells from **cumulus** outputs according to user-provided criteria. It then performs batch correction, dimension reduction, diffusion map calculation, graph-based clustering and 2D visualization calculation (e.g. t-SNE/UMAP/FLE).
2. **de_analysis** (optional). In this step, **cumulus_subcluster** calculates potential markers for each cluster by performing a variety of differential expression (DE) analysis. The available DE tests include Welch's t test, Fisher's exact test, and Mann-Whitney U test. **cumulus_subcluster** can also calculate the area under ROC curve (AUROC) values for putative markers. If the samples are human or mouse immune cells, **cumulus_subcluster** can optionally annotate putative cell types for each cluster based on known markers.
3. **plot** (optional). In this step, **cumulus_subcluster** can generate the following 5 types of figures based on the **subcluster** step results:
   - **composition** plots which are bar plots showing the cell compositions (from different conditions) for each cluster. This type of plots is useful to fast assess library quality and batch effects.
   - **tsne**, **fitsne**, and **net_tsne**: t-SNE like plots based on different algorithms, respectively. Users can specify different cell attributes (e.g. cluster labels, conditions) for coloring side-by-side.
   - **umap** and **net_umap**: UMAP like plots based on different algorithms, respectively. Users can specify different cell attributes (e.g. cluster labels, conditions) for coloring side-by-side.
   - **fle** and **net_fle**: FLE (Force-directed Layout Embedding) like plots based on different algorithms, respectively. Users can specify different cell attributes (e.g. cluster labels, conditions) for coloring side-by-side.
   - **diffmap** plots which are 3D interactive plots showing the diffusion maps. The 3 coordinates are the first 3 PCs of all diffusion components.

cumulus_subcluster's inputs

cumulus_subcluster shares many inputs/outputs with **cumulus**, we will only cover inputs/outputs that are specific to **cumulus_subcluster** in this section.

Note that we will make the required inputs/outputs bold and all other inputs/outputs are optional.

Name	Description	Example	Default
input_h5ad	Google bucket URL of input h5ad file containing *cumulus* results	"gs://fc-e0000000-0000-0000-0000-000000000000/my_results_dir/my_results.h5ad"	
output_name	This is the prefix for all output files. It should contain the Google bucket URL, subdirectory name and output name prefix	"gs://fc-e0000000-0000-0000-0000-000000000000/my_results_dir/my_results_sub"	
subset_selections	Specify which cells will be included in the subcluster analysis. This field contains one or more <subset_selection> strings separated by ';'. Each <subset_selection> string takes the format of 'attr:value,...,value', which means select cells with attr in the values. If multiple <subset_selection> strings are specified, the subset of cells selected is the intersection of these strings	"louvain_labels:3,6" or "louvain_labels:3,6;Donor:1,2"	
calculate_pseudotime	Calculate diffusion-based pseudotimes based on <roots>. <roots> should be a comma-separated list of cell barcodes	"sample_1-ACCCGGGTTT-1,sample_1-TCCCGGGAAA-2"	None
num_cpu	Number of cpus per cumulus job	32	64
memory	Memory size string	"200G"	"200G"
disk_space	Total disk space in GB	100	100
preemptible	Number of preemptible tries	2	2

For other **cumulus_subcluster** inputs, please refer to cumulus cluster inputs list for details. Notice that some inputs (as listed below) in **cumulus cluster** inputs list are DISABLED for **cumulus_subcluster**:

- cite_seq
- cite_seq_capping
- output_filtration_results
- plot_filtration_results
- plot_filtration_figsize
- output_seurat_compatible
- batch_group_by
- min_genes
- max_genes
- min_umis
- max_umis
- mito_prefix
- percent_mito
- gene_percent_cells

- min_genes_on_raw
- counts_per_cell_after

cumulus_subcluster's outputs

Name	Type	Description
output_h5ad	File	h5ad-formatted HDF5 file containing all results (output_name.h5ad). If `perform_de_analysis` is on, this file should be the same as *output_de_h5ad*. To load this file in Python, it's similar as in cumulus cluster outputs section. Besides, for subcluster results, there is a new cell attributes in `data.obs['pseudo_time']`, which records the inferred pseudotime for each cell.
output_loom_file	File	Generated loom file (output_name.loom)
output_parquet_file	File	Generated PARQUET file that contains metadata and expression levels for every gene (output_name.parquet)
output_de_h5ad	File	Generated h5ad-formatted results with DE results updated (output_name.h5ad)
output_de_xlsx	File	Generated Spreadsheet reporting DE results (output_name.de.xlsx)
output_pdfs	Array[File]	Generated pdf files
output_htmls	Array[File]	Generated html files

9.8.4 Load Cumulus results into Seurat

First, you need to set `output_seurat_compatible` field to `true` in cumulus cluster inputs to generate a Seurat-compatible output file `output_name.seurat.h5ad`, in addition to the normal result `output_name.h5ad`.

