

Package ‘SFEData’

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Title Example SpatialFeatureExperiment datasets

Version 1.4.0

Description Example spatial transcriptomics datasets with Simple Feature annotations as SpatialFeatureExperiment objects. Technologies include Visium, slide-seq, Nanostring CoxMX, Vizgen MERFISH, and 10X Xenium. Tissues include mouse skeletal muscle, human melanoma metastasis, human lung, breast cancer, and mouse liver.

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BiermannMelaMetasData *Melanoma metastasis slide-seq2 data*

Description

This function can download one of the human melanoma brain metastasis (MBM) samples and one of the melanoma extracranial metastasis (ECM) samples from the paper Dissecting the treatment-naive ecosystem of human melanoma brain metastasis, [Biermann et al.](#) The datasets are GSM6025935_MBM05_rep1 and GSM6025946_ECM01_rep1. The raw counts and cell metadata were downloaded from GEO. The raw counts, QC metrics such as number of UMIs and genes detected per barcode, and centroid coordinates as sf POINT geometry, are included in the SFE object.

Usage

```
BiermannMelaMetasData(dataset = datasets, force = FALSE, verbose = TRUE)
```

Arguments

| | |
|---------|---|
| dataset | Which dataset to use, must be one of "MBM05_rep1" and "ECM01_rep1". |
| force | Logical, whether to force redownload if the files are already present. Defaults to FALSE. |
| verbose | Whether to display progress of download. |

Value

A SpatialFeatureExperiment object.

Examples

```
sfe <- BiermannMelaMetasData()
```

HeNSCLCData

Nanostring FFPE CosMX human NSCLC data

Description

One of the CosMX example datasets for human non small cell lung cancer (NSCLC, Lung5_Rep1) from the [Nanostring website](#) was downloaded and formatted into an SFE object. The dataset is described in the paper High-plex Multiomic Analysis in FFPE at Subcellular Level by Spatial Molecular Imaging, [He et al.](#) Since there's no easy way to get the cell segmentation polygon coordinates from the Nanostring website, the polygon coordinates were downloaded from Seurat's vignette. The raw count matrix, QC metrics, cell segmentation in one z-plane, and other cell attributes such as area, aspect ratio, mean DAPI level, mean immunofluorescence signal, and etc. are included in the SFE object.

Usage

```
HeNSCLCData(dataset = datasets, force = FALSE, verbose = TRUE)
```

Arguments

| | |
|---------|---|
| dataset | Which dataset to use, for now can only be "Lung5_Rep1". |
| force | Logical, whether to force redownload if the files are already present. Defaults to FALSE. |
| verbose | Whether to display progress of download. |

Value

A SpatialFeatureExperiment object.

JanesickBreastData

Xenium FFPE human breast cancer data

Description

This dataset was downloaded from the [10X website](#), and described in the paper High resolution mapping of the breast cancer tumor microenvironment using integrated single cell, spatial and in situ analysis of FFPE tissue, [Janesick et al.](#) The dataset might not be representative of later Xenium data. There are two samples, which can both be downloaded with this package. For each sample, the raw gene counts, QC metrics, cell and nuclei segmentation polygons in one z-plane, and cell centroids are included in the SFE object. The two samples are in separate SFE objects. A small number of nuclei polygons are invalid due to self-intersection; these cases were resolved by making a buffer of distance 0 and then removing the holes. Additional cell metadata provided by 10X, such as cell area, are also included.

Usage

```
JanesickBreastData(dataset = datasets, force = FALSE, verbose = TRUE)
```

Arguments

| | |
|---------|---|
| dataset | Which dataset to use, must be one of "rep1" and "rep2". |
| force | Logical, whether to force redownload if the files are already present. Defaults to FALSE. |
| verbose | Whether to display progress of download. |

Details

As the SFE and Voyager packages are in the experimental stage and they were originally developed and tested on relatively small Visium datasets, they are not yet very scalable to larger smFISH datasets. While 10X provided transcript spot locations, these are not included in the SFE objects for now as we are not sure if spatstat can work with such a large dataset for spatial point process analyses, nor does SFE integrate with spatstat. In a future version of this package, the transcript locations might be added as a separate dataset, but this is not guaranteed.

Value

A SpatialFeatureExperiment object.

LohoffGastrulationData

seqFISH mouse gastrulation dataset

Description

This dataset was downloaded from the [companion website](#) titled, Integration of spatial and single-cell transcriptomic data elucidates mouse organogenesis [Lohoff et al.](#) There are three biological replicates available in this dataset, each representing a different embryo. For each dataset, the raw gene counts, metadata, and cell segmentation in one z-plane are provided in the SFE object. Segmentation data were not provided for provided for all cells in the count matrix for embryos 1 and 2. In these cases, the segmentation data are represented by empty polygons.

Usage

```
LohoffGastrulationData(dataset = datasets, force = FALSE, verbose = TRUE)
```

Arguments

| | |
|---------|---|
| dataset | Which dataset to use, must be one of "rep1", "rep2", and "rep3". |
| force | Logical, whether to force redownload if the files are already present. Defaults to FALSE. |
| verbose | Whether to display progress of download. |

Details

While the authors provided the spot location for each mRNA molecule, these are not included in the SFE objects.

