Package ‘SpatialFeatureExperiment’

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Type Package

Title Integrating SpatialExperiment with Simple Features in sf

Version 1.11.0

Description A new S4 class integrating Simple Features with the R package sf to bring geospatial data analysis methods based on vector data to spatial transcriptomics. Also implements management of spatial neighborhood graphs and geometric operations. This pakage builds upon SpatialExperiment and SingleCellExperiment, hence methods for these parent classes can still be used.

Imports Biobase, BiocGenerics (>= 0.51.2), BiocNeighbors, BiocParallel, data.table, DropletUtils, EBImage, grDevices, lifecycle, Matrix, methods, rjson, rlang, S4Vectors, sf, sfheaders, SingleCellExperiment, SpatialExperiment, spatialreg, spdep (>= 1.1-7), SummarizedExperiment, stats, terra, utils, zeallot

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addVisiumSpotPoly — *Add Visium spot polygons to colGeometry*

Description

For adding the spot polygons to SFE objects converted from SPE.

Usage

```
addVisiumSpotPoly(x, spotDiameter)
```

Arguments

x	A SpatialFeatureExperiment object.
spotDiameter	Spot diameter for technologies with arrays of spots of fixed diameter per slide, such as Visium, ST, DBiT-seq, and slide-seq. The diameter must be in the same unit as the coordinates in the *Geometry arguments. Ignored for geometries that are not POINT or MULTIPOINT.

Value

A SFE object with a new colGeometry called spotPoly, which has polygons of the spots.

Examples

```
library(SpatialExperiment)
example(read10xVisium)
# There can't be suplicate barcodes
colnames(spe) <- make.unique(colnames(spe), sep = "-")
rownames(spatialCoords(spe)) <- colnames(spe)
sfe <- toSpatialFeatureExperiment(spe)
# A hypothetical spot diameter; check the scalefactors_json.json file for
# actual diameter in pixels in full resolution image.
sfe <- addVisiumSpotPoly(sfe, spotDiameter = 80)
```

affineImg — *Affine transformation of images*

Description

This function performs affine transformation on images, with any matrix and translation vector.

Usage

```
## S4 method for signature 'SpatRasterImage'
affineImg(x, M, v, maxcell = 1e+07, ...)

## S4 method for signature 'BioFormatsImage'
affineImg(x, M, v, ...)

## S4 method for signature 'ExtImage'
affineImg(x, M, v, ...)
```

Arguments

x	An object of class *Image as implemented in this package.
M	A 2x2 numeric matrix for the linear transformation in the xy plane.
v	A numeric vector of length 2 for translation in the xy plane.
maxcell	Max number of pixels to load SpatRasterImage into memory. The default 1e7 is chosen because this is the approximate number of pixels in the medium resolution image at resolution = 4L in Xenium OME-TIFF to make different methods of this function consistent.
...	Ignored. It's there so different methods can all be passed to the same lapply in the method for SFE objects. Some methods have extra arguments.

Value

SpatRasterImage will be converted to ExtImage. Otherwise *Image object of the same class. For BioFormatsImage, the transformation info is stored and will be applied when the image is loaded into memory as ExtImage.

See Also

Other image methods: SFE-image, cropImg(), dim,BioFormatsImage-method, dim,ExtImage-method, ext(), imgRaster(), imgSource(), mirrorImg(), rotateImg(), scaleImg(), translateImg(), transposeImg()

---

aggBboxes *Aggregate bounding boxes*

Description

To find the bounding box of multiple bounding boxes.

Usage

```
aggBboxes(bboxes)
```

Arguments

bboxes	Either a matrix with 4 rows whose columns are the different bounding boxes, with row names "xmin", "xmax", "ymin", and "ymax" in any order, or a list of bounding boxes which are named numeric vectors.

Value

A named numeric vector for the total bounding box.

Examples

```
bboxes <- list(c(xmin = 5, xmax = 10, ymin = 2, ymax = 20),
c(xmin = 8, xmax = 18, ymin = 0, ymax = 15))
bbox_all <- aggBboxes(bboxes)
```