Notice that python, the anndata python library with version at least `0.6.22.post1`, and the `reticulate` R library are required to load the result into Seurat.

Execute the R code below to load the results into Seurat (working with both Seurat v2 and v3):

```
source("https://raw.githubusercontent.com/klarman-cell-observatory/cumulus/master/
↪workflows/cumulus/h5ad2seurat.R")
ad <- import("anndata", convert = FALSE)
test_ad <- ad$read_h5ad("output_name.seurat.h5ad")
result <- convert_h5ad_to_seurat(test_ad)
```

The resulting Seurat object `result` has three data slots:

- **raw.data** records filtered raw count matrix.
- **data** records filtered and log-normalized expression matrix.
- **scale.data** records variable-gene-selected, standardized expression matrix that are ready to perform PCA.

9.8.5 Visualize Cumulus results in Python

Ensure you have Pegasus installed.

Download your analysis result data, say `output_name.h5ad`, from Google bucket to your local machine.

Load the output:

```
import pegasus as pg
adata = pg.read_input("output_name.h5ad")
```

Violin plot of the computed quality measures:

```
fig = pg.violin(adata, keys = ['n_genes', 'n_counts', 'percent_mito'], by = 'passed_qc
↪')
fig.savefig('output_file.qc.pdf', dpi = 500)
```

t-SNE plot colored by louvain cluster labels and channel:

```
fig = pg.embedding(adata, basis = 'tsne', keys = ['louvain_labels', 'Channel'])
fig.savefig('output_file.tsne.pdf', dpi = 500)
```

t-SNE plot colored by genes of interest:

```
fig = pg.embedding(adata, basis = 'tsne', keys = ['CD4', 'CD8A'])
fig.savefig('output_file.genes.tsne.pdf', dpi = 500)
```

For other embedding plots using FIt-SNE (`fitsne`), Net t-SNE (`net_tsne`), CITE-Seq FIt-SNE (`citeseq_fitsne`), UMAP (`umap`), Net UMAP (`net_umap`), FLE (`fle`), or Net FLE (`net_fle`) coordinates, simply substitute its basis name for `tsne` in the code above.

Composition plot on louvain cluster labels colored by channel:

```
fig = pg.composition_plot(adata, by = 'louvain_labels', condition = 'Channel')
fig.savefig('output_file.composition.pdf', dpi = 500)
```

9.9 Demuxlet

This workflow runs demuxlet to deconvolute sample identity when multiple samples are pooled by barcoded single-cell sequencing.

1. Align your single-cell sequencing data (for example using the cellranger or drop_seq workflows).
2. Create a sample sheet.

   Please note that the columns in the tab separated file must be in the order shown below and does not contain a header line.

Column	Description
Name	Sample name.
BAM	Location of the BAM file in the cloud (gs:// URL).
Barcodes	Location of the valid cellular barcodes file in the cloud (gs:// URL).
VCF	Location of the VCF file to use for this sample in the cloud (gs:// URL).

   Example:

```
sample-1,gs://fc-e0000000/sample-1/out/possorted_genome_bam.bam,gs://fc-
↪e0000000/sample-1/out/filtered_feature_bc_matrix/barcodes.tsv.gz,gs://
↪fc-e0000000/sample-1.vcf
sample-2,gs://fc-e0000000/sample-2/out/possorted_genome_bam.bam,gs://fc-
↪e0000000/sample-2/out/filtered_feature_bc_matrix/barcodes.tsv.gz,gs://
↪fc-e0000000/sample-2.vcf
```

3. Upload your sample sheet to the workspace bucket.

   Example:

