

# Package ‘scRNAseq’

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**Title** Collection of Public Single-Cell RNA-Seq Datasets

**Version** 2.22.0

**Date** 2025-01-18

**Description** Gene-level counts for a collection of public scRNA-seq datasets,  
provided as SingleCellExperiment objects with cell- and gene-level metadata.

**License** CC0

**NeedsCompilation** no

**Depends** SingleCellExperiment

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DelayedArray, GenomicRanges, SummarizedExperiment,  
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AnnotationDbi, ensemblDb, GenomicFeatures, alabaster.base,  
alabaster.matrix, alabaster.sce, gypsum, jsonlite, DBI, RSQLite

**Suggests** BiocStyle, knitr, rmarkdown, testthat, jsonvalidate,  
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**VignetteBuilder** knitr

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SequencingData, RNASeqData, SingleCellData

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scRNAseq-package

*Collection of Public Single-Cell RNA-Seq Datasets*

## Description

Gene-level counts for a collection of public scRNA-seq datasets, provided as SingleCellExperiment objects with cell- and gene-level metadata.

## Details

This package contains a collection of three publicly available single-cell RNA-seq datasets.

The dataset fluidigm contains 65 cells from Pollen et al. (2014), each sequenced at high and low coverage.

The dataset th2 contains 96 T helper cells from Mahata et al. (2014).

The dataset allen contains 379 cells from the mouse visual cortex. This is a subset of the data published in Tasic et al. (2016).

See the package vignette for details on the pre-processing of the data.

**Author(s)**

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Maintainer: Aaron Lun <infinite.monkeys.with.keyboards@gmail.com>

**References**

Pollen, Nowakowski, Shuga, Wang, Leyrat, Lui, Li, Szpankowski, Fowler, Chen, Ramalingam, Sun, Thu, Norris, Lebofsky, Toppani, Kemp II, Wong, Clerkson, Jones, Wu, Knutsson, Alvarado, Wang, Weaver, May, Jones, Unger, Kriegstein, West. Low-coverage single-cell mRNA sequencing reveals cellular heterogeneity and activated signaling pathways in developing cerebral cortex. *Nature Biotechnology*, 32, 1053-1058 (2014).

Mahata, Zhang, Kolodziejczyk, Proserpio, Haim-Vilmovsky, Taylor, Hebenstreit, Dingler, Moignard, Gottgens, Arlt, McKenzie, Teichmann. Single-Cell RNA Sequencing Reveals T Helper Cells Synthesizing Steroids De Novo to Contribute to Immune Homeostasis. *Cell Reports*, 7(4): 1130–1142 (2014).

Tasic, Menon, Nguyen, Kim, Jarsky, Yao, Levi, Gray, Sorensen, Dolbeare, Bertagnolli, Goldy, Shapovalova, Parry, Lee, Smith, Bernard, Madisen, Sunkin, Hawrylycz, Koch, Zeng. Adult mouse cortical cell taxonomy revealed by single cell transcriptomics. *Nature Neuroscience*, 19, 335–346 (2016).

---

AztekinTailData	<i>Obtain the Aztekin tail data</i>
-----------------	-------------------------------------

---

**Description**

Obtain the *Xenopus* tail single-cell RNA-seq data from Aztekin et al. (2019).

**Usage**

```
AztekinTailData(legacy = FALSE)
```

**Arguments**

legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.
--------	---

**Details**

Column metadata is provided in the same form as supplied in E-MTAB-7761. This contains information such as the treatment condition, batch, putative cell type, putative cell cycle phase.

The UMAP results are available as the "UMAP" entry in the [reducedDims](#).

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/aztekin-tail`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Aztekin C et al. (2019). Identification of a regeneration-organizing cell in the *Xenopus* tail. *Science* 364(6441), 653-658

**Examples**

```
sce <- AztekinTailData()
```

---

BacherTCellData	<i>Obtain the Bacher T cell data</i>
-----------------	--------------------------------------

---

**Description**

Obtain the human COVID T cell single-cell RNA-seq dataset from Bacher et al. (2020).

**Usage**

```
BacherTCellData(  
  filtered = TRUE,  
  ensembl = FALSE,  
  location = TRUE,  
  legacy = FALSE  
)
```

**Arguments**

filtered	Logical scalar indicating whether to filter out cells that were not used by the authors.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata is scraped from GEO, using both the author-supplied TSV of per-cell annotations and the sample-level metadata. This contains information such as the diagnosis, severity, WHO class, clustering and clonotype.

If `filtered=TRUE`, only the cells used by the authors in their final analysis are returned. Otherwise, an additional `filtered` field will be present in the [colData](#), indicating whether the cell was retained by the authors.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/bacher-tcell`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Bacher P et al. (2020). Low avidity T cell responses to SARS-CoV-2 in unexposed individuals and severe COVID-19 *Immunity* 53, 1258-1271

**Examples**

```
if (.Machine$sizeof.pointer > 4) { # too large for 32-bit machines!
  sce <- BacherTCellData()
}
```

---

BachMammaryData

*Obtain the Bach mammary data*

---

**Description**

Obtain the mouse mammary gland single-cell RNA-seq data from Bach et al. (2017).

**Usage**

```
BachMammaryData(  
  samples = c("NP_1", "NP_2", "G_1", "G_2", "L_1", "L_2", "PI_1", "PI_2"),  
  location = TRUE,  
  legacy = FALSE  
)
```

**Arguments**

<code>samples</code>	A character vector with at least one element, specifying which samples(s) to retrieve.
<code>location</code>	Logical scalar indicating whether genomic coordinates should be returned.
<code>legacy</code>	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata is extracted from the sample annotation in GSE106273, and refers to the developmental stage of the mammary gland.

If multiple samples are specified in `samples`, the count matrices will be cbinded together. Cells originating from different samples are identifiable by the "Sample" field in the column metadata.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scrNAseq/bach-mammary`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Bach K et al. (2017). Differentiation dynamics of mammary epithelial cells revealed by single-cell RNA sequencing. *Nat Commun.* 8(1), 2128

**Examples**

```
sce <- BachMammaryData(samples="NP_1")
```

---

BaronPancreasData	<i>Obtain the Baron pancreas data</i>
-------------------	---------------------------------------

---

## Description

Obtain the human/mouse pancreas single-cell RNA-seq data from Baron et al. (2017).

## Usage

```
BaronPancreasData(  
  which = c("human", "mouse"),  
  ensembl = FALSE,  
  location = TRUE,  
  legacy = FALSE  
)
```

## Arguments

which	String specifying the species to get data for.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column metadata is provided in the same form as supplied in GSE84133. This contains information such as the cell type labels and donor ID (for humans) or strain (for mouse).

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/baron-pancreas`.

## Value

A [SingleCellExperiment](#) object with a single matrix of read counts.

## Author(s)

Aaron Lun



## References

Baron M et al. (2017). Single-cell transcriptomic map of the human and mouse pancreas reveals inter- and intra-cell population structure. *Cell Syst.* 3(4), 346-360.

## Examples

```
sce.human <- BaronPancreasData()

sce.mouse <- BaronPancreasData("mouse")
```

---

BhaduriOrganoidData      *Obtain the Bhaduri cortical organoid data*

---

## Description

Obtain the human cortical organoid single-cell RNA-seq dataset from Bhaduri et al. (2020).

## Usage

```
BhaduriOrganoidData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

## Arguments

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column data contains sample-level information. In theory, there is also cell-level metadata for this dataset but it could not be unambiguously mapped to the column names.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/bhaduri-organoid`.

## Value

A [SingleCellExperiment](#) object with a single matrix of normalized expression values.

**Author(s)**

Aaron Lun

**References**

Bhaduri A et al. (2020). Cell stress in cortical organoids impairs molecular subtype specification. *Nature* 578(7793), 142-148.

**Examples**

```
if (.Machine$sizeof.pointer > 4) { # too large for 32-bit machines!
  sce <- BhaduriOrganoidData()
}
```

---

BuettnerESCData

*Obtain the Buettner ESC data*

---

**Description**

Obtain the mouse embryonic stem cell single-cell RNA-seq data from Buettner et al. (2015).

**Usage**

```
BuettnerESCData(remove.htseq = TRUE, location = TRUE, legacy = FALSE)
```

**Arguments**

<code>remove.htseq</code>	Logical scalar indicating whether HT-seq alignment statistics should be removed.
<code>location</code>	Logical scalar indicating whether genomic coordinates should be returned.
<code>legacy</code>	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Rows corresponding to HT-seq's alignment statistics are removed by default. These can be retained by setting `remove.htseq=FALSE`.

Column metadata contains the experimentally determined cell cycle phase for each cell.

Counts for ERCC spike-ins are stored in the "ERCC" entry in the [altExps](#).

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/buettner-esc`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of read counts.

**Author(s)**

Aaron Lun

**References**

Buettner F et al. (2015). Computational analysis of cell-to-cell heterogeneity in single-cell RNA-sequencing data reveals hidden subpopulations of cells. *Nat. Biotechnol.* 33(2), 155-160.

**Examples**

```
sce <- BuettnerESData()
```

BunisHSPCData

*Obtain the Bunis haematopoietic stem and progenitor cell data***Description**

Obtain the human fetal, newborn, and adult haematopoietic stem and progenitor cell single-cell RNA-seq dataset from Bunis et al. (2021).

**Usage**

```
BunisHSPCData(filtered = TRUE, legacy = FALSE)
```

**Arguments**

filtered	Logical scalar or "cells" indicating whether to: <ul style="list-style-type: none"> <li>• TRUE: filter out cells that were not used by the authors.</li> <li>• "cells": filter out empty droplets as filtered out by cell ranger.</li> <li>• FALSE: no filtering</li> </ul>
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata is recreated from GEO using the author-supplied TSV of per-cell annotations, or retrieved from a processed version of the data shared by authors via figshare. This contains information such as the tissue & sample of origin, age group, likely cell type, and Developmental Stage Scoring. Within DevStageScoring element of the column metadata are the applied results ('<cell\_type>\_scores') of random forest regression trained on the fetal (score = 0) and adult (score = 1) cells of individual cell types indicated by ('<cell\_type>\_inTraining').

If filtered=TRUE, only the cells used by the authors in their final analysis are returned. Otherwise, an additional retained field will be present in the `colData`, indicating whether the cell was retained by the authors.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/bunis-hspc`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Daniel Bunis

**References**

Bunis DG et al. (2021). Single-Cell Mapping of Progressive Fetal-to-Adult Transition in Human Naive T Cells *Cell Rep.* 34(1): 108573

**Examples**

```
sce <- BunishSPCData()
```

---

CampbellBrainData	<i>Obtain the Campbell brain data</i>
-------------------	---------------------------------------

---

**Description**

Obtain the mouse brain single-cell RNA-seq data from Campbell et al. (2017).

**Usage**

```
CampbellBrainData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

**Arguments**

ensembl	Logical scalar indicating whether the row names of the returned object should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata is provided in the same form as supplied in GSE93374. This contains information such as the diet of the mice, sex and proposed cell type for each cell.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/campbell-brain`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Campbell R et al. (2017). A molecular census of arcuate hypothalamus and median eminence cell types. *Nat. Neurosci.* 20, 484-496.