aggregate,SpatialFeatureExperiment-method

Aggregate data in SFE using geometry

Description

Gene expression and numeric columns of `colData` will be aggregated with the function specified in `FUN`, according to another geometry supplied and a geometry predicate (such as `st_intersects`). For example, when the predicate is `st_intersects` and a spatial grid is used to aggregate, then the data associated with all cells that intersect with each grid cell will be aggregated with `FUN`, such as `mean` or `sum`. The categorical columns will be collected into list columns, and logical columns will be converted into numeric before applying `FUN`.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
aggregate(
  x,
  by = NULL,
  FUN = sum,
  sample_id = "all",
  colGeometryName = 1L,
  rowGeometryName = NULL,
  cellsize = NULL,
  square = TRUE,
  flat_topped = FALSE,
  new_geometry_name = "bins",
  join = st_intersects,
  sparse = FALSE,
  BPPARAM = SerialParam()
)
```

Arguments

`x`	An SFE object to be aggregated.
`by`	A `sf` data frame whose geometry column is used for aggregation or `sfc` or for multiple samples a list of `sfc` whose names are the sample IDs. For multiple samples, the `sf` data frame must have a column `sample_id` to indicate which geometry for which sample. This argument is optional if `cellsize` is specified.
`FUN`	Function to aggregate the numerical columns in `colData` and the gene count matrix. This can be `sum`, `mean`, or any function that takes a numeric matrix as input and returns a numeric vector whose length is same as the number of rows in the input matrix, such as `rowMedians`. See package `matrixStats`. Depending on the function used for aggregation, numeric columns of `colData` may need to be interpreted differently after aggregation. Aggregation is not done when aggregating by transcript spots in `rowGeometry`. When it's sum or mean, matrix multiplication is used for aggregation rather than calling the sum or mean function itself; this is much faster than looping through the bins and calling the function on each of them.
`sample_id`	Which samples to aggregate, defaults to "all".

colGeometryName
: Which colGeometry to spatially aggregate the data, by default the first one.

rowGeometryName
: Which rowGeometry to spatially aggregate

cellsize
: numeric of length 1 or 2 with target cellsize: for square or rectangular cells the width and height, for hexagonal cells the distance between opposite edges (edge length is cellsize/sqrt(3)). A length units object can be passed, or an area unit object with area size of the square or hexagonal cell.

square
: logical; if FALSE, create hexagonal grid

flat_topped
: logical; if TRUE generate flat topped hexagons, else generate pointy topped

new_geometry_name
: Name to give to the new colGeometry in the output. Defaults to "bins".

join
: logical spatial predicate function to use if by is a simple features object or geometry; see st_join

sparse
: Logical, whether the gene count matrix from aggregating transcript spots should be sparse. When the bins are large, the matrix will not be very sparse so using sparse matrix will not save memory, but when the bins are small, sparsity is worth it.

BPPARAM
: A BiocParallelParam object specifying parallel computing when aggregating data with functions other than sum and mean when aggregating cells. When aggregating transcript spots, this specifies parallel computing over genes. Defaults to SerialParam().

Details

For smFISH-based data where the transcript spots are available, the transcript spots can be used instead of cells to aggregate the gene count matrix, in which case all assays other than counts will be dropped and FUN only applies to colData because the transcript spots are simply counted.

What this function does is similar to SEraster but more general because any geometry and more aggregation function can be used, not just regular grids, and the aggregation can be performed on the transcript spots.

Value

An SFE object with colGeometry the same as the geometry specified in by or same as the grid specified in cellsize. rowGeometries and rowData remain the same as in the input x. reducedDims, localResults, colFeatureData (and its colGeometry, annotGeometry, and reducedDim counterparts), and spatialGraphs are dropped because those results no longer apply after aggregation.

Note

For developers: When debugging this function after calling devtools::load_all("."), you may get an error that comes from S3 dispatch of aggregate.Vector from the S4Vectors package. When that happens, either restart the R session, or run setGeneric("aggregate", function(x, ...) standardGeneric("aggregate")) in the console to make an S4 generic as done in the terra package to prioritize S4 dispatch.

Examples

```
# example code
```

aggregateTx *Aggregate transcript spots from file*

Description

This function reads the transcript spot file from the standard output of the commercial technologies (not GeoParquet) for spatial aggregation where the spots are assigned to polygons such as cells or spatial bins. Presets for Xenium, MERFISH, and CosMX are available. For Vizgen and Xenium, the images can be added when `add_images = TRUE`.

Usage

```
aggregateTx(
  file,
  df = NULL,
  by = NULL,
  sample_id = "sample01",
  spatialCoordsNames = c("X", "Y", "Z"),
  gene_col = "gene",
  phred_col = "qv",
  min_phred = 20,
  flip_geometry = FALSE,
  cellsize = NULL,
  square = TRUE,
  flat_topped = FALSE,
  new_geometry_name = "bins",
  unit = "micron",
  sparse = FALSE,
  BPPARAM = SerialParam(),
  .orig_nrows = NULL
)