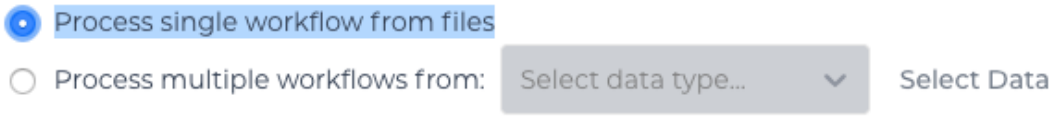
   ```
   gsutil cp /foo/bar/projects/sample_sheet.tsv gs://fc-e0000000/
   ```

4. Import *demuxlet* workflow to your workspace.

   See the Terra documentation for adding a workflow. The workflow is under `Broad Methods Repository` with the name "**cumulus/demuxlet**".

   Next, in the workflow page, click the `Export to Workspace...` button, and select the workspace you want to export to in the drop-down menu.

5. In your workspace, open `demuxlet` in `WORKFLOWS` tab. Select `Process single workflow from files` as below

   and click the `Save` button.

9.9.1 Inputs

Please see the description of important inputs below.

Column	Description
tsv_file	Four column tab-separated file without a header with name, coordinate sorted bam, barcodes, and vcf
min_MQ	Minimum mapping quality to consider (default 20)
alpha	Grid of alpha to search for (default [0.1, 0.2, 0.3, 0.4, 0.5]).
min_TD	Minimum distance to the tail (default 0)
tag_group	Tag representing readgroup or cell barcodes, in the case to partition the BAM file into multiple groups (default "CB")
tag_UMI	Tag representing UMIs (default "UB"")
field	FORMAT field to extract the genotype, likelihood, or posterior from (default "GT")
geno_error	Offset of genotype error rate (default 0.1)

9.9.2 Outputs

The demuxlet output file contains the best guess of the sample identity, with detailed statistics to reach to the best guess.

9.10 Run Terra pipelines via command line

You can run Terra pipelines via the command line by installing the **altocumulus** package.

9.10.1 Install `altocumulus` for Broad users

Request an UGER node:

```
reuse UGER
qrsh -q interactive -l h_vmem=4g -pe smp 8 -binding linear:8 -P regevlab
```

The above command requests an interactive shell using the regevlab project with 4G memory per thread, 8 threads. Feel free to change the memory, thread, and project parameters.

Add conda to your path:

```
reuse Anaconda3
```

Activate the alto virtual environment:

```
source activate /seq/regev_genome_portal/conda_env/cumulus
```

9.10.2 Install `altocumulus` for non-Broad users

1. Make sure you have `conda` installed. If you haven't installed conda, use the following commands to install it on Linux:

   ```
   wget https://repo.continuum.io/miniconda/Miniconda3-latest-Linux-x86_64.sh .
   bash Miniconda3-latest-Linux-x86_64.sh -p /home/foo/miniconda3
   mv Miniconda3-latest-Linux-x86_64.sh /home/foo/miniconda3
   ```

   where `/home/foo/miniconda3` should be replaced by your own folder holding Miniconda3.

Or use the following commdands for MacOS installation:

```
curl -O curl -O https://repo.anaconda.com/miniconda/Miniconda3-latest-MacOSX-x86_64.sh
bash Miniconda3-latest-MacOSX-x86_64.sh -p /Users/foo/miniconda3
mv Miniconda3-latest-MacOSX-x86_64.sh /Users/foo/miniconda3

where ``/Users/foo/miniconda3`` should be replaced by your own folder holding␣
↪Miniconda3.
```

1. Create a conda environment named "alto" and install `altocumulus`:

   ```
   conda create -n alto -y pip
   source activate alto
   git clone https://github.com/klarman-cell-observatory/altocumulus.git
   cd altocumulus
   pip install -e .
   ```

When the installation is done, type `alto fc_run -h` in terminal to see if you can see the help information.

9.10.3 Run Terra workflows via `alto fc_run`

alto fc_run runs a Terra method. Features:

- Uploads local files/directories in your inputs to a Google Cloud bucket updates the file paths to point to the Google Cloud bucket.

  Your sample sheet can point to local file paths. In this case, `alto run` will take care of uploading directories (e.g. fastq directories) and modifying the sample sheet to point to a Google Cloud bucket.

- Creates or uses an existing workspace.
- Uses the latest version of a method unless the method version is specified.

Options

Required options are in bold.