**Examples**

```
sce <- CampbellBrainData()
```

---

ChenBrainData	<i>Obtain the Chen brain data</i>
---------------	-----------------------------------

---

**Description**

Obtain the mouse brain single-cell RNA-seq data from Chen et al. (2017).

**Usage**

```
ChenBrainData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

**Arguments**

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata is provided in the same form as supplied in GSE87544. This contains the putative cell type assigned by the original authors.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/chen-brain`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Chen R et al. (2017). Single-Cell RNA-Seq reveals hypothalamic cell diversity. *Cell Rep.* 18, 3227-3241.

**Examples**

```
sce <- ChenBrainData()
```

---

countErccMolecules	<i>Obtain ERCC molecule counts from concentrations</i>
--------------------	--

---

**Description**

Compute the number of molecules for each transcript in the ERCC spike-in mixture, based on their published concentration as well as the volume of the diluted mixture added to each cell.

**Usage**

```
countErccMolecules(volume, dilution, mix = c("1", "2"), ...)
```

**Arguments**

volume	Numeric scalar specifying the added volume (in microliters) of ERCC spike-in mixture.
dilution	Numeric scalar specifying the dilution factor used for the added volume of the spike-in mixture.
mix	String specifying whether to compute the number of molecules for mix 1 or 2.
...	Further arguments to pass to <a href="#">fetchDataset</a> .

**Value**

A [DataFrame](#) object with one row per ERCC spike-in transcript. This contains the estimated concentration and molecule count for each transcript.

**Author(s)**

Aaron Lun, based on code from Alan O'Callaghan

**Examples**

```
countErccMolecules(volume = 9, dilution = 300000)
```

---

DarmanisBrainData	<i>Obtain the Darmanis brain data</i>
-------------------	---------------------------------------

---

**Description**

Obtain the human brain single-cell RNA-seq dataset from Darmanis et al. (2015).

**Usage**

```
DarmanisBrainData(
  ensembl = FALSE,
  location = TRUE,
  remove.htseq = TRUE,
  legacy = FALSE
)
```

**Arguments**

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
remove.htseq	Logical scalar indicating whether HT-seq alignment statistics should be removed.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata is scraped from GEO and includes patient information, tissue of origin and likely cell type.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. This is only performed when `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/darmanis-brain`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Darmanis S et al. (2015). A survey of human brain transcriptome diversity at the single cell level. *Proc Natl Acad Sci USA* 112, 7285-90.

**Examples**

```
sce <- DarmanisBrainData()
```

---

ERCCSpikeInConcentrations

*Obtain ERCC concentrations*

---

**Description**

Obtain ERCC spike-in concentrations from the Thermo Fisher Scientific website.

**Usage**

```
ERCCSpikeInConcentrations(  
  volume = NULL,  
  dilution = NULL,  
  mix = c("1", "2"),  
  legacy = FALSE  
)
```

**Arguments**

volume	Numeric scalar specifying the added volume (in nanoliters) of ERCC spike-in mixture. Only used if dilution is specified.
dilution	Numeric scalar specifying the dilution factor used for the added volume of the spike-in mixture. Only used if volume is specified.
mix	String specifying whether to compute the number of molecules for mix 1 or 2. Only used if both dilution and volume are specified.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

If volume and dilution are specified, an additional column is added to the output specifying the number of molecules of spike-in transcript for the specified mix.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/ercc-concentrations`.



**Value**

A [DataFrame](#) object with one row per ERCC spike-in transcript. This contains information such as the spike-in concentration in each mix.

**Author(s)**

Alan O’Callaghan

**Examples**

```
df <- ERCCSpikeInConcentrations()
```

---

```
ErnstSpermatogenesisData
```

*Obtain the Ernst spermatogenesis data*

---

**Description**

Obtain the mouse spermatogenesis single-cell RNA-seq dataset from Ernst et al. (2019).

**Usage**

```
ErnstSpermatogenesisData(
  method = c("emptyDrops", "Cellranger"),
  location = TRUE,
  legacy = FALSE
)
```

**Arguments**

method	String indicating which cell caller to obtain results for.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

This study contains two analyses done with datasets from different cell calling algorithms. One uses Cellranger version 2 while the other uses emptyDrops from **DropletUtils**.

Column metadata includes sample information, per-cell QC metrics and cell type labels. In particular, the sample label specifies the developmental stage of the mouse.

Note that method="Cellranger" contains additional data for Tc1 mice. These mice have an additional human chromosome 21 inserted alongside the usual mouse chromosomes.

If location=TRUE, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/ernst-spermatogenesis`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Ernst C et al. (2019). Staged developmental mapping and X chromosome transcriptional dynamics during mouse spermatogenesis. *Nat Commun* 10, 1251

**Examples**

```
if (.Machine$sizeof.pointer > 4) { # too large for 32-bit machines!
  sce <- ErnstSpermatogenesisData()
}
```

---

fetchDataset

*Fetch a dataset from the gypsum backend*

---

**Description**

Fetch a dataset (or its metadata) from the gypsum backend.

**Usage**

```
fetchDataset(
  name,
  version,
  path = NA,
  package = "scRNAseq",
  cache = cacheDirectory(),
  overwrite = FALSE,
  realize.assays = FALSE,
  realize.reduced.dims = TRUE,
  ...
)

fetchMetadata(
  name,
  version,
  path = NA,
  package = "scRNAseq",
  cache = cacheDirectory(),
  overwrite = FALSE
)
```

**Arguments**

name	String containing the name of the dataset.
version	String containing the version of the dataset.
path	String containing the path to a subdataset, if name consists of multiple sub-datasets. Defaults to NA if no subdatasets are present.
package	String containing the name of the package.
cache, overwrite	Arguments to pass to <a href="#">saveVersion</a> or <a href="#">saveFile</a> .
realize.assays, realize.reduced.dims	Logical scalars indicating whether to realize assays and reduced dimensions into memory. Dense and sparse <a href="#">ReloadedArray</a> objects are converted into ordinary arrays and dgCMatix objects, respectively.
...	Further arguments to pass to <a href="#">readObject</a> .

**Value**

fetchDataset returns the dataset as a [SummarizedExperiment](#) or one of its subclasses.

fetchMetadata returns a named list of metadata for the specified dataset.

**Author(s)**

Aaron Lun

**See Also**

<https://github.com/ArtifactDB/bioconductor-metadata-index>, on the expected schema for the metadata.

[saveDataset](#) and [uploadDirectory](#), to save and upload a dataset.

[surveyDatasets](#) and [listVersions](#), to get possible values for name and version.

**Examples**

```
fetchDataset("zeisel-brain-2015", "2023-12-14")
fetchMetadata("zeisel-brain-2015", "2023-12-14")
```

---

FletcherOlfactoryData *Obtain the Fletcher Olfactory data*

---

**Description**

Obtain the mouse olfactory epithelial HBC stem cell differentiation dataset from Fletcher et al. (2017).

**Usage**

```
FletcherOlfactoryData(
  filtered = TRUE,
  ensembl = FALSE,
  location = TRUE,
  legacy = FALSE
)
```

**Arguments**

filtered	Logical scalar indicating whether to filter out cells that were not used by the authors.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata is scraped from GEO, using both the author-supplied “phenoData” per-cell annotations and the author-supplied “protocolData” per-cell annotations. The former includes information about the animals and the instruments, while the latter contains QC statistics.

We also included the clustering results from the authors’ analysis.

If `filtered=TRUE`, only the cells used by the authors in their cluster analysis are returned. Otherwise, the cells not used by the authors will have NA in the clustering columns of the `colData`.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/fletcher-olfactory`.

**Value**

A `SingleCellExperiment` object with a single matrix of read counts.

**Author(s)**

Davide Risso

**References**

Fletcher R et al. (2017). Deconstructing olfactory stem cell trajectories at single-cell resolution. *Cell Stem Cell* 20, 817-30.

**Examples**

```
sce <- FletcherOlfactoryData()
```

---

GiladiHSCData

*Obtain the Giladi HSC data*


---

**Description**

Obtain the mouse haematopoietic stem cell single-cell RNA-seq and CRISPR-seq dataset from Giladi et al. (2018).

**Usage**