aggregateTxTech(
  data_dir,
  df = NULL,
  by = NULL,
  tech = c("Vizgen", "Xenium", "CosMX"),
  sample_id = "sample01",
  image = NULL,
  min_phred = 20,
  flip = c("geometry", "image", "none"),
  max_flip = "50 MB",
  cellsize = NULL,
  square = TRUE,
  flat_topped = FALSE,
  new_geometry_name = "bins",
  sparse = FALSE,
  BPPARAM = SerialParam()
)
```

Arguments

- `file` — File with the transcript spot coordinates. Should be one row per spot when read into R and should have columns for coordinates on each axis, gene the transcript is assigned to, and optionally cell the transcript is assigned to. Must be csv, tsv, or parquet.
- `df` — If the file is already loaded into memory, a data frame (sf) with columns for the x, y, and optionally z coordinates and gene assignment of each transcript spot. If specified, then argument `file` will be ignored.
- `by` — A `sfc` or `sf` object for spatial aggregation.
- `sample_id` — Which sample in the SFE object the transcript spots should be added to.
- `spatialCoordsNames` — Column names for the x, y, and optionally z coordinates of the spots. The defaults are for Vizgen.
- `gene_col` — Column name for genes.
- `phred_col` — Column name for Phred scores of the spots.
- `min_phred` — Minimum Phred score to keep spot. By default 20, the conventional threshold indicating "acceptable", meaning that there's 1 chance that the spot was decoded in error.
- `flip_geometry` — Logical, whether to flip the transcript spot geometries to match the images if added later.
- `cellsize` — numeric of length 1 or 2 with target cellsize: for square or rectangular cells the width and height, for hexagonal cells the distance between opposite edges (edge length is cellsize/sqrt(3)). A length units object can be passed, or an area unit object with area size of the square or hexagonal cell.
- `square` — logical; if `FALSE`, create hexagonal grid
- `flat_topped` — logical; if `TRUE` generate flat topped hexagons, else generate pointy topped
- `new_geometry_name` — Name to give to the new `colGeometry` in the output. Defaults to "bins".
- `unit` — Unit the coordinates are in, either microns or pixels in full resolution image.
- `sparse` — Logical, whether the gene count matrix from aggregating transcript spots should be sparse. When the bins are large, the matrix will not be very sparse so using sparse matrix will not save memory, but when the bins are small, sparsity is worth it.
- `BPPARAM` — bpparam object to specify parallel computing over genes. If a lot of memory is used, then stick to 'SerialParam()'.
- `.orig_nrows` — Only used internally in the SFE method of `aggregate`
- `data_dir` — Top level output directory.
- `tech` — Which technology whose output to read, must be one of "Vizgen", "Xenium", or "CosMX" though more technologies may be added later.
- `image` — String, which image(s) to add to the output SFE object. Not applicable to CosMX. See `readVizgen` and `readXenium` for options and multiple images can be specified. If `NULL`, then the default from the read function for the technology will be used.
- `flip` — Logical, whether to flip the geometry to match image. Here the y coordinates are simply set to -y, so the original bounding box is not preserved. This is consistent with `readVizgen` and `readXenium`.
- `max_flip` — Maximum size of the image allowed to flip the image. Because the image will be loaded into memory to be flipped. If the image is larger than this size then the coordinates will be flipped instead.

Value

A SFE object with count matrix for number of spots of each gene in each geometry. Geometries with no spot are removed.

Note

The resulting SFE object often includes geometries (e.g. grid cells) outside tissue, because there can be transcript spots detected outside the tissue. Also, bins at the edge of the tissue that don't fully overlap with the tissue will have lower transcript counts; this may have implications to downstream spatial analyses.

---

annotGeometries *Annotation geometry methods*

Description

"Annotation geometry" refers to Simple Feature (`sf`) geometries NOT associated with rows (features, genes) or columns (cells or spots) of the gene count matrix in the `SpatialFeatureExperiment` object. So there can be any number of rows in the `sf` data frame specifying the geometry. Examples of such geometries are tissue boundaries, pathologist annotation of histological regions, and objects not characterized by columns of the gene count matrix (e.g. nuclei segmentation in a Visium dataset where the columns are Visium spots). This page documents getters and setters for the annotation geometries. Internally, annotation geometries are stored in `int_metadata`.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
annotGeometries(x)

## S4 replacement method for signature 'SpatialFeatureExperiment'
annotGeometries(x, translate = TRUE, ...) <- value

## S4 method for signature 'SpatialFeatureExperiment'
annotGeometryNames(x)

## S4 replacement method for signature 'SpatialFeatureExperiment,character'
annotGeometryNames(x) <- value

## S4 method for signature 'SpatialFeatureExperiment'
annotGeometry(x, type = 1L, sample_id = NULL)

## S4 replacement method for signature 'SpatialFeatureExperiment'
annotGeometry(x, type = 1L, sample_id = NULL, translate = TRUE, ...) <- value

tissueBoundary(x, sample_id = 1L)

tissueBoundary(x, sample_id = 1L, translate = TRUE, ...) <- value
```

Arguments

x	A SpatialFeatureExperiment object.
translate	Logical. Only used if removeEmptySpace has been run of the SFE object. If that's the case, this argument indicates whether the new value to be assigned to the geometry is in the coordinates prior to removal of empty space so it should be translated to match the new coordinates after removing empty space. Default to TRUE.
...	spatialCoordsNames, spotDiameter, geometryType passed to df2sf. Defaults are the same as in df2sf. For dimGeometries<- only: geometryType can be a character vector of the geometry type of each data frame in the list of the same length as the list if the data frames specify different types of geometries.
value	Value to set. For annotGeometry, must be a sf data frame, or an ordinary data frame that can be converted to a sf data frame (see df2sf). For annotGeometries, must be a list of such sf or ordinary data frames. There must be a column sample_id to indicate the sample the geometries are for, and the sample_id must also appear in colData.
type	An integer specifying the index or string specifying the name of the *Geometry to query or replace. If missing, then the first item in the *Geometries will be returned or replaced.
sample_id	Sample ID to get or set geometries.

Details

Wrapper for getter and setter of special geometry:

tisseuBoundary Boundary of the tissue of interest, including holes. This is usually of geometry type MULTIPOLYGON, though geometries in annotGeometries can have any type supported by sf.