Name	Description
-m <METHOD> –method <METHOD>	Specify a Terra workflow <*METHOD*> to use. <*METHOD*> is of format *Namespace/Name* (e.g. `cumulus/cellranger_workflow`). A snapshot version number can optionally be specified (e.g. `cumulus/cellranger_workflow/4`); otherwise the latest snapshot of the method is used.
-w <WORKSPACE> –workspace <WORKSPACE>	Specify which Terra workspace <*WORKSPACE*> to use. <*WORKSPACE*> is also of format *Namespace/Name* (e.g. `foo/bar`). The workspace will be created if it does not exist.
-i <WDL_INPUTS> –inputs <WDL_INPUTS>	Specify the WDL input JSON file to use. It can be a local file, a JSON string, or a Google bucket URL directing to a remote JSON file.
–bucket-folder <folder>	Store inputs to <folder> under workspace's google bucket.
-o <updated_json> –upload <updated_json>	Upload files/directories to Google bucket of the workspace, and generate an updated input JSON file (with local paths replaced by Google bucket URLs) to <updated_json> on local machine.

Example

This example shows how to use `alto fc_run` to run cellranger_workflow to extract gene-count matrices from sequencing output.

1. Prepare your sample sheet `example_sample_sheet.csv` as the following:

```
Sample,Reference,Flowcell,Lane,Index,Chemistry
sample_1,GRCh38,/my-local-path/flowcell1,1-2,SI-GA-A8,threeprime
sample_2,GRCh38,/my-local-path/flowcell1,3-4,SI-GA-B8,threeprime
sample_3,mm10,/my-local-path/flowcell1,5-6,SI-GA-C8,fiveprime
sample_4,mm10,/my-local-path/flowcell1,7-8,SI-GA-D8,fiveprime
sample_1,GRCh38,/my-local-path/flowcell2,1-2,SI-GA-A8,threeprime
sample_2,GRCh38,/my-local-path/flowcell2,3-4,SI-GA-B8,threeprime
sample_3,mm10,/my-local-path/flowcell2,5-6,SI-GA-C8,fiveprime
sample_4,mm10,/my-local-path/flowcell2,7-8,SI-GA-D8,fiveprime
```

   where `/my-local-path` is the top-level directory of your BCL files on your local machine.

   Note that `sample_1`, `sample_2`, `sample_3`, and `sample_4` are sequenced on 2 flowcells.

2. Prepare your JSON input file `inputs.json` for cellranger_workflow:

```
{
        "cellranger_workflow.input_csv_file" : "/my-local-path/sample_sheet.csv",
        "cellranger_workflow.output_directory" : "gs://url/outputs",
        "cellranger_workflow.delete_input_bcl_directory": true
}
```

   where `gs://url/outputs` is the folder on Google bucket of your workspace to hold output.

3. Run the following command to kick off your Terra workflow:

```
alto fc_run -m cumulus/cellranger_workflow -i inputs.json -w myworkspace_
↪namespace/myworkspace_name -o inputs_updated.json
```

   where `myworkspace_namespace/myworkspace_name` should be replaced by your workspace namespace and name.

Upon success, `alto fc_run` returns a URL pointing to the submitted Terra job for you to monitor.

If for any reason, your job failed. You could rerun it without uploading files again via the following command:

```
alto fc_run -m cumulus/cellranger_workflow -i inputs_updated.json -w myworkspace_
↪namespace/myworkspace_name
```

because `inputs_updated.json` is the updated version of `inputs.json` with all local paths being replaced by their corresponding Google bucket URLs after uploading.

9.11 Examples

9.11.1 Example of Cell-Hashing and CITE-Seq Analysis on Cloud

In this example, you'll learn how to perform Cell-Hashing and CITE-Seq analysis using **cumulus** on Terra.

0. Workspace and Data Preparation

After registering on Terra and creating a workspace there, you'll need the following two information:

- **Terra workspace name**. This is shown on your Terra workspace webpage, with format *"<workspace-namespace>/<workspace-name>"*. Let it be `ws-lab/ws-01` in this example, which means that your workspace has namespace **ws-lab** and name **ws-01**.
- The corresponding **Google Cloud Bucket location** of your workspace. You can check it by clicking the link under **"Google Bucket"** title on your Terra workspace webpage. Let it be `gs://fc-e0000000-0000-0000-0000-000000000000` in this example.