```
GiladiHSCData(
  mode = c("rna", "crispr"),
  filtered = TRUE,
  ensembl = FALSE,
  location = TRUE,
  legacy = FALSE
)
```

**Arguments**

mode	Character vector specifying which modalities should be returned.
filtered	Logical scalar indicating whether to filter out cells that were not used by the authors.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers, when mode contains "rna".
location	Logical scalar indicating whether genomic coordinates should be returned, when mode contains "rna".
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata is scraped from GEO using the author-supplied TSV of per-cell annotations. This contains information such as the batch of origin for each cell plus an array of FACS measurements per cell.

If filtered=TRUE, only the cells used by the authors in their final analysis are returned. Otherwise, an additional filtered field will be present in the [colData](#), indicating whether the cell was retained by the authors.

If ensembl=TRUE, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated

IDs is retained. For row names with multiple semi-colon-delimited symbols, the last symbol is used for matching against the Ensembl annotation.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. This is only relevant when `ensembl=TRUE`.

If `mode` contains multiple modalities, the intersection of cells that are present in both modalities is returned. This is because not all cells have data across both modalities. If `mode` contains only one modality, all cells for that modality are returned.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/giladi-hsc`.

### Value

A `SingleCellExperiment` object with a matrix of UMI counts for the scRNA-seq or CRISPR-seq data. Alternatively, an object with both count matrices, where the second modality is stored as an alternative Experiment.

### Author(s)

Aaron Lun

### References

Giladi A et al. (2018). Single-cell characterization of haematopoietic progenitors and their trajectories in homeostasis and perturbed haematopoiesis. *Nat Cell Biol* 20, 836-846

### Examples

```
if (.Machine$sizeof.pointer > 4) { # too large for 32-bit machines!
  sce <- GiladiHSCData()
}
```

---

GrunHSCData

*Obtain the Grun HSC data*

---

### Description

Obtain the mouse haematopoietic stem cell single-cell RNA-seq data from Grun et al. (2016).

### Usage

```
GrunHSCData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

### Arguments

<code>ensembl</code>	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
<code>location</code>	Logical scalar indicating whether genomic coordinates should be returned.
<code>legacy</code>	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Row metadata contains the symbol and chromosomal location for each gene. Column metadata contains the extraction protocol used for each sample, as described in GSE76983.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scrNaseq/grun-hsc`.

**Value**

A `SingleCellExperiment` object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Grun D et al. (2016). De novo prediction of stem cell identity using single-cell transcriptome data. *Cell Stem Cell* 19(2), 266-277.

**Examples**

```
sce <- GrunHSCData()
```

---

GrunPancreasData	<i>Obtain the Grun pancreas data</i>
------------------	--------------------------------------

---

**Description**

Obtain the human pancreas single-cell RNA-seq data from Grun et al. (2016).

**Usage**

```
GrunPancreasData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

**Arguments**

<code>ensembl</code>	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
<code>location</code>	Logical scalar indicating whether genomic coordinates should be returned.
<code>legacy</code>	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Row metadata contains fields for the symbol and chromosomal location of each gene, as derived from the row names.

Column metadata is derived from the column names of the count matrix with the sample annotations in GSE81076. This includes the donor identity for each cell and the type of sample.

The "ERCC" entry in the [altExps](#) contains count data for the ERCC spike-in transcripts.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/grun-pancreas`.

## Value

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

## Author(s)

Aaron Lun, using additional metadata obtained by Vladimir Kiselev.

## References

Grun D et al. (2016). De novo prediction of stem cell identity using single-cell transcriptome data. *Cell Stem Cell* 19(2), 266-277.

## Examples

```
sce <- GrunPancreasData()
```

---

HeOrganAtlasData

*Obtain the He organ atlas data*

---

## Description

Obtain the human cortex single-nuclei RNA-seq data from Hu et al. (2017).



**Usage**

```
HeOrganAtlasData(
  tissue = c("Bladder", "Blood", "Common.bile.duct", "Esophagus", "Heart", "Liver",
    "Lymph.node", "Marrow", "Muscle", "Rectum", "Skin", "Small.intestine", "Spleen",
    "Stomach", "Trachea"),
  ensembl = FALSE,
  location = TRUE,
  legacy = FALSE
)
```

**Arguments**

tissue	Character vector specifying the tissues to return.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column data contains the tissue of origin, a variety of per-cell QC metrics well as some cell type annotations. The reclustered annotations required some assembly:

- `reclustered.broad` was generated based on whether the barcode was present in each `*_meta.data.txt` file at <https://github.com/bei-lab/scRNA-AHCA>.
- For each barcode that was present in one of those files, `reclustered.fine` was generated based on the label in the annotation field inside that file.

If multiple tissues are requested, counts are only reported for the intersection of genes across all tissues. This is because the gene annotation in the original count matrices differs across tissues.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/he-organ-atlas`.

**Value**

A `SingleCellExperiment` object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

## References

He S et al. (2020). Single-cell transcriptome profiling of an adult human cell atlas of 15 major organs. *Genome Biol* 21, 1:294.

## Examples

```
if (.Machine$sizeof.pointer > 4) { # too large for 32-bit machines!
  sce <- HeOrganAtlasData()
}
```

---

HermannSpermatogenesisData

*Obtain the Hermann spermatogenesis data*

---

## Description

Obtain the mouse spermatogenesis single-cell RNA-seq data from Hermann et al. (2018).

## Usage

```
HermannSpermatogenesisData(strip = FALSE, location = TRUE, legacy = FALSE)
```

## Arguments

strip	Logical scalar indicating whether to strip the .X notation from the row names.
location	Logical scalar indicating whether genomic coordinates should be returned. Only used if strip=TRUE.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column metadata contains cell types provided by the data generators at <https://data.mendeley.com/datasets/kxd5f8vpt4/1#file-fe79c10b-c42e-472e-9c7e-9a9873d9b3d8>.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for scRNAseq/hermann-spermatogenesis.

## Value

A [SingleCellExperiment](#) object with two matrices, containing spliced and unspliced counts, respectively.

## Author(s)

Charlotte Soneson

## References

Hermann B.P. et al. (2018). The Mammalian Spermatogenesis Single-Cell Transcriptome, from Spermatogonial Stem Cells to Spermatids. *Cell Rep.* 25(6), 1650-1667.e8.

## Examples

```
sce <- HermannSpermatogenesisData()
```

---

HuCortexData	<i>Obtain the Hu cortex data</i>
--------------	----------------------------------

---

## Description

Obtain the mouse cortex single-nuclei RNA-seq data from Hu et al. (2017).

## Usage

```
HuCortexData(
  mode = c("ctx", "3T3"),
  samples = NULL,
  ensembl = FALSE,
  location = TRUE,
  legacy = FALSE
)
```

## Arguments

mode	Character vector indicating whether to return data for the 3T3 cells or the mouse cortex.
samples	Character vector indicating whether to return data for specific samples, see Details. If specified, this overrides mode.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column metadata includes the mode and sample corresponding to each cell/nuclei. Available samples are:

- "cell-3T3" and "nuclei-3T3", generated from the 3T3 cell line.
- "nuclei-ctx-X", nuclei generated from the cortex of animal number X (from 1 to 13).

- "nuclei-ctx-salineX" or "nuclei-ctx-PTZX", nuclei generated from the cortex of saline- or PTZ-treated mice. X represents the replicate number and can be 1 or 2.

If multiple modes are requested, counts are only reported for the intersection of genes across all modes. This is because the gene annotation in the original count matrices differs across modes.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/wu-kidney`.

### Value

A `SingleCellExperiment` object with a single matrix of read counts.

### Author(s)

Aaron Lun

### References

Hu P et al. (2017). Dissecting cell-type composition and activity-dependent transcriptional state in mammalian brains by massively parallel single-nucleus RNA-seq. *Mol. cell* 68, 1006-1015.

### Examples

```
sce <- HuCortexData("3T3")
```

---

JessaBrainData

*Obtain the Jessa brain data*

---

### Description

Obtain the mouse brain single-cell RNA-seq dataset from Jessa et al. (2019).

### Usage

```
JessaBrainData(filtered = TRUE, location = TRUE, legacy = FALSE)
```

### Arguments

<code>filtered</code>	Logical scalar indicating whether to filter out cells that were not used by the authors.
<code>location</code>	Logical scalar indicating whether genomic coordinates should be returned.
<code>legacy</code>	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

If `filtered=TRUE`, only the cells used by the authors in their final analysis are returned. Otherwise, an additional `filtered` field will be present in the `colData`, indicating whether the cell was retained by the authors.

The column data contains sample of origin, some QC metrics and various cluster assignments for each cell. Cluster assignments starting with `Sample_*` are derived from per-sample analyses and cannot be compared sensibly across samples. Other clusterings (`Forebrain_*` and `Pons_*`) are derived from joint analyses across all samples involving the named tissue.

The `reducedDims` of the output contains various dimensionality reduction results. Coordinates for entries prefixed with `Sample_*` were generated from per-sample analyses and cannot be compared across samples. Coordinates for entries prefixed with `Forebrain_*` and `Pons_*` were generated from joint analyses from the corresponding tissue.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/jessa-brain`.

**Value**

A `SingleCellExperiment` object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Jessa S et al. (2019). Stalled developmental programs at the root of pediatric brain tumors *Nat Genet* 51, 1702-1713

**Examples**

```
if (.Machine$sizeof.pointer > 4) { # too large for 32-bit machines!
  sce <- JessaBrainData()
}
```

---

KolodziejczykESCData    *Obtain the Kolodziejczyk ESC data*

---

**Description**

Obtain the mouse embryonic stem cell single-cell RNA-seq data from Kolodziejczyk et al. (2015).

**Usage**