Value

Getters for multiple geometries return a named list. Getters for names return a character vector of the names. Getters for single geometries return an sf data frame. Setters return an SFE object.

Examples

```
# Example dataset
library(SFEData)
sfe_small <- McKellarMuscleData(dataset = "small")

# Get all annotation geometries, returning a named list
annotGeometries(sfe_small)

# Set all annotation geometries, in a named list
toy <- readRDS(system.file("extdata/sfe_toy.rds",
    package = "SpatialFeatureExperiment"
))
ag <- readRDS(system.file("extdata/ag.rds",
    package = "SpatialFeatureExperiment"
))
annotGeometries(toy) <- list(hull = ag)
```

```
# Get names of annotation geometries
annotGeometryNames(sfe_small)

# Set names of annotation geometries
annotGeometryNames(toy) <- "foo"

# Get a specific annotation geometry by name
# sample_id is optional when there is only one sample present
nuclei <- annotGeometry(sfe_small, type = "nuclei", sample_id = "Vis5A")

# Get a specific annotation geometry by index
tb <- annotGeometry(sfe_small, type = 1L)

# Set a specific annotation geometry
annotGeometry(sfe_small, type = "nuclei2") <- nuclei

# Special convenience function for tissue boundaries
# Getter
tb <- tissueBoundary(sfe_small, sample_id = "Vis5A")
# Setter
tissueBoundary(sfe_small, sample_id = "Vis5A") <- tb
```

---

annotOp *Binary operations for geometry of each cell/spot and annotation*

Description

Just like annotPred, but performs the operation rather than predicate. For example, this function would return the geometry of the intersections between each Visium spot and the tissue boundary for each sample, rather than whether each Visium spot intersects the tissue boundary. In case one cell/spot gets broken up into multiple geometries, the union of those geometries will be taken, so each cell/spot will only get one geometry.

Usage

```
annotOp(
  sfe,
  colGeometryName = 1L,
  annotGeometryName = 1L,
  sample_id = "all",
  op = st_intersection
)
```

Arguments

sfe	An SFE object.
colGeometryName	Name of column geometry for the predicate.
annotGeometryName	Name of annotation geometry for the predicate.
sample_id	Which sample(s) to operate on. Can be "all" to indicate all samples.
op	A binary operation function for the geometries. Defaults to st_intersection.

Value

A `sf` data frame with `geometry` column containing the geometries and corresponding column names of sfe as row names. There is no guarantee that the returned geometries are valid or preserve the geometry class (e.g. when the intersection of polygons result into a line of a point).

See Also

annotPred

Examples

```
library(SFEData)
sfe <- McKellarMuscleData("small")
# Get the intersection of myofibers with each Visium spot
myofibers_on_spots <- annotOp(sfe, "spotPoly",
    annotGeometryName = "myofiber_simplified"
)
```

---

annotPred — *Binary predicates for geometry of each cell/spot and annotation*

Description

This function finds binary predicates for the geometry of each cell/spot (i.e. `colGeometry`) and an annotation geometry for each sample. For example, whether each Visium spot intersects with the tissue boundary in each sample.

Usage

```
annotPred(
  sfe,
  colGeometryName = 1L,
  annotGeometryName = 1L,
  sample_id = "all",
  pred = st_intersects,
  yx = FALSE
)

annotNPred(
  sfe,
  colGeometryName = 1L,
  annotGeometryName = 1L,
  sample_id = "all",
  pred = st_intersects
)
```

Arguments

`sfe`	An SFE object.
`colGeometryName`	Name of column geometry for the predicate.

annotGeometryName
: Name of annotation geometry for the predicate.

sample_id
: Which sample(s) to operate on. Can be "all" to indicate all samples.

pred
: Predicate function to use, defaults to st_intersects.

yx
: Whether to do pred(y, x) instead of pred(x, y). For symmetric predicates, the results should be the same. When x has a large number of geometries and y has few, pred(y, x) is much faster than pred(x, y) for st_intersects, st_disjoint, and st_is_within_distance.

Value

For annotPred, a logical vector of the same length as the number of columns in the sample(s) of interest, with barcodes (or corresponding column names of sfe) as names. For annotNPred, a numeric vector of the same length as the number of columns in the sample(s) of interest with barcodes as names, indicating the number of geometries in the annotGeometry of interest returns TRUE for the predicate for each each geometry in the colGeometry of interest.

See Also

annotOp

Examples

```
library(SFEData)
sfe <- McKellarMuscleData("small")
# Whether each spot is in tissue
in_tissue <- annotPred(sfe, "spotPoly", annotGeometryName = "tissueBoundary")
# How many nuclei are there in each Visium spot
n_nuclei <- annotNPred(sfe, "spotPoly", annotGeometryName = "nuclei")
```

---

annotSummary *Summarize attributes of an annotGeometry for each cell/spot*

Description

In SFE objects, the annotation geometries don't have to correspond to the dimensions of the gene count matrix, so there generally is no one to one mapping between annotation geometries and cells/spots. However, it may be interesting to relate attributes of annotation geometries to cell/spots so the attributes can be related to gene expression. This function summarizes attributes of an annotGeometry for each cell/spot by a geometric predicate with a colGeometry.