Then upload your BCL directories to Google bucket of your workspace using gsutil:

```
gsutil -m cp -r /my-local-path/BCL/* gs://fc-e0000000-0000-0000-0000-000000000000/
↪data-source
```

where `/my-local-path/BCL` is the path to the top-level directory of your BCL files on your local machine, and `data-source` is the folder on Google bucket to hold the uploaded data.

---

1. Extract Gene-Count Matrices

First step is to extract gene-count matrices from sequencing output.

You need two original files from your dataset to start:

- Cell-Hashing Index CSV file, say its filename is `cell_hashing_index.csv`, of format *feature_barcode,feature_name*. See an example below:

  ```
  AATCATCACAAGAAA,CB1
  GGTCACTGTTACGTA,CB2
  ... ...
  ```

  where each line is a pair of feature barcode and feature name of a sample.

- CITE-Seq Index CSV file, say its filename is `cite_seq_index.csv`, of the same format as above. See an example below:

  ```
  TTACATGCATTACGA,CD19
  GCATTAGCATGCAGC,HLA-ABC
  ... ...
  ```

  where each line is a pair of Barcode and Specificity of an Antibody.

Then upload them to your Google Bucket using gsutil. Assuming both files are in folder `/Users/foo/data-source` on your local machine, type the following command to upload:

```
gsutil -m cp -r /Users/foo/data-source gs://fc-e0000000-0000-0000-0000-000000000000/
↪data-source
```

where option `-m` means copy in parallel, `-r` means copy the directory recursively, and `gs://fc-e0000000-0000-0000-0000-000000000000/data-source` is your working directory at cloud side, which can be changed at your will.

Next, create a sample sheet, `cellranger_sample_sheet.csv`, on your local machine with content below:

```
Sample,Reference,Flowcell,Lane,Index,DataType,FeatureBarcodeFile
sample_control,GRCh38,gs://fc-e0000000-0000-0000-0000-000000000000/data-source,2,SI-
↪GA-F1,rna
sample_cc,GRCh38,gs://fc-e0000000-0000-0000-0000-000000000000/data-source,3,SI-GA-A1,
↪rna
sample_cell_hashing,GRCh38,gs://fc-e0000000-0000-0000-0000-000000000000/data-source,3,
↪ATTACTCG,adt,cell_hashing_index.csv
sample_cite_seq,GRCh38,gs://fc-e0000000-0000-0000-0000-000000000000/data-source,3,
↪CGTGAT,adt,cite_seq_index.csv
```

For the details on how to prepare this sample sheet, please refer to Step 3 of Cell Ranger sample sheet instruction.

When you are done with the sample sheet, upload it to Google bucket:

```
gsutil cp cellranger_sample_sheet.csv gs://fc-e0000000-0000-0000-0000-000000000000/my-
↪dir/
```

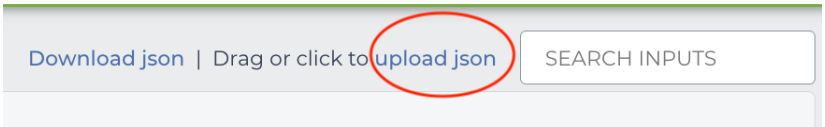
Now we are ready to set up **cellranger_workflow** workflow for this phase. If your workspace doesn't have this workflow, import it to your workspace by following Step 5 and 6 of cellranger_workflow documentation.

Then prepare a JSON file, `cellranger_inputs.json`, which is used to set up the workflow inputs:

```
{
        "cellranger_workflow.input_csv_file" : "gs://fc-e0000000-0000-0000-0000-
↪000000000000/my-dir/cellranger_sample_sheet.csv",
        "cellranger_workflow.output_directory" : "gs://fc-e0000000-0000-0000-0000-
↪000000000000/my-dir"
}
```

where `gs://fc-e0000000-0000-0000-0000-000000000000/my-dir` is the remote directory in which the output of cellranger_workflow will be generated. For the details on the options above, please refer to Cell Ranger workflow inputs.

When you are done with the JSON file, on cellranger_workflow workflow page, upload `cellranger_inputs.json` by clicking `upload json` link as below:



Then Click `SAVE` button to save the inputs, and click `RUN ANALYSIS` button as below to start the job:

When the execution is done, all the output results will be in folder `gs://fc-e0000000-0000-0000-0000-000000000000/my-dir`.