```
KolodziejczykESCData(remove.htseq = TRUE, location = TRUE, legacy = FALSE)
```

**Arguments**

<code>remove.htseq</code>	Logical scalar indicating whether HT-seq alignment statistics should be removed.
<code>location</code>	Logical scalar indicating whether genomic coordinates should be returned.
<code>legacy</code>	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata is generated from the column names, and contains the culture conditions and the plate of origin for each cell.

Count data for ERCC spike-ins are stored in the "ERCC" entry in the [altExps](#).

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/kolodziejczyk-esc`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of read counts.

**Author(s)**

Aaron Lun

**References**

Messmer T et al. (2019). Transcriptional heterogeneity in naive and primed human pluripotent stem cells at single-cell resolution. *Cell Rep* 26(4), 815-824.e4

**Examples**

```
sce <- KolodziejczykESCData()
```

---

KotliarovPBMCDData

---

*Obtain the Kotliarov CITE-seq data*


---

**Description**

Obtain the Kotliarov PBMC CITE-seq data from Kotliarov et al. (2020).

**Usage**

```
KotliarovPBMCDData(
  mode = c("rna", "adt"),
  ensembl = FALSE,
  location = TRUE,
  legacy = FALSE
)
```

**Arguments**

mode	Character vector specifying whether to return either or both the RNA and ADT counts.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

This dataset contains 20 samples from 2 experimental batches, where each batch contains 5 high and 5 low responders. The 10 samples per batch were mixed and distributed across the 6 lanes using a cell hashing approach.

The column metadata contains the following fields:

- sample\*: identifiers for the sample of origin for each cell.
- adjmfc.time: type of responder for each sample.
- tenx\_lane: 10X lane from which each cell was collected.
- batch: the batch of origin.
- barcode\_check: barcode identifier.
- hash\_\* and hto\_\* columns: **HTOdemux** outputs.
- DEMUXLET.\* columns: **demuxlet** outputs.
- joint\_classification\_global: **HTOdemux** and **demuxlet** joint classification.
- nGene: number of genes as defined from **Seurat**'s CreateSeuratObject.
- nUMI: number of UMIs as defined from **Seurat**'s CreateSeuratObject.
- pctMT: percent of mitochondrial reads as defined from **Seurat**'s CreateSeuratObject.

Note, no filtering has been performed based on the quality control metrics.

If ensembl=TRUE, the gene symbols in the RNA data are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If location=TRUE, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) for the RNA data. Note that this is only performed if ensembl=TRUE.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/kotliarov-pbmc`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts corresponding to the first mode, with an optional alternative Experiment if there is a second mode.

**Author(s)**

Stephany Orjuela, with modifications from Aaron Lun

**References**

Kotliarov et al. (2020). Broad immune activation underlies shared set point signatures for vaccine responsiveness in healthy individuals and disease activity in patients with lupus. *Nat. Med.* 26, 618–629

**Examples**

```
sce <- KotliarovPBMCData()
```

---

LaMannoBrainData	<i>Obtain the La Manno brain data</i>
------------------	---------------------------------------

---

**Description**

Obtain the mouse/human brain scRNA-seq data from La Manno et al. (2016).

**Usage**

```
LaMannoBrainData(
  which = c("human-es", "human-embryo", "human-ips", "mouse-adult", "mouse-embryo"),
  ensembl = FALSE,
  location = TRUE,
  legacy = FALSE
)
```

**Arguments**

which	A string specifying which dataset should be obtained.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.



## Details

Column metadata is provided in the same form as supplied in the supplementary tables in GSE71585. This contains information such as the time point and cell type.

The various settings of which will obtain different data sets.

- "human-es", human embryonic stem cells.
- "human-embryo", human embryo midbrain.
- "human-ips", human induced pluripotent stem cells.
- "mouse-adult", mouse adult dopaminergic neurons.
- "mouse-embryo", mouse embryo midbrain.

Unfortunately, each of these datasets uses a different set of features. If multiple datasets are to be used simultaneously, users will have to decide how to merge them, e.g., by taking the intersection of common features across all datasets.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/lamanno-brain`.

## Value

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

## Author(s)

Aaron Lun

## References

La Manno A et al. (2016). Molecular diversity of midbrain development in mouse, human, and stem cells. *Cell* 167(2), 566-580.

## Examples

```
sce.h.es <- LaMannoBrainData()

sce.h.em <- LaMannoBrainData("human-embryo")

sce.h.ip <- LaMannoBrainData("human-ips")

sce.m.ad <- LaMannoBrainData("mouse-adult")

sce.m.em <- LaMannoBrainData("mouse-embryo")
```

---

LawlorPancreasData	<i>Obtain the Lawlor pancreas data</i>
--------------------	--

---

## Description

Provides the human pancreas single-cell RNA-seq data from Lawlor et al. (2017).

## Usage

```
LawlorPancreasData(legacy = FALSE)
```

## Arguments

legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.
--------	---

## Details

Column metadata is provided in the same form as supplied in GSE86469. This contains information such as the cell type labels and patient status.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/lawlor-pancreas`.

## Value

A [SingleCellExperiment](#) object with a single matrix of read counts.

## Author(s)

Aaron Lun

## References

Lawlor N et al. (2017). Single-cell transcriptomes identify human islet cell signatures and reveal cell-type-specific expression changes in type 2 diabetes. *Genome Res.* 27(2), 208-222.

## Examples

```
sce <- LawlorPancreasData()
```

---

LedergorMyelomaData	<i>Obtain the Ledergor Myeloma data</i>
---------------------	---

---

## Description

Obtain the human multiple myeloma single-cell RNA-seq data from Ledergor et al. (2018).

## Usage

```
LedergorMyelomaData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

## Arguments

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column metadata was created from the sample metadata file in GSE117156. It contains an 'Experiment\_ID' column, from which the tissue and subject of origin were extracted, as well as the condition and treatment status of the subject.

If ensembl=TRUE, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If location=TRUE, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output. Note that this is only performed if ensembl=TRUE.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/ledergor-myeloma`.

## Value

A [SingleCellExperiment](#) object with a single matrix of read counts.

## Author(s)

Milan Malfait

## References

Ledergor G et al. (2018) Single cell dissection of plasma cell heterogeneity in symptomatic and asymptomatic myeloma. *Nat Med.* 24(12), 1867–1876.

**Examples**

```
sce <- LedergerMyelomaData()
```

---

LengESCData

---

*Obtain the Leng ESC data*


---

**Description**

Obtain the human embryonic stem cell single-cell RNA-seq data from Leng et al. (2015).

**Usage**

```
LengESCData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

**Arguments**

ensembl	Logical scalar indicating whether gene symbols should be converted to Ensembl annotation.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata contains the cell line, experiment number and experimentally determined cell cycle phase for each cell.

If ensembl=TRUE, the gene symbols in the published annotation are converted to Ensembl. If location=TRUE, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for scRNAseq/leng-esc.

**Value**

A [SingleCellExperiment](#) object with a single matrix of normalized expected read counts.

**Author(s)**

Aaron Lun

**References**

Leng F et al. (2015). Oscope identifies oscillatory genes in unsynchronized single-cell RNA-seq experiments. *Nat. Methods* 12(10), 947-950.

**Examples**

```
sce <- LengESCData()
```

---

listDatasets	<i>List all available datasets</i>
--------------	------------------------------------

---

**Description**

Summary information for all available datasets in the **scRNAseq** package.

**Usage**

```
listDatasets()
```

**Details**

A study may contribute multiple datasets if they cannot be reasonably combined (e.g., different species). The reported number of cells refers only to the dataset as it is stored in **scRNAseq**; this may be different to the number of cells used by the authors in their analysis, e.g., due to filtering.

**Value**

A [DataFrame](#) where each row corresponds to a dataset, containing the fields:

- Reference, a Markdown-formatted citation to scripts/ref.bib in the **scRNAseq** installation directory.
- Taxonomy, an identifier for the organism.
- Part, the part of the organism being studied.
- Number, the total number of cells in the dataset.
- Call, the relevant R call required to construct the dataset.

**Author(s)**

Aaron Lun

**Examples**

```
listDatasets()
```

---

listPaths	<i>List all paths for a dataset</i>
-----------	-------------------------------------

---

## Description

List the available paths to subdatasets within a version of a given dataset.

## Usage

```
listPaths(  
  name,  
  version,  
  package = "scRNAseq",  
  cache = cacheDirectory(),  
  overwrite = FALSE,  
  include.metadata = FALSE  
)
```

## Arguments

name	String containing the name of the dataset.
version	String containing the version of the dataset.
package	String containing the name of the package.
cache, overwrite	Arguments to pass to <a href="#">saveVersion</a> or <a href="#">saveFile</a> .
include.metadata	Logical scalar indicating whether to report the metadata for each subdataset.

## Value

If `include.metadata=FALSE`, a character vector containing the paths to subdatasets within the specified version of the dataset. If name does not contain any subdatasets, NA is returned.

Otherwise, a [DFrame](#) is returned containing the metadata for each subdataset, e.g., the title and description. More details can be found in the Bioconductor metadata schema at <https://github.com/ArtifactDB/bioconductor-metadata-index>.

## Author(s)

Aaron Lun

## Examples

```
listPaths("he-organs-2020", "2023-12-21")  
listPaths("he-organs-2020", "2023-12-21", include.metadata=TRUE)  
listPaths("zeisel-brain-2015", "2023-12-14") # no subdatasets  
listPaths("zeisel-brain-2015", "2023-12-14", include.metadata=TRUE)
```

---

listVersions	<i>List available versions</i>
--------------	--------------------------------

---

**Description**

List the available and latest versions for a dataset.

**Usage**

```
listVersions(name)

fetchLatestVersion(name)
```

**Arguments**

name	String containing the name of the dataset.
------	--

**Value**

For listVersions, a character vector containing the names of the available versions of the dataset.  
For fetchLatestVersion, a string containing the name of the latest version.

**Author(s)**

Aaron Lun

**Examples**