Usage

```
annotSummary(
  sfe,
  colGeometryName = 1L,
  annotGeometryName = 1L,
  annotColNames = 1L,
  sample_id = "all",
  pred = st_intersects,
  summary_fun = mean
)
```

Arguments

sfe	An SFE object.
colGeometryName	Name of column geometry for the predicate.
annotGeometryName	Name of annotation geometry for the predicate.
annotColNames	Character, column names of the annotGeometry of interest, to indicate the columns to summarize. Columns that are absent from the annotGeometry are removed. The column cannot be "geometry" or "barcode".
sample_id	Which sample(s) to operate on. Can be "all" to indicate all samples.
pred	Predicate function to use, defaults to st_intersects.
summary_fun	Function for the summary, defaults to mean.

Value

A data frame whose row names are the relevant column names of sfe, and each column of which is the summary of each column specified in annotColName.

Examples

```
library(SFEData)
sfe <- McKellarMuscleData("small")
s <- annotSummary(sfe, "spotPoly", "myofiber_simplified",
    annotColNames = c("area", "convexity")
)
```

---

bbox,SpatialFeatureExperiment-method

Find bounding box of SFE objects

Description

Find bounding box of the union of all colGeometries and annotGeometries of each sample in the SFE object. This can be used to remove empty space so the tissue and geometries have one corner at the origin so all samples will be on comparable coordinates.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
bbox(sfe, sample_id = "all", include_images = FALSE, include_row = TRUE)
```

Arguments

sfe	A SpatialFeatureExperiment object.
sample_id	Sample(s) whose bounding box(es) to find. The bounding box would be for the union of all colGeometries and annotGeometries associated with each sample.
include_images	Logical, whether the bounding boxes should include image extents. Defaults to FALSE because often the image has a lot of empty space surrounding the tissue.
include_row	Logical, whether the bounding boxes should include rowGeometries, defaults to TRUE.

Value

For one sample, then a named vector with names `xmin`, `ymin`, `xmax`, and `ymax` specifying the bounding box. For multiple samples, then a matrix whose columns are samples and whose rows delineate the bounding box.

Examples

```
library(SFEData)
sfe <- McKellarMuscleData("small")
bbox(sfe, sample_id = "Vis5A")
```

---

bbox_center — *Find center of bounding box*

Description

Get x-y coordinates of the center of any bounding box

Usage

```
bbox_center(bbox)
```

Arguments

`bbox`	A numeric vector of length 4 with names xmin, xmax, ymin, ymax, in any order.

Value

A numeric vector of length 2.

Examples

```
bbox <- c(xmin = 0, xmax = 100, ymin = 0, ymax = 80)
bbox_center(bbox)
```

BioFormatsImage — *On disk representation of BioFormats images in SFE object*

Description

‘r lifecycle::badge("experimental")‘ At present, the `BioFormatsImage` is designed for OME-TIFF from Xenium and has not been tested on other formats that can be read with `BioFormats`. The image is not loaded into memory, and when it is, the the `BioFormatsImage` object is converted into `ExtImage` because the loaded image is of a class that inherits from `Image`. The `ExtImage` class is a thin wrapper inheriting from `VirtualSpatialImage` so it's compatible with `SpatialExperiment` from which SFE is derived. This class might drastically change as it matures, say to accommodate other formats supported by `BioFormats` and to store the transformation matrix rather than loading image into memory upon transform.

Usage

```
## S4 method for signature 'BioFormatsImage'
show(object)

BioFormatsImage(
  path,
  ext = NULL,
  isFull = TRUE,
  origin = c(0, 0),
  transformation = list()
)
```

Arguments

`object`	A `BioFormatsImage` object.
`path`	Path to an OME-TIFF image file.
`ext`	Numeric vector with names "xmin", "xmax", "ymin", "ymax" in microns indicating the spatial extent covered by the image. If `NULL`, then the extent will be inferred from the metadata, from physical pixel size and the number of pixels.
`isFull`	Logical, if the extent specified in `ext` is the full extent. If `ext = NULL` so it will be inferred from metadata then `isFull = TRUE` will be set internally.
`origin`	Origin of the whole image in the x-y plane, defaults to `c(0,0)`. This is shifted when the image is translated. This is not the same as xmin and xmax. For example, when the extent is only part of the whole image and the whole image itself can be spatially translated, the origin is needed to determine which part of the whole image this extent corresponds to.
`transformation`	Named list specifying affine transformation. The list can have names "name" and named parameter of the transformation, e.g. `list(name = "mirror", direction = "vertical")`, "rotate" and degrees = 90 (clockwise), and "scale" and factor = 2. The list can also have names "M" for a 2x2 linear transformation matrix in the xy plane and "v" for a translation vector of length 2 to specify general affine transformation.

Details

Spatial extent is inferred from OME-TIFF metadata if not specified. Physical pixel size from the metadata is used to make the extent in micron space. If physical pixel size is absent from metadata, then the extent will be in pixel space, which might mean that the image will not align with the geometries because often the geometry coordinates are in microns, so a warning is issued in this case.

Affine transformations can be specified in the `transformation` argument, either by name or by directly specifying the matrix. The transformations specified by name will always preserve the center of the image. When named transformations are chained, name and parameter will be converted to matrix and translation vector the second time a transformation is specified. If the subsequent transformation happens to restore the image to its original place, then transformation specifications will be removed.

Value

A `BioFormatsImage` object.

See Also

[isFull()], [origin()]

BioFormatsImage-getters

Other BioFormatsImage *getters*

Description

isFULL indicates if the extent is the full extent of the image. origin gets the x-y coordinates of the origin of the image, i.e. the smallest possible x-y coordinate values within the full image.

Usage