You'll need 4 files for the next phases. 3 are from the output:

- RNA count matrix of the sample group of interest: `gs://fc-e0000000-0000-0000-0000-000000000000/my-dir/sample_cc/raw_feature_bc_matrix.h5`;
- Cell-Hashing Antibody count matrix: `gs://fc-e0000000-0000-0000-0000-000000000000/my-dir/sample_cell_hashing/sample_cell_hashing.csv`;

- CITE-Seq Antibody count matrix: `gs://fc-e0000000-0000-0000-0000-000000000000/my-dir/sample_cite_seq/sample_cite_seq.csv`.

Besides, create a sample sheet, `citeseq_antibody_control.csv`, with content as the following example:

```
Antibody,Control
CD3-0034,Mouse_IgG1
CD4-0045,Mouse_IgG1
... ...
```

where each line is a pair of Antibody name and the Control group name to which it is assigned. You should be able to get this information from your experiment setting or the original dataset.

Copy or upload them to `gs://fc-e0000000-0000-0000-0000-000000000000/my-dir`.

---

2. Demultiplex Cell-Hashing Data

1. Prepare a sample sheet, `cell_hashing_sample_sheet.csv`, with the following content:

   ```
   OUTNAME,RNA,ADT,TYPE
   exp,gs://fc-e0000000-0000-0000-0000-000000000000/my-dir/raw_feature_bc_matrix.h5,
   ↪gs://fc-e0000000-0000-0000-0000-000000000000/my-dir/sample_cell_hashing.csv,
   ↪cell-hashing
   ```

   where **OUTNAME** specifies the subfolder and file names of output, which is free to change, **RNA** and **ADT** columns specify the RNA and ADT meta-data of samples, and **TYPE** is `cell-hashing` for this phase.

   Then upload it to Google bucket:

   ```
   gsutil cp cell_hashing_sample_sheet.csv gs://fc-e0000000-0000-0000-0000-
   ↪000000000000/my-dir/
   ```

2. If your workspace doesn't have **cumulus_hashing_cite_seq** workflow, import it to your workspace by following Step 5 and 6 of cumulus_hashing_cite_seq documentation.

3. Prepare an input JSON file, `cell_hashing_inputs.json` with the following content to set up cumulus_hashing_cite_seq workflow inputs:

   ```
   {
           "cumulus_hashing_cite_seq.input_sample_sheet" : "gs://fc-e0000000-0000-
   ↪0000-0000-000000000000/my-dir/cell_hashing_sample_sheet.csv",
           "cumulus_hashing_cite_seq.output_directory" : "gs://fc-e0000000-0000-0000-
   ↪0000-000000000000/my-dir/",
           "cumulus_hashing_cite_seq.demuxEM_min_num_genes" : 500,
           "cumulus_hashing_cite_seq.demuxEM_generate_diagnostic_plots" : true
   }
   ```

   For the details on these options, please refer to cell-hashing/nuclei-hashing inputs.

4. On the page of cumulus_hashing_cite_seq workflow, upload `cell_hashing_inputs.json` by clicking `upload json` link. Save the inputs, and click `RUN ANALYSIS` button to start the job.

When the execution is done, you'll get a processed file, `exp_demux.h5sc`, stored on cloud `gs://fc-e0000000-0000-0000-0000-000000000000/my-dir/exp/`.

---

3. Merge RNA and ADT Matrices for CITE-Seq Data

1. Prepare a sample sheet, `cite_seq_sample_sheet.csv`, with the following content:

```
OUTNAME,RNA,ADT,TYPE
exp_raw,gs://fc-e0000000-0000-0000-0000-000000000000/my-dir/exp/exp_demux.h5sc,
↪gs://fc-e0000000-0000-0000-0000-000000000000/my-dir/sample_cite_seq.csv,cite-seq
```

   The structure of sample sheet here is the same as Phase 2. The difference is that you are now using the demultiplexed output `h5sc` file from Phase 2 as **RNA** here, and the sample **TYPE** is now `cite-seq`.