```
listVersions("zeisel-brain-2015")
fetchLatestVersion("zeisel-brain-2015")
```

---

LunSpikeInData	<i>Obtain the Lun spike-in data</i>
----------------	-------------------------------------

---

**Description**

Obtain the spike-in single-cell RNA-seq data from Lun et al. (2017).

**Usage**

```
LunSpikeInData(
  which = c("416b", "tropho"),
  split.oncogene = FALSE,
  location = TRUE,
  legacy = FALSE
)
```

## Arguments

<code>which</code>	String specifying whether the 416B or trophoblast data should be obtained.
<code>split.oncogene</code>	Logical scalar indicating whether the oncogene should be split to a separate <a href="#">altExp</a> .
<code>location</code>	Logical scalar indicating whether genomic coordinates should be returned.
<code>legacy</code>	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Row data contains a single "Length" field describing the total exonic length of each feature.

Column metadata is provided in the same form as supplied in E-MTAB-5522. This contains information such as the cell type, plate of origin, spike-in addition order and oncogene induction.

Two sets of spike-ins were added to each cell in each dataset. These are available as the "SIRV" and "ERCC" entries in the [altExps](#).

If `split.oncogene=TRUE` and `which="416b"`, the CBFB-MYH11-mcherry oncogene is moved to extra "oncogene" entry in the [altExps](#).

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/lun-spikein`.

## Value

A [SingleCellExperiment](#) object with a single matrix of read counts.

## Author(s)

Aaron Lun

## References

Lun ATL et al. (2017). Assessing the reliability of spike-in normalization for analyses of single-cell RNA sequencing data. *Genome Res.* 27(11), 1795-1806.

## Examples

```
sce <- LunSpikeInData()

sce <- LunSpikeInData("tropho")
```



---

MacoskoRetinaData	<i>Obtain the Macosko retina data</i>
-------------------	---------------------------------------

---

## Description

Obtain the mouse retina single-cell RNA-seq data from Macosko et al. (2016).

## Usage

```
MacoskoRetinaData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

## Arguments

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column metadata contains the cluster identity as reported in the paper. Note that some cells will have NA identities as they are present in the count matrix but not in the metadata file. These are presumably low-quality cells that were discarded prior to clustering.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/macosko-retina`.

## Value

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

## Author(s)

Aaron Lun

## References

Macosko E et al. (2016). Highly parallel genome-wide expression profiling of individual cells using nanoliter droplets. *Cell* 161(5), 1202-1214.

**Examples**

```
sce <- MacoskoRetinaData()
```

---

MairPBMCDData

---

*Obtain the Mair CITE-seq data*


---

**Description**

Obtain the Mair PBMC targeted CITE-seq data from Mair et al. (2020).

**Usage**

```
MairPBMCDData(
  mode = c("rna", "adt"),
  ensembl = FALSE,
  location = TRUE,
  legacy = FALSE
)
```

**Arguments**

mode	Character vector specifying whether to return either or both the RNA and ADT counts.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata contains the donor identity and cartridge of origin. Some libraries may also be classified as multiplets or have undeterminate origins after hash tag debarcoding.

If `ensembl=TRUE`, the gene symbols in the RNA data are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) for the RNA data. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/mair-pbmc`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts corresponding to the first mode, with an optional alternative Experiment if there is a second mode.

**Author(s)**

Stephany Orjuela, with modifications from Aaron Lun

**References**

Mair C et al. (2020). A targeted multi-omic analysis approach measures protein expression and low-abundance transcripts on the single-cell level. *Cell Rep.* 31, 107499

**Examples**

```
sce <- MairPBMCData()
```

---

MarquesBrainData	<i>Obtain the Marques brain data</i>
------------------	--------------------------------------

---

**Description**

Obtain the mouse brain single-cell RNA-seq data from Marques et al. (2016).

**Usage**

```
MarquesBrainData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

**Arguments**

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata is provided in the same form as supplied in GSE75330. This contains information such as the cell type and age/sex of the mouse of origin for each cell.

Note that some genes may be present in multiple rows corresponding to different genomic locations. These additional rows are identified by a `_loc[2-9]` suffix in their row names. Users may wish to consider either removing them or merging them, e.g., with `scater::sumCountsAcrossFeatures`.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained. All searching is performed after removing the `_loc[2-9]` suffix.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/marques-brain`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Marques A et al. (2016). Oligodendrocyte heterogeneity in the mouse juvenile and adult central nervous system. *Science* 352(6291), 1326-1329.

**Examples**

```
sce <- MarquesBrainData()
```

---

MessmerESCData

*Obtain the Messmer ESC data*

---

**Description**

Obtain the human embryonic stem cell single-cell RNA-seq data from Messmer et al. (2019).

**Usage**

```
MessmerESCData(location = TRUE, legacy = FALSE)
```

**Arguments**

location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Row data contains a single "Length" field describing the total exonic length of each feature.

Column metadata is provided in the same form as supplied in E-MTAB-6819. This contains information such as the cell phenotype (naive or primed) and the batch of origin. Note that counts for technical replicates have already been summed together.

Count data for ERCC spike-ins are stored in the "ERCC" entry of the [altExps](#).

If location=TRUE, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for scRNAseq/messmer-esc.

**Value**

A [SingleCellExperiment](#) object with a single matrix of read counts.

**Author(s)**

Aaron Lun

**References**

Messmer T et al. (2019). Transcriptional heterogeneity in naive and primed human pluripotent stem cells at single-cell resolution. *Cell Rep* 26(4), 815-824.e4

**Examples**

```
sce <- MessmerESData()
```

---

MuraroPancreasData	<i>Obtain the Muraro pancreas data</i>
--------------------	--

---

**Description**

Obtain the human pancreas single-cell RNA-seq data from Muraro et al. (2016).

**Usage**

```
MuraroPancreasData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

**Arguments**

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Row data contains fields for the symbol and chromosomal location of each gene.

Column metadata is derived from the columns of the count matrix provided in GSE85241, with additional cell type labels obtained from the authors (indirectly, via the Hemberg group). Some cells have NA labels and were presumably removed prior to downstream analyses.

Count data for ERCC spike-ins are stored in the "ERCC" entry of the [altExps](#).

If ensembl=TRUE, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/muraro-pancreas`.

### Value

A `SingleCellExperiment` object with a single matrix of UMI counts.

### Author(s)

Aaron Lun, using additional metadata obtained by Vladimir Kiselev.

### References

Muraro MJ et al. (2016). A single-cell transcriptome atlas of the human pancreas. *Cell Syst.* 3(4), 385-394.

### Examples

```
sce <- MuraroPancreasData()
```

---

NestorowaHSCData	<i>Obtain the Nestorowa HSC data</i>
------------------	--------------------------------------

---

### Description

Obtain the mouse haematopoietic stem cell single-cell RNA-seq data from Nestorowa et al. (2015).

### Usage

```
NestorowaHSCData(remove.htseq = TRUE, location = TRUE, legacy = FALSE)
```

### Arguments

<code>remove.htseq</code>	Logical scalar indicating whether HT-seq alignment statistics should be removed.
<code>location</code>	Logical scalar indicating whether genomic coordinates should be returned.
<code>legacy</code>	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Rows corresponding to HT-seq's alignment statistics are removed by default. These can be retained by setting `remove.htseq=FALSE`.

Column metadata includes the cell type mapping, as described on the website (see References), and the FACS expression levels of selected markers. Note that these are stored as nested matrices within the `colData`.

Diffusion map components are provided as the "diffusion" entry in the `reducedDims`.

Counts for ERCC spike-ins are stored in the "ERCC" entry in the `altExps`.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/nestorowa-hsc`.

## Value

A `SingleCellExperiment` object with a single matrix of read counts.

## Author(s)

Aaron Lun

## References

Nestorowa S et al. (2016). A single-cell resolution map of mouse hematopoietic stem and progenitor cell differentiation *Blood* 128, e20-e31.

Gene and protein expression in adult haematopoiesis: Data. [http://blood.stemcells.cam.ac.uk/single\\_cell\\_atlas.html#data](http://blood.stemcells.cam.ac.uk/single_cell_atlas.html#data).

## Examples

```
sce <- NestorowaHSCData()
```

---

NowakowskiCortexData    *Obtain the Nowakowski cortex data*

---

## Description

Obtain the human cortex single-cell RNA-seq dataset from Nowakowski et al. (2017).

## Usage

```
NowakowskiCortexData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

**Arguments**

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata includes the presumed cell type (WGCNAcluster), patient and tissue region of origin. A variety of dimensionality reduction results are also provided.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. This is only performed when `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/nowakowski-cortex`.

**Value**

A `SingleCellExperiment` object with a single matrix of TPMs. The `reducedDims` contains an assortment of dimensionality reduction results.

**Author(s)**

Aaron Lun

**References**

Nowakowski S et al. (2017). Spatiotemporal gene expression trajectories reveal developmental hierarchies of the human cortex. *Science* 358, 1318-1323.

**Examples**

```
sce <- NowakowskiCortexData()
```



---

PaulHSCData*Obtain the Paul HSC data*

---

**Description**

Obtain the mouse haematopoietic stem cell single-cell RNA-seq data from Paul et al. (2015).

**Usage**

```
PaulHSCData(  
  ensembl = FALSE,  
  discard.multiple = TRUE,  
  location = TRUE,  
  legacy = FALSE  
)
```

**Arguments**

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
discard.multiple	Logical scalar indicating whether ambiguous rows should be discarded.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata includes the plate and the mouse of origin, fluorescence intensities from indexed sorting and the number of cells in each well.

Some of the original rownames are concatenated symbols from multiple genes. We consider these rows to represent ambiguously assigned counts and discard them if `discard.multiple=TRUE`.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/nestorowa-hsc`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of read counts.

**Author(s)**

Aaron Lun

**References**

Paul F et al. (2015). Transcriptional heterogeneity and lineage commitment in myeloid progenitors. *Cell* 163, 1663-77.

**Examples**