```
## S4 method for signature 'BioFormatsImage'
isFull(x)

## S4 method for signature 'BioFormatsImage'
origin(x)

## S4 method for signature 'BioFormatsImage'
transformation(x)
```

Arguments

x	A BioFormatsImage object.

Value

For isFull: Logical scalar indicating whether the extent is the full extent. For origin: Numeric vector of length 2. For transformation, a list.

---

cbind,SpatialFeatureExperiment-method

Concatenate SpatialFeatureExperiment objects

Description

On top of the cbind method of SpatialExperiment, this method is needed to properly merge the spatialGraphs field in the different SFE objects. rowGeometries and annotGeometries also need to be combined properly.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
cbind(..., deparse.level = 1)
```

Arguments

`...`	SFE objects to cbind.
`deparse.level`	See ?`rbind`.

Value

A combined SFE object.

Examples

```
library(SFEData)
sfe_small <- McKellarMuscleData(dataset = "small")
sfe_small2 <- McKellarMuscleData(dataset = "small2")
sfe2 <- cbind(sfe_small, sfe_small2)
```

`changeSampleIDs` *Change sample IDs*

Description

Change sample IDs in all fields of the SFE object where sample IDs are present, not just the colData.

Usage

```
changeSampleIDs(sfe, replacement)
```

Arguments

`sfe`	A `SpatialFeatureExperiment` object.
`replacement`	A named character vector whose names are the existing sample IDs to be changed and whose values are the corresponding replacements.

Value

An SFE object.

Examples

```
library(SFEData)
sfe <- McKellarMuscleData(dataset = "small")
sfe <- changeSampleIDs(sfe, c(Vis5A = "sample01"))
sampleIDs(sfe)
```

colFeatureData *Get global spatial analysis results and metadata of colData, rowData, and geometries*

Description

Results of spatial analyses on columns in colData, rowData, and geometries are stored in their metadata. The colFeaturedata function allows the users to more directly access these results.

Usage

```
colFeatureData(sfe)

rowFeatureData(sfe)

geometryFeatureData(sfe, type, MARGIN = 2L)

reducedDimFeatureData(sfe, dimred)
```

Arguments

sfe	An SFE object.
type	Which geometry, can be name (character) or index (integer)
MARGIN	Integer, 1 means rowGeometry, 2 means colGeometry, and 3 means annotGeometry. Defaults to 2, colGeometry.
dimred	Name of a dimension reduction, can be seen in reducedDimNames.

Value

A DataFrame.

See Also

getParams

Examples

```
library(SpatialFeatureExperiment)
library(SingleCellExperiment)
library(SFEData)
library(Voyager)
sfe <- McKellarMuscleData("small")
colGraph(sfe, "visium") <- findVisiumGraph(sfe)
# Moran's I for colData
sfe <- colDataMoransI(sfe, "nCounts")
colFeatureData(sfe)
```

colGeometries *Column geometry getters and setters*

Description

colGeometries are geometries that correspond to columns of the gene count matrix, such as Visium spots or cells. Same as dimGeometry(x,MARGIN = 2L, ...), with convenience wrappers for getters and setters of special geometries:

spotPoly Polygons of spots from technologies such as Visium, ST, and slide-seq, which do not correspond to cells. Centroids of the polygons are stored in spatialCoords of the underlying SpatialExperiment object.

ROIPoly Polygons of regions of interest (ROIs) from technologies such as laser capture microdissection (LCM) and GeoMX DSP. These should correspond to columns of the gene count matrix.

cellSeg Cell segmentation polygons. If the columns of the gene count matrix are single cells, then this is stored in colGeometries. Otherwise, this is stored in annotGeometries.

nucSeg Similar to cellSeg, but for nuclei rather than whole cell.

Usage

```
colGeometry(x, type = 1L, sample_id = 1L, withDimnames = TRUE)

colGeometry(
  x,
  type = 1L,
  sample_id = 1L,
  withDimnames = TRUE,
  translate = TRUE
) <- value

colGeometries(x, withDimnames = TRUE)

colGeometries(x, withDimnames = TRUE, translate = TRUE) <- value

colGeometryNames(x)

colGeometryNames(x) <- value

spotPoly(x, sample_id = 1L, withDimnames = TRUE)

spotPoly(x, sample_id = 1L, withDimnames = TRUE, translate = TRUE) <- value

centroids(x, sample_id = 1L, withDimnames = TRUE)

centroids(x, sample_id = 1L, withDimnames = TRUE, translate = TRUE) <- value

ROIPoly(x, sample_id = 1L, withDimnames = TRUE)

ROIPoly(x, sample_id = 1L, withDimnames = TRUE, translate = TRUE) <- value
```

```
cellSeg(x, sample_id = 1L, withDimnames = TRUE)

cellSeg(x, sample_id = 1L, withDimnames = TRUE, translate = TRUE) <- value

nucSeg(x, sample_id = 1L, withDimnames = TRUE)

nucSeg(x, sample_id = 1L, withDimnames = TRUE, translate = TRUE) <- value
```