   Then upload it to Google bucket:

```
gsutil cp cite_seq_sample_sheet.csv gs://fc-e0000000-0000-0000-0000-000000000000/
↪my-dir/
```

2. Prepare an input JSON file, `cite_seq_inputs.json`, in the same directory as above, with the following content:

```
{
        "cumulus_hashing_cite_seq.input_sample_sheet" : "gs://fc-e0000000-0000-
↪0000-0000-000000000000/my-dir/cite_seq_sample_sheet.csv",
        "cumulus_hashing_cite_seq.output_directory" : "gs://fc-e0000000-0000-0000-
↪0000-000000000000/my-dir/",
        "cumulus_hashing_cite_seq.antibody_control_csv" : "gs://fc-e0000000-0000-
↪0000-0000-000000000000/my-dir/citeseq_antibody_control.csv"
}
```

   For the details on these options, please refer to cell-hashing/nuclei-hashing inputs.

3. On **cumulus_hashing_cite_seq** workflow page, clear all previous inputs, and then upload `cite_seq_inputs.json` by clicking `upload json` link. Save the new inputs, and click `RUN ANALYSIS` button to start the job.

When the execution is done, you'll get a merged raw matrices file, `exp_raw.h5sc`, stored on cloud `gs://fc-e0000000-0000-0000-0000-000000000000/my-dir/exp_raw`.

---

4. Data Analysis

1. Prepare a sample sheet, `cumulus_count_matrix.csv`, with the following content:

```
Sample,Location
exp,gs://fc-e0000000-0000-0000-0000-000000000000/my-dir/exp_raw/exp_raw.h5sc
```

   This sample sheet describes the metadata for each 10x channel (as one row in the sheet). **Sample** specifies the name for each channel, which can be renamed; **Location** specifies the file location, which is the output of Phase 3.

   Then upload it to Google bucket:

```
gsutil cp cumulus_count_matrix.csv gs://fc-e0000000-0000-0000-0000-000000000000/
↪my-dir/
```

   Alternative, if you have only one count matrix for analysis, which is the case here, you can skip this step. See this manual for input file formats that cumulus currently supports.

2. If your workspace doesn't have **cumulus** workflow, import it to your workspace by following Step 2 and 3 of cumulus documentation.

3. Prepare a JSON file, `cumulus_inputs.json` with the following content to set up **cumulus** workflow inputs:

```
{
        "cumulus.input_file" : "gs://fc-e0000000-0000-0000-0000-000000000000/my-
↪dir/cumulus_count_matrix.csv",
        "cumulus.output_name" : "gs://fc-e0000000-0000-0000-0000-000000000000/my-
↪dir/results/exp_merged_out",
        "cumulus.num_cpu" : 8,
        "cumulus.select_only_singlets" : true,
        "cumulus.cite_seq" : true,
        "cumulus.run_louvain" : true,
        "cumulus.find_markers_lightgbm" : true,
        "cumulus.remove_ribo" : true,
        "cumulus.mwu" : true,
        "cumulus.annotate_cluster" : true,
        "cumulus.plot_fitsne" : "louvain_labels,assignment",
        "cumulus.plot_citeseq_fitsne" : "louvain_labels,assignment",
        "cumulus.plot_composition" : "louvain_labels:assignment"
}
```

   Alternatively, if you have only one count matrix for analysis and has skipped Step 1, directly set its location in `cumulus.input_file` parameter above. For this example, it is:

```
{
        "cumulus.input_file" : "gs://fc-e0000000-0000-0000-0000-000000000000/my-
↪dir/exp_raw/exp_raw.h5sc",
        ... ...
}
```

   All the rest parameters remain the same.

   Notice that for some file formats, `cumulus.genome` is required.

   A typical cumulus pipeline consists of 4 steps, which is given here. For the details of options above, please refer to cumulus inputs.

4. On the page of cumulus workflow, upload `cumulus_inputs.json` by clicking `upload json` link. Save the inputs, and click `RUN ANALYSIS` button to start the job.

When the execution is done, you'll get the following results stored on cloud `gs://fc-e0000000-0000-0000-0000-000000000000/my-dir/results/` to check:

- `exp_merged_out.h5sc`: The aggregated count matrix data. This file doesn't exist if your `cumulus.input_file` parameter is not a sample sheet.
- `exp_merged_out.h5ad`: The processed RNA matrix data.
- `exp_merged_out.filt.xlsx`: The Quality-Control (QC) summary of the raw data.
- `exp_merged_out.filt.{UMI, gene, mito}.pdf`: The QC plots of the raw data.
- `exp_merged_out.de.xlsx`: Differential Expression analysis result.
- `exp_merged_out.markers.xlsx`: Result on cluster-specific markers predicted by gradient boosting machine.
- `exp_merged_out.anno.txt`: Cell type annotation output.
- `exp_merged_out.fitsne.pdf`: FIt-SNE plot.