```
sce <- PaulHSCData()
```

---

polishDataset

*Polish dataset for saving*


---

**Description**

Prepare a [SummarizedExperiment](#) or [SingleCellExperiment](#) to be saved with [saveDataset](#). This performs some minor changes to improve storage efficiency.

**Usage**

```
polishDataset(
  x,
  strip.inner.names = TRUE,
  reformat.assay.by.density = 0.3,
  attempt.integer.conversion = TRUE,
  remove.altexp.coldata = TRUE,
  forbid.nested.altexp = TRUE
)
```

**Arguments**

**x** A [SummarizedExperiment](#) or one of its subclasses.

**strip.inner.names** Logical scalar indicating whether to strip redundant names from internal objects, e.g., dimnames of the assays, row names of reduced dimensions, column names of alternative experiments. This saves some space in the on-disk representation.

**reformat.assay.by.density** Numeric scalar indicating whether to optimize assay formats based on the density of non-zero values. Assays with densities above this number are converted to ordinary dense arrays (if they are not already), while those with lower densities are converted to sparse matrices. This can be disabled by setting it to NULL.

`attempt.integer.conversion`  
 Logical scalar indicating whether to convert double-precision assays containing integer values to actually have the integer type. This can improve efficiency of downstream applications by avoiding the need to operate in double precision.

`remove.altexp.coldata`  
 Logical scalar indicating whether column data for alternative experiments should be removed. This defaults to TRUE as the alternative experiment column data is usually redundant with that of the main experiment.

`forbid.nested.altexp`  
 Logical scalar indicating whether nested alternative experiments (i.e., alternative experiments of alternative experiments) should be forbidden. This defaults to TRUE as nested alternative experiments are usually the result of some mistake in `altExp` preparation.

**Value**

A modified copy of `x`.

**Author(s)**

Aaron Lun

**Examples**

```
mat <- matrix(rpois(1000, lambda=0.2), 100, 10) * 1.0
rownames(mat) <- sprintf("GENE_%i", seq_len(nrow(mat)))
colnames(mat) <- head(LETTERS, 10)

library(SingleCellExperiment)
sce <- SingleCellExperiment(list(counts=mat))
str(assay(sce, withDimnames=FALSE))

polished <- polishDataset(sce)
str(assay(polished, withDimnames=FALSE))
```

---

PollenGliaData

---

*Obtain the Pollen radial glia data*


---

**Description**

Obtain the human radial glia single-cell RNA-seq dataset from Pollen et al. (2017).

**Usage**

```
PollenGliaData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

**Arguments**

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata includes the anatomical source, sample of origin, presumed cell type and assorted alignment statistics.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. This is only performed when `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/pollen-glia`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of read counts.

**Author(s)**

Aaron Lun

**References**

Pollen A et al. (2017). Molecular identity of human outer radial glia during cortical development. *Cell* 163, 55-67.

**Examples**

```
sce <- PollenGliaData()
```

---

reexports

*Objects exported from other packages*


---

**Description**

These objects are imported from other packages. Follow the links below to see their documentation.

**gypsum** [defineTextQuery](#), [gsc](#)

---

ReprocessedAllenData    *Reprocessed single-cell data sets*

---

## Description

Obtain the legacy count matrices for three publicly available single-cell RNA-seq datasets. Raw sequencing data were downloaded from NCBI's SRA or from EBI's ArrayExpress, aligned to the relevant genome build and used to quantify gene expression.

## Usage

```
ReprocessedAllenData(  
  assays = NULL,  
  ensembl = FALSE,  
  location = TRUE,  
  legacy = FALSE  
)
```

```
ReprocessedTh2Data(  
  assays = NULL,  
  ensembl = FALSE,  
  location = TRUE,  
  legacy = FALSE  
)
```

```
ReprocessedFluidigmData(  
  assays = NULL,  
  ensembl = FALSE,  
  location = TRUE,  
  legacy = FALSE  
)
```

## Arguments

assays	Character vector specifying one or more assays to return. Choices are "tophat_counts", "cufflinks_fpkms", "rsem_counts" and "rsem_tpm". If NULL, all assays are returned.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

ReprocessedFluidigmData returns a dataset of 65 human neural cells from Pollen et al. (2014), each sequenced at high and low coverage (SRA accession SRP041736).

ReprocessedTh2Data returns a dataset of 96 mouse T helper cells from Mahata et al. (2014), obtained from ArrayExpress accession E-MTAB-2512. Spike-in counts are stored in the "ERCC" entry of the [altExps](#).

ReprocessedAllenData return a dataset of 379 mouse brain cells from Tasic et al. (2016). This is a re-processed subset of the data from [TasicBrainData](#), and contains spike-in information stored as in the [altExps](#).

In each dataset, the first columns of the `colData` are sample quality metrics from FastQC and Picard. The remaining fields were obtained from the original study in their GEO/SRA submission and/or as Supplementary files in the associated publication. These two categories of `colData` are distinguished by a `which_qc` element in the [metadata](#), which contains the names of the quality-related columns in each object.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/legacy-allen`, `scRNAseq/legacy-fluidigm` or `scRNAseq/legacy-th2`.

## Value

A [SingleCellExperiment](#) object containing one or more expression matrices of counts and/or TPMs, depending on assays.

## Pre-processing details

FASTQ files were either obtained directly from ArrayExpress, or converted from SRA files (downloaded from the Sequence Read Archive) using the SRA Toolkit.

Reads were aligned with TopHat (v. 2.0.11) to the appropriate reference genome (GRCh38 for human samples, GRCm38 for mouse). RefSeq mouse gene annotation (GCF\_000001635.23\_GRCm38.p3) was downloaded from NCBI on Dec. 28, 2014. RefSeq human gene annotation (GCF\_000001405.28) was downloaded from NCBI on Jun. 22, 2015.

`featureCounts` (v. 1.4.6-p3) was used to compute gene-level read counts. `Cufflinks` (v. 2.2.0) was used to compute gene-level FPKMs. Reads were also mapped to the transcriptome using `RSEM` (v. 1.2.19) to compute read counts and TPM's.

`FastQC` (v. 0.10.1) and `Picard` (v. 1.128) were used to compute sample quality control (QC) metrics. However, no filtering on the QC metrics has been performed for any dataset.

## References

Pollen AA et al. (2014). Low-coverage single-cell mRNA sequencing reveals cellular heterogeneity and activated signaling pathways in developing cerebral cortex. *Nat. Biotechnol.* 32(10), 1053-8.

Mahata B et al. (2014). Single-cell RNA sequencing reveals T helper cells synthesizing steroids de novo to contribute to immune homeostasis. *Cell Rep*, 7(4), 1130-42.

Tasic A et al. (2016). Adult mouse cortical cell taxonomy revealed by single cell transcriptomics. *Nat. Neurosci.* 19(2), 335-46.

## Examples

```
sce <- ReprocessedAllenData()
```

---

RichardTCellData	<i>Obtain the Richard T cell data</i>
------------------	---------------------------------------

---

## Description

Obtain the mouse CD8+ T cell single-cell RNA-seq data from Richard et al. (2018).

## Usage

```
RichardTCellData(location = TRUE, legacy = FALSE)
```

## Arguments

location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column metadata is provided in the same form as supplied in E-MTAB-6051. This contains information such as the stimulus, time after stimulation, age of the mice and sequencing batch.

Count data for ERCC spike-ins are stored in the "ERCC" entry of the [altExps](#).

If location=TRUE, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/richard-tcell`.

## Value

A [SingleCellExperiment](#) object with a single matrix of read counts.

## Author(s)

Aaron Lun

## References

Richard AC et al. (2018). T cell cytolytic capacity is independent of initial stimulation strength. *Nat. Immunol.* 19(8), 849-858.

## Examples

```
sce <- RichardTCellData()
```

---

RomanovBrainData	<i>Obtain the Romanov brain data</i>
------------------	--------------------------------------

---

## Description

Obtain the mouse brain single-cell RNA-seq dataset from Romanov et al. (2017).

## Usage

```
RomanovBrainData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

## Arguments

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column metadata is provided in the same form as supplied in GSE74672. This contains information such as the reporter gene expressed in each cell, the mouse line, dissection type and so on.

Counts for ERCC spike-ins are stored in the "ERCC" entry of the [altExps](#). Note that some of the spike-in rows have NA observations for some (but not all) cells.

If ensembl=TRUE, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If location=TRUE, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output. Note that this is only performed if ensembl=TRUE.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for scrnaseq/romanov-brain.

## Value

A [SingleCellExperiment](#) object with a single matrix of UMI counts.



**Author(s)**

Aaron Lun, based on code by Vladimir Kiselev and Tallulah Andrews.

**References**

Romanov RA et al. (2017). Molecular interrogation of hypothalamic organization reveals distinct dopamine neuronal subtypes. *Nat. Neurosci.* 20, 176-188.

**Examples**

```
sce <- RomanovBrainData()
```

---

saveDataset	<i>Save a dataset to disk</i>
-------------	-------------------------------

---

**Description**

Save a single-cell dataset to disk, usually in preparation for upload.

**Usage**

```
saveDataset(x, path, metadata)
```

**Arguments**

x	A <a href="#">SummarizedExperiment</a> or one of its subclasses.
path	String containing the path to a new directory in which to save x. Any existing directory is removed before saving x.
metadata	Named list containing metadata for this dataset, see the schema returned by <a href="#">fetchMetadataSchema()</a> . Note that the <code>applications.takane</code> property will be automatically added by this function and does not have to be supplied.

**Value**

x and its metadata are saved into path, and NULL is invisibly returned.

**Author(s)**

Aaron Lun

**See Also**

<https://github.com/ArtifactDB/bioconductor-metadata-index>, on the expected schema for the metadata.

[polishDataset](#), to polish x before saving it.

[uploadDirectory](#), to upload the saved contents.

## Examples