Arguments

x	A SpatialFeatureExperiment object.
type	An integer specifying the index or string specifying the name of the *Geometry to query or replace. If missing, then the first item in the *Geometries will be returned or replaced.
sample_id	Sample ID to get or set geometries.
withDimnames	Logical. If TRUE, then the dimnames (colnames or rownames) of the gene count matrix should correspond to row names of the sf data frames of interest.
translate	Logical. Only used if removeEmptySpace has been run of the SFE object. If that's the case, this argument indicates whether the new value to be assigned to the geometry is in the coordinates prior to removal of empty space so it should be translated to match the new coordinates after removing empty space. Default to TRUE.
value	Value to set. For dimGeometry, must be a sf data frame with the same number of rows as size in the dimension of interest, or an ordinary data frame that can be converted to such a sf data frame (see df2sf). For dimGeometries, must be a list of such sf or ordinary data frames.

See Also

[dimGeometries()], [rowGeometries()]

Examples

```
library(SFEData)
sfe <- McKellarMuscleData(dataset = "small")
cgs <- colGeometries(sfe)
spots <- spotPoly(sfe)
```

---

containsOutOfMemoryData,SpatialFeatureExperiment-method

Whether an SFE object contains out of memory data

Description

Out of memory data, such as DelayedArray, some SpatRasterImage objects, and BioFormatsImage, will break if saved as RDS. This method of containsOutOfMemoryData checks if an SFE object has out of memory data, specifically the images. Having out of memory data will result into an error when saveRDS is called; we recommend using the alabaster.sfe package instead.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
containsOutOfMemoryData(object)
```

Arguments

object	An SFE object

Value

TRUE or FALSE

Examples

```
outdir <- system.file("extdata", package = "SpatialFeatureExperiment")
samples <- file.path(outdir, paste0("sample0", 1:2))
sfe <- read10xVisiumSFE(samples, type = "sparse", data = "filtered")
containsOutOfMemoryData(sfe)
```

---

crop — *Crop an SFE object with a geometry*

Description

Returns an SFE object whose specified `colGeometry` returns `TRUE` with a geometric predicate function (usually intersects) with another geometry of interest. This can be used to subset an SFE object with a tissue boundary or histological region polygon, or crop away empty spaces. After cropping, not only will the cells/spots be subsetted, but also all geometries will be cropped.

Usage

```
crop(
  x,
  y = NULL,
  colGeometryName = 1L,
  sample_id = "all",
  op = st_intersection,
  keep_whole = "none",
  cover = FALSE
)
```

Arguments

x	An SFE object.
y	An object of class `sf`, `sfg`, `sfc` with which to crop the SFE object, or a bounding box with the format of the output of `bbox,SpatialFeatureExperiment-method`.
colGeometryName	Column geometry to used to indicate which cells/spots to keep.

`sample_id` Samples to crop. Optional when only one sample is present. Can be multiple samples, or "all", which means all samples. For multiple samples, `sf` data frame `y` may have column `sample_id` indicating which geometry subsets which sample or matrix `y` may indicate sample specific bounding boxes in its column names. Only samples included in the indicated sample IDs are subsetted. If sample is not indicated in `y`, then the same geometry or bounding box is used to subset all samples specified in the `sample_id` argument.

`op` A geometric operation function to crop the geometries in the SFE object. Only `st_intersection` and `st_difference` are allowed. If "intersection", then only things inside `y` is kept after cropping. If "difference", then only things outside `y` is kept.

`keep_whole` Character vector, can be one or more of "col" and "annot" to keep whole items from `colGeometries` or `annotGeometries`, keeping geometries that partially intersect with `y` whole. This can greatly speed up code while not breaking geometries into multiple pieces. Can also be "none" so all geometries are actually cropped.

`cover` Logical, whether the geometries in `x` must be entirely covered by `y` if `op = st_intersection` or whether `x` must be entirely outside `y` if `op = st_difference`. Only relevant when `keep_whole != "none"`.

Details

3D geometries are allowed, but geometric operations can only be performed in x and y but not z.

Value

An SFE object. There is no guarantee that the geometries after cropping are still all valid or preserve the original geometry class.

Examples

```
library(SFEData)
sfe <- McKellarMuscleData("small")
# Subset sfe to only keep spots on tissue
sfe_on_tissue <- crop(sfe, tissueBoundary(sfe),
    colGeometryName = "spotPoly",
    sample_id = "Vis5A"
)
```

`cropImg` *Crop images*

Description

Crop images of class `*Image` in this package with a bounding box.

Usage

```
## S4 method for signature 'SpatRasterImage'
cropImg(x, bbox, filename = "")