- `exp_merged_out.citeseq.fitsne.pdf`: CITE-Seq FIt-SNE plot.
- `exp_merged_out.louvain_labels.assignment.composition.pdf`: Composition plot.

You can directly go to your Google Bucket to view or download these results.

(optional) Run Terra Workflows in Command Line

For Phase 1, 2, and 3, besides uploading sample sheets and setting-up workflow inputs on workflow pages, you can also start the workflow execution via command line using **altocumulus** tool.

First, install *altocumulus* by following altocumulus installation instruction.

1. For Phase 1 above, when you are done with creating a sample sheet `cellranger_sample_sheet.csv` on your local machine, in the same directory, prepare JSON file `cellranger_inputs.json` as below:

```
{
        "cellranger_workflow.input_csv_file" : "cellranger_sample_sheet.csv",
        ... ...
}
```

   where all the rest parameters remain the same as in Phase 1. Import **cellranger_workflow** workflow to your workspace as usual.

   Now run the following command in the same directory on your local machine:

```
alto fc_run -m cumulus/cellranger_workflow -w ws-lab/ws-01 --bucket-folder my-dir␣
↪-i cellranger_input.json -o cellranger_input_updated.json
```

   Notice that if the execution failed, you could rerun the execution by setting `cellranger_input_updated.json` for `-i` option to use the sample sheet already uploaded to Google bucket. Similarly below.

2. For Phase 2 above, similarly, in the same directory of your `cell_hashing_sample_sheet.csv` file, prepare JSON file `cell_hashing_inputs.json` as below:

```
{
        "cumulus_hashing_cite_seq.input_sample_sheet" : "cell_hashing_sample_
↪sheet.csv",
        ... ...
}
```

   where all the rest parameters remain the same as in Phase 2. Import **cumulus_hashing_cite_seq** workflow to your workspace as usual.

   Run the following command in the same directory on your local machine:

```
alto fc_run -m cumulus/cumulus_hashing_cite_seq -w ws-lab/ws-01 --bucket-folder␣
↪my-dir -i cell_hashing_inputs.json -o cell_hashing_inputs_updated.json
```

3. For Phase 3 above, similarly, in the same directory of your `cite_seq_sample_sheet.csv` file, prepare JSON file `cite_seq_inputs.json` as below:

```
{
        "cumulus_hashing_cite_seq.input_sample_sheet" : "cite_seq_sample_sheet.csv
↪",
        ... ...
}
```

where all the rest parameters remain the same as in Phase 3.

Run the following command in the same directory on your local machine:

```
alto fc_run -m cumulus/cumulus_hashing_cite_seq -w ws-lab/ws-01 --bucket-folder␣
↪my-dir -i cite_seq_inputs.json -o cite_seq_inputs_updated.json
```

4. For Phase 4 above, similarly, in the same directory of your `cumulus_count_matrix.csv` file, prepare JSON file `cumulus_inputs.json` as below:

```
{
        "cumulus.input_file" : "cumulus_count_matrix.csv",
        ... ...
}
```

where all the rest parameters remain the same as in Phase 4.

Alternatively, if your input is not a sample sheet, simply set your `cumulus_inputs.json` as:

```
{
        "cumulus.input_file" : "gs://fc-e0000000-0000-0000-0000-000000000000/my-
↪dir/exp_raw/exp_raw.h5sc",
        ... ...
}
```

where all the rest parameters remain the same.

Run the following command in the same directory of your `cumulus_inputs.json` file:

```
alto fc_run -m cumulus/cumulus -w ws-lab/ws-01 --bucket-folder my-dir/results -i␣
↪cumulus_inputs.json -o cumulus_inputs_updated.json
```

Examples using Terra to perform single-cell sequencing analysis are provided here. Please click the topics on the left panel under title **"Examples"** to explore.

9.12 Contribution

9.13 Contact us

If you have any questions related to Cumulus, please feel free to contact us via Cumulus Support Google Group.