```
library(SingleCellExperiment)
sce <- SingleCellExperiment(list(counts=matrix(rpois(1000, lambda=1), 100, 10)))
rownames(sce) <- sprintf("GENE_%i", seq_len(nrow(sce)))
colnames(sce) <- head(LETTERS, 10)

meta <- list(
  title="My dataset",
  description="This is my dataset",
  taxonomy_id="10090",
  genome="GRCh38",
  sources=list(list(provider="GEO", id="GSE12345")),
  maintainer_name="Shizuka Mogami",
  maintainer_email="mogami.shizuka@765pro.com"
)

tmp <- tempfile()
saveDataset(sce, tmp, meta)
list.files(tmp, recursive=TRUE)
alabaster.base::readObject(tmp)
```

---

searchDatasets	<i>Search dataset metadata</i>
----------------	--------------------------------

---

## Description

Search for datasets of interest based on matching text in the associated metadata.

## Usage

```
searchDatasets(
  query,
  cache = cacheDirectory(),
  overwrite = FALSE,
  latest = TRUE
)
```

## Arguments

query	String containing a query in a human-readable syntax or a <a href="#">gypsum.search.clause</a> , see Examples.
cache, overwrite	Arguments to pass to <a href="#">fetchMetadataDatabase</a> .
latest	Whether to only consider the latest version of each dataset.

## Details

The returned DataFrame contains the usual suspects like the title and description for each dataset, the number of rows and columns, the organisms and genome builds involved, whether the dataset has any pre-computed reduced dimensions, and so on. More details can be found in the Bioconductor metadata schema at <https://github.com/ArtifactDB/bioconductor-metadata-index>.

If a dataset contains multiple subdatasets, each subdataset is reported as a separate row in the DataFrame. The location of subdataset is provided in the path column. If a dataset does not contain any subdatasets, the path entry will be set to NA.

## Value

A [DataFrame](#) where each row corresponds to a (sub)dataset, containing various columns of meta-data. Some columns may be lists to capture 1:many mappings.

## Author(s)

Aaron Lun

## See Also

[surveyDatasets](#), to easily obtain a listing of all available datasets.

[translateTextQuery](#), for details on the human-readable query syntax.

## Examples

```
searchDatasets("brain")[,c("name", "title")]
searchDatasets("Neuro%")[,c("name", "title")]
searchDatasets("taxonomy_id:10090")[,c("name", "title")]
searchDatasets("(genome: GRCm38 AND neuro%) OR pancrea%")[,c("name", "title")]

# We can also use gypsum search clauses via the gsc() function.
# Here, 'asset' is analogous to the 'name' of the dataset.
searchDatasets(gsc(asset="he-organs-2020"))[,c("path")]
searchDatasets(
  (gsc(asset="%brain%", partial=TRUE) |
   gsc(asset="%neur%", partial=TRUE) |
   gsc(text="%neur%", partial=TRUE) |
   gsc(text="%brain%", partial=TRUE)) &
  gsc(field="taxonomy_id", text="10090"))
)[,c("name", "title")]
```

---

SegerstolpePancreasData

*Obtain the Segerstolpe pancreas data*


---

## Description

Download the human pancreas single-cell RNA-seq (scRNA-seq) dataset from Segerstolpe et al. (2016)

## Usage

```
SegerstolpePancreasData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

## Arguments

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Row data contains fields for the gene symbol and RefSeq transcript IDs corresponding to each gene. The rows of the output object are named with the symbol, but note that these are not unique.

Column metadata were extracted from the Characteristics fields of the SDRF file for ArrayExpress E-MTAB-5061. This contains information such as the cell type labels and patient status.

Count data for ERCC spike-ins are stored in the "ERCC" entry of the [altExps](#). Estimated numbers of spike-in molecules are provided in the [rowData](#) of this entry. Note that these concentrations are incorrect for donor H1, as 100 uL of spike-in mixture were added for this donor, rather than 25 uL for all others.

If ensembl=TRUE, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If location=TRUE, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output. Note that this is only performed if ensembl=TRUE.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/segerstolpe-pancreas`.

## Value

A [SingleCellExperiment](#) object with a single matrix of read counts.

## Author(s)

Aaron Lun

## References

Segerstolpe A et al. (2016). Single-cell transcriptome profiling of human pancreatic islets in health and type 2 diabetes. *Cell Metab.* 24(4), 593-607.

## Examples

```
sce <- SegerstolpePancreasData()
```

---

ShekharRetinaData	<i>Obtain the Shekhar retina data</i>
-------------------	---------------------------------------

---

## Description

Obtain the mouse retina single-cell RNA-seq dataset from Shekhar et al. (2016).

## Usage

```
ShekharRetinaData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

## Arguments

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column metadata contains the cluster identities as reported in the paper. Note that some cells will have NA identities as they are present in the count matrix but not in the metadata file. These are presumably low-quality cells that were discarded prior to clustering.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/shekhar-retina`.

## Value

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Shekhar K et al. (2016). Comprehensive classification of retinal bipolar neurons by single-cell transcriptomics. *Cell* 166(5), 1308-1323.

**Examples**

```
sce <- ShekharRetinaData()
```

---

StoeckiusHashingData    *Obtain the Stoeckius cell hashing data*

---

**Description**

Obtain the (mostly human) cell hashing single-cell RNA-seq data from Stoeckius et al. (2018).

**Usage**

```
StoeckiusHashingData(
  type = c("pbmc", "mixed"),
  mode = NULL,
  ensembl = FALSE,
  location = TRUE,
  strip.metrics = TRUE,
  legacy = FALSE
)
```

**Arguments**

type	String specifying the dataset to obtain.
mode	String specifying the data modalities to obtain, see Details.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
strip.metrics	Logical scalar indicating whether quality control metrics should be removed from the HTO/ADT counts.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

When `type="pbmc"`, the mode can be one or more of:

- "human", the RNA counts for human genes.
- "mouse", the RNA counts for mouse genes. Present as the PBMC dataset is actually a mixture of human PBMCs and unlabelled mouse cells.
- "hto", the HTO counts.
- "adt1", counts for the first set of ADTs (immunoglobulin controls).
- "adt2", counts for the second set of ADTs (cell type-specific markers).

If `mode=NULL`, the default is to use "human", "mouse" and "hto".

When `type="mixed"`, the mode can be one or more of:

- "rna", the RNA counts for the genes;
- "hto", the HTO counts.

If `mode=NULL`, the default is to use "rna" and "hto".

If `ensembl=TRUE`, gene symbols for the RNA counts are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE` and only for the RNA counts.

For the HTO and ADT matrices, some rows correspond to quality control metrics. If `strip.metrics=TRUE`, these rows are removed so that only data for actual HTOs or ADTs are present.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scrnaseq/nestorowa-hsc`.

## Value

A [SingleCellExperiment](#) object with a matrix of UMI counts corresponding to the first mode, plus any number of alternative Experiments containing the remaining modes. If multiple modes are specified, the output object only contains the intersection of their column names.

## Author(s)

Aaron Lun

## References

Stoeckius et al. (2018). Cell Hashing with barcoded antibodies enables multiplexing and doublet detection for single cell genomics. *Genome Biol.* 19, 224.

**Examples**

```
sce.pbmc <- StoeckiusHashingData()
sce.pbmc

sce.mixed <- StoeckiusHashingData(type="mixed")
sce.mixed
```

---

surveyDatasets	<i>Survey of dataset metadata</i>
----------------	-----------------------------------

---

**Description**

Metadata survey for all available datasets in the **scRNAseq** package.

**Usage**

```
surveyDatasets(cache = cacheDirectory(), overwrite = FALSE, latest = TRUE)
```

**Arguments**

cache, overwrite	Arguments to pass to <a href="#">fetchMetadataDatabase</a> .
latest	Whether to only consider the latest version of each dataset.

**Details**

The returned `DataFrame` contains the usual suspects like the title and description for each dataset, the number of rows and columns, the organisms and genome builds involved, whether the dataset has any pre-computed reduced dimensions, and so on. More details can be found in the Bioconductor metadata schema at <https://github.com/ArtifactDB/bioconductor-metadata-index>.

If a dataset contains multiple subdatasets, each subdataset is reported as a separate row in the `DataFrame`. The location of subdataset is provided in the `path` column. If a dataset does not contain any subdatasets, the `path` entry will be set to `NA`.

**Value**

A [DataFrame](#) where each row corresponds to a (sub)dataset, containing various columns of metadata. Some columns may be lists to capture 1:many mappings.

**Author(s)**

Aaron Lun

**See Also**

[searchDatasets](#), to search on the metadata for specific datasets.



**Examples**

```
surveyDatasets()
```

---

TasicBrainData	<i>Obtain the Tasic brain data</i>
----------------	------------------------------------

---

**Description**

Obtain the mouse brain single-cell RNA-seq data from Tasic et al. (2015).

**Usage**

```
TasicBrainData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

**Arguments**

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata is provided in the same form as supplied in GSE71585. This contains information such as the reporter gene expressed in each cell, the mouse line, dissection type and so on.

Count data for ERCC spike-ins are stored in the "ERCC" entry of the [altExps](#). Note that some of the spike-in rows have NA observations for some (but not all) cells.

The last 9 columns (containing \_CTX\_ in their names) correspond to no-cell control libraries.

If ensembl=TRUE, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If location=TRUE, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output. Note that this is only performed if ensembl=TRUE.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for scrNAseq/tasic-brain.

**Value**

A [SingleCellExperiment](#) object with a single matrix of read counts.

**Author(s)**

Aaron Lun

## References

Tasic A et al. (2016). Adult mouse cortical cell taxonomy revealed by single cell transcriptomics. *Nat. Neurosci.* 19(2), 335-46.