Value

A SpatialFeatureExperiment object.

McKellarMuscleData *Download McKellar et al. mouse skeletal muscle data*

Description

In the first version of this package, only the first time point, 2 days after notexin injury, is available. We may add the later time points in later versions of this package.

Usage

```
McKellarMuscleData(dataset = datasets, force = FALSE, verbose = TRUE)
```

Arguments

| | |
|---------|--|
| dataset | Which dataset to use. Whether the full dataset ("full"), the first small subset ("small"), or the second small subset ("small2"). The second small subset has a different <code>sample_id</code> . |
| force | Logical, whether to force redownload if the files are already present. Defaults to FALSE. |
| verbose | Whether to display progress of download. |

Details

All datasets are SpatialFeatureExperiment (SFE) objects, with a counts assay for the raw gene counts. Column metadata includes total UMI counts (`nCounts`) and number of genes (`nGenes`) detected per spot. Row metadata includes means, variances, and CV2 of each gene in the full dataset. Column geometry includes Visium spot polygons (`spotPoly`). Annotation geometry includes tissue boundary (`tissueBoundary`), myofiber segmentation (full resolution `myofiber_full` and simplified `myofiber_simplified`), nuclei segmentation (`nuclei`), and nuclei centroids (`nuclei_centroid`).

Myofibers were segmented manually with the LabKit ImageJ plugin on a 4x downsized H&E image, downsized so the image can be loaded into LabKit, and the `terra` R package was used to convert the TIFF segmentation masks into polygons. Coordinates in the SFE objects are in pixels in the full resolution H&E image. Hence the coordinates of the myofiber segmentations were scaled up to match the other coordinates. The full resolution myofiber segmentation looks pixelated; the `mapshaper` R package was used to simplify polygons while conserving contiguity. Morphological (area, perimeter, eccentricity, angle) and Haralick (see `EImage::computeFeatures.haralick`) metrics were computed for the myofibers with the `EImage` R package.

Nuclei were segmented with StarDist. About 3000 nuclei from randomly selected regions in the H&E image from this and later time points were manually annotated with LabKit to train the StarDist model, which was then used to segment all nuclei. OpenCV was used to convert segmentation masks into polygons and compute morphological metrics.

Tissue boundary was obtained by first thresholding the H&E image by grayscale intensity and then converting the mask into polygons with OpenCV. Small pieces which are debris were removed.

Value

A `SpatialFeatureExperiment` object.

Examples

```
sfe <- McKellarMuscleData("small")
```

SFEData

Example SpatialFeatureExperiment datasets

Description

Example spatial transcriptomics datasets with **Simple Features** annotations as `SpatialFeatureExperiment` objects.

Datasets

Full Visium dataset of the first time point, including spots outside tissue. ([McKellarMuscleData](#))

Small subset of the full Visium dataset for function examples. ([McKellarMuscleData](#))

A second small subset of the full Visium dataset with a different `sample_id` used for function examples involving multiple samples. ([McKellarMuscleData](#))

Slide-seq2 human melanoma brain metastasis dataset ([BiermannMelaMetasData](#))

Slide-seq2 human melanoma extracranial metastasis dataset ([BiermannMelaMetasData](#))

10X Xenium formalin fixed paraffin embedded (FFPE) Xenium dataset for human breast cancer (2 biological replica, [JanesickBreastData](#))

Nanostring CosMX FFPE human non small cell lung cancer data ([HeNSCLCData](#))

Vizgen MERFISH mouse liver data ([VizgenLiverData](#))

| | |
|-----------------|--|
| VizgenLiverData | <i>Vizgen MERFISH mouse liver data</i> |
|-----------------|--|

Description

This is one of the example datasets from Vizgen's website, downloaded from [here](#). The gene count matrix, cell metadata provided by Vizgen, QC metrics, and cell segmentation in one z-plane are included in the SFE object. While it appears that 7 z-planes are in the cell boundaries in the hdf5 files on the website, all 7 z-planes are the same, so only one was used here.

Usage

```
VizgenLiverData(dataset = datasets, force = FALSE, verbose = TRUE)
```

Arguments

| | |
|---------|---|
| dataset | Which dataset to use, for now can only be "Liver1Slice1". |
| force | Logical, whether to force redownload if the files are already present. Defaults to FALSE. |
| verbose | Whether to display progress of download. |

Details

This is the largest SFE example dataset thus far. While the SFE object can fit into memory due to the relatively small number of genes, we are considering making a HDF5Array version of this example dataset. Furthermore, the geometries of the large number of cells can also consume a lot of memory. We are considering using [Apache Sedona](#) and possibly [SQLDataFrame](#) for on disk geometries and geometric operations in a future version of SpatialFeatureExperiment and Voyager.

While Vizgen provides transcript spot locations, we don't yet know what to do with the huge dataset, so this is not included in the SFE object.

Value

A SpatialFeatureExperiment object.

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