## Examples

```
sce <- TasicBrainData()
```

---

UsoskinBrainData	<i>Obtain the Usoskin brain data</i>
------------------	--------------------------------------

---

## Description

Obtain the mouse brain single-cell RNA-seq dataset from Usoskin et al. (2015).

## Usage

```
UsoskinBrainData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

## Arguments

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column metadata is provided in the same form as supplied in External Table 2 of <http://linnarssonlab.org/drg/>. This contains information such as the library of origin and the cell type.

The count matrix contains information for repeats, marked with `r_` prefixes in the row names; as well as mitochondrial transcripts, marked with `mt-` prefixes.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/usoskin-brain`.

## Value

A [SingleCellExperiment](#) object with a single matrix of RPMs.

**Author(s)**

Aaron Lun

**References**

Usoskin A et al. (2015). Unbiased classification of sensory neuron types by large-scale single-cell RNA sequencing. *Nat. Neurosci.* 18(1), 145-53.

**Examples**

```
sce <- UsoskinBrainData()
```

---

WuKidneyData

---

*Obtain the Wu kidney data*


---

**Description**

Obtain the mouse kidney single-nuclei RNA-seq data from Wu et al. (2019).

**Usage**

```
WuKidneyData(
  mode = c("healthy", "disease"),
  ensembl = FALSE,
  location = TRUE,
  legacy = FALSE
)
```

**Arguments**

mode	String indicating whether to return data for healthy and/or diseased donors.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata includes the single-cell technology and whether they came from a diseased or healthy individual.

If mode specifies both healthy and disease donors, counts are only reported for the intersection of genes that are present for both donors. This is because the original count matrices had differences in their annotation.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/wu-kidney`.

### Value

A `SingleCellExperiment` object with a single matrix of read counts.

### Author(s)

Aaron Lun

### References

Wu H et al. (2019). Advantages of single-nucleus over single-cell RNA sequencing of adult kidney: rare cell types and novel cell states revealed in fibrosis. *J. Am. Soc. Nephrol.* 30, 23-32.

### Examples

```
sce <- WuKidneyData("disease")
```

---

XinPancreasData	<i>Obtain the Xin pancreas data</i>
-----------------	-------------------------------------

---

### Description

Obtain the human pancreas single-cell RNA-seq dataset from Xin et al. (2016).

### Usage

```
XinPancreasData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

### Arguments

<code>ensembl</code>	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
<code>location</code>	Logical scalar indicating whether genomic coordinates should be returned.
<code>legacy</code>	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Row data contains fields for the Entrez ID and symbol for each gene. Column metadata was obtained from the authors (indirectly, via the Hemberg group) and contains information such as the cell type labels and donor status.

If `ensembl=TRUE`, the Entrez IDs are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/xin-pancreas`.

**Value**

A `SingleCellExperiment` object with a single matrix of RPKMs.

**Author(s)**

Aaron Lun, using additional metadata obtained by Vladimir Kiselev.

**References**

Xin A et al. (2016). RNA sequencing of single human islet cells reveals type 2 diabetes genes. *Cell Metab.* 24(4), 608-615.

**Examples**

```
sce <- XinPancreasData()
```

---

ZeiselBrainData	<i>Obtain the Zeisel brain data</i>
-----------------	-------------------------------------

---

**Description**

Obtain the mouse brain single-cell RNA-seq dataset from Zeisel et al. (2015).

**Usage**

```
ZeiselBrainData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

**Arguments**

<code>ensembl</code>	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
<code>location</code>	Logical scalar indicating whether genomic coordinates should be returned.
<code>legacy</code>	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Row data contains a single "featureType" field describing the type of each feature (endogenous genes, mitochondrial genes, spike-in transcripts and repeats). Spike-ins and repeats are stored as separate entries in the [altExps](#).

Column metadata is provided in the same form as supplied in <http://linnarssonlab.org/cortex/>. This contains information such as the cell diameter and the published cell type annotations.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output.

Spike-in metadata is added using [ERCCSpikeInConcentrations](#), with molecule counts computed using a volume of 9 nL per cell at a dilution of 1:20000.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/zeisel-brain`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Zeisel A et al. (2015). Brain structure. Cell types in the mouse cortex and hippocampus revealed by single-cell RNA-seq. *Science* 347(6226), 1138-42.

**Examples**

```
sce <- ZeiselBrainData()
```

---

ZeiselNervousData	<i>Obtain the Zeisel nervous system data</i>
-------------------	--

---

**Description**

Obtain the mouse nervous system single-cell RNA-seq dataset from Zeisel et al. (2018).

**Usage**

```
ZeiselNervousData(location = TRUE, legacy = FALSE)
```

**Arguments**

location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Row data contains the gene symbol as well as some relevant per-gene statistics, e.g., the squared coefficient of variance, mean, and whether it was selected for downstream analyses.

Column data contains a wide variety of fields including patient-level information, sample-level sequencing statistics and many flavors of cell type classification. Note that many numeric columns may have NA values if they could not be successfully parsed from the source file.

If location=TRUE, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/zeisel-nervous`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Zeisel A et al. (2018). Molecular architecture of the mouse nervous system. *Cell* 174(4), 999-1014.

**Examples**

```
if (.Machine$sizeof.pointer > 4) { # too large for 32-bit machines!
  sce <- ZeiselNervousData()
}
```

---

ZhaoImmuneLiverData	<i>Obtain the Zhao immune liver data</i>
---------------------	--

---

**Description**

Obtain the human liver immune single-cell RNA-seq data from Zhao et al. (2020).

**Usage**

```
ZhaoImmuneLiverData(location = TRUE, filter = FALSE, legacy = FALSE)
```

**Arguments**

location	Logical scalar indicating whether genomic coordinates should be returned.
filter	Logical scalar indicating if the filtered subset should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

**Details**

Column metadata contains various cell labels as provided by the authors. Some of these labels required assembly on our part:

- The broad label was assigned to each barcode based on whether that barcode was present in each \*\_identities.tsv.gz in GSE125188's supplementary files.
- For each cell barcode that was present in one of these files, the fine label was generated from the Group annotations inside that file.

We guessed the sample for each cell by assuming that the GEM group numbers match the order of samples in GSE125188. We also assumed that “donor 4” is a typo, given that the paper only mentions 3 donors.

If location=TRUE, the coordinates of the Ensembl gene models are stored in the [rowRanges](#) of the output. Note that this is only performed if ensembl=TRUE.

If filter=TRUE, only cells that have been used in the original analysis are returned. Otherwise, the cells used are specified in the retained column of the [colData](#).

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/zhao-immune-liver`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

**Author(s)**

Aaron Lun

**References**

Zhao J et al. (2020). Single-cell RNA sequencing reveals the heterogeneity of liver-resident immune cells in human. *Cell Discov* 6, 22.

**Examples**

```
sce.zhao <- ZhaoImmuneLiverData()
```



---

ZhongPrefrontalData	Obtain the Zhong prefrontal cortex data
---------------------	---

---

## Description

Obtain the human prefrontal cortex single-cell RNA-seq dataset from Zhong et al. (2018).

## Usage

```
ZhongPrefrontalData(ensembl = FALSE, location = TRUE, legacy = FALSE)
```

## Arguments

ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column metadata is scraped from GEO and includes week of gestation, gender and likely cell type. If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. This is only performed when `ensembl=TRUE`.

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scrNaseq/zhong-prefrontal`.

## Value

A [SingleCellExperiment](#) object with a single matrix of UMI counts.

## Author(s)

Aaron Lun

## References

Zhong S et al. (2018). A single-cell RNA-seq survey of the developmental landscape of the human prefrontal cortex. *Nature* 555, 524-528.

## Examples

```
sce <- ZhongPrefrontalData()
```

---

ZilionisLungData

---

*Obtain the Zilionis lung cancer data*


---

## Description

Obtain the human/mouse lung cancer single-cell RNA-seq data from Zilionis et al. (2019).

## Usage

```
ZilionisLungData(
  which = c("human", "mouse"),
  ensembl = FALSE,
  location = TRUE,
  filter = FALSE,
  legacy = FALSE
)
```

## Arguments

which	String specifying the species to get data for.
ensembl	Logical scalar indicating whether the output row names should contain Ensembl identifiers.
location	Logical scalar indicating whether genomic coordinates should be returned.
filter	Logical scalar indicating if the filtered subset should be returned.
legacy	Logical scalar indicating whether to pull data from ExperimentHub. By default, we use data from the gypsum backend.

## Details

Column metadata is provided and contains information on the library, donor ID/animal ID, replicate and tissue.

If `ensembl=TRUE`, the gene symbols are converted to Ensembl IDs in the row names of the output object. Rows with missing Ensembl IDs are discarded, and only the first occurrence of duplicated IDs is retained.

If `location=TRUE`, the coordinates of the Ensembl gene models are stored in the `rowRanges` of the output. Note that this is only performed if `ensembl=TRUE`.

If `filter=TRUE`, only cells that have been used in the original analysis are returned. The cells used are specified in the `Used` column of the `colData`.

The `reducedDim` contains coordinates of SPRING representations. This may be filled with NAs for SPRING coordinates computed on a subset of cells (specified in `colData`).

All data are downloaded from ExperimentHub and cached for local re-use. Specific resources can be retrieved by searching for `scRNAseq/zilionis-lung`.

**Value**

A [SingleCellExperiment](#) object with a single matrix of read counts.

**Author(s)**

Jens Preussner

**References**

Zilionis R et al. (2019). Single-cell transcriptomics of human and mouse lung cancers reveals conserved myeloid populations across individuals and species. *Immunity* 50(5), 1317-1334.

**Examples**

```
sce.human <- ZilionisLungData()

sce.mouse <- ZilionisLungData("mouse")
```

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