## S4 method for signature 'BioFormatsImage'
cropImg(x, bbox)

## S4 method for signature 'ExtImage'
cropImg(x, bbox)
```

Arguments

x	An object of class *Image as implemented in this package.
bbox	Numeric vector with names "xmin", "xmax", "ymin", "ymax", in any order, to specify the bounding box.
filename	Output file name for transformed SpatRaster.

Value

Image of the same class as input but cropped. For BioFormatsImage, the image is not loaded into memory; only the extent is changed.

See Also

Other image methods: SFE-image, affineImg(), dim,BioFormatsImage-method, dim,ExtImage-method, ext(), imgRaster(), imgSource(), mirrorImg(), rotateImg(), scaleImg(), translateImg(), transposeImg()

---

df2sf — *From ordinary data frame to sf to construct SFE object*

Description

While the SpatialFeatureExperiment constructor and *Geometry replacement methods can convert properly formatted ordinary data frames into sf objects which are used to store the geometries internally, the user might want to do the conversion, check if the geometry is valid, and inspect and fix any invalid geometries.

Usage

```
df2sf(
  df,
  spatialCoordsNames = c("x", "y"),
  spotDiameter = NA,
 geometryType = c("POINT", "LINESTRING", "POLYGON", "MULTIPOINT", "MULTILINESTRING",
    "MULTIPOLYGON"),
  group_col = "group",
  id_col = "ID",
  subid_col = "subID",
  check = TRUE,
  ...
)
```

Arguments

- `df` — An ordinary data frame, i.e. not `sf`. Or a matrix that can be converted to a data frame.
- `spatialCoordsNames` — Column names in `df` that specify spatial coordinates.
- `spotDiameter` — Spot diameter for technologies with arrays of spots of fixed diameter per slide, such as Visium, ST, DBiT-seq, and slide-seq. The diameter must be in the same unit as the coordinates in the *Geometry arguments. Ignored for geometries that are not POINT or MULTIPOINT.
- `geometryType` — Type of geometry to convert the ordinary data frame to. If the geometry in `df` is de facto points, then this argument will be ignored and the returned `sf` will have geometry type POINT.
- `group_col` — Column to indicate which coordinates for which MULTI geometry, such as to identify which MULTIPOLYGON or MULTIPOINT.
- `id_col` — Column to indicate coordinates for which geometry, within a MULTI geometry if applicable, such as to identify which POLYGON or which polygon within a MULTIPOLYGON.
- `subid_col` — Column to indicate coordinates for holes in polygons.
- `check` — Logical, whether to check the input data frame for issues related to constructing the geometry of interese such as number of vertices per geometry. If `FALSE`, it will save a bit of time, which is useful when the input is already known to be good.
- `...` — Other arguments passed to ‘sf::st_buffer‘, mainly to make polygon shapes, eg Visium spot ‘endCapStyle = "ROUND"‘ and VisiumHD bin ‘endCapStyle = "SQUARE"‘

Value

An `sf` object.

Examples

```
# Points, use spotDiameter to convert to circle polygons
# This is done to Visium spots
pts_df <- readRDS(system.file("extdata/pts_df.rds",
    package = "SpatialFeatureExperiment"
))
sf_use <- df2sf(pts_df, geometryType = "POINT", spotDiameter = 0.1)
# Linestring
ls_df <- readRDS(system.file("extdata/ls_df.rds",
    package = "SpatialFeatureExperiment"
))
sf_use <- df2sf(ls_df, geometryType = "LINESTRING")
# Polygon
pol_df <- readRDS(system.file("extdata/pol_df.rds",
    package = "SpatialFeatureExperiment"
))
sf_use <- df2sf(pol_df,
    geometryType = "POLYGON",
    spatialCoordsNames = c("V1", "V2")
)
# Multipolygon
```

```
mpol_df <- readRDS(system.file("extdata/mpol_df.rds",
    package = "SpatialFeatureExperiment"
))
sf_use <- df2sf(mpol_df,
    geometryType = "MULTIPOLYGON",
    spatialCoordsNames = c("V1", "V2")
)
# Multiple sample_ids present
multipts_df <- readRDS(system.file("extdata/multipts_df.rds",
    package = "SpatialFeatureExperiment"
))
sf_use <- df2sf(multipts_df, geometryType = "MULTIPOINT")
```

`dim,BioFormatsImage-method`

Find dimension of BioFormatsImage

Description

This is different from other classes. The metadata is read where the dimensions in pixels can be found. The image itself is not read into memory here.

Usage

```
## S4 method for signature 'BioFormatsImage'
dim(x)
```

Arguments

x	A `BioFormatsImage` object.

Value

An integer vector of length 5 showing the number of rows and columns in the full resolution image. The 5 dimensions are in the order of XYCZT: x, y, channel, z, and time. This is not changed by transformations. Use `ext` to see the extent after transformation.

See Also

Other image methods: `SFE-image`, `affineImg()`, `cropImg()`, `dim,ExtImage-method`, `ext()`, `imgRaster()`, `imgSource()`, `mirrorImg()`, `rotateImg()`, `scaleImg()`, `translateImg()`, `transposeImg()`

dim,ExtImage-method *Find dimensions of ExtImage*

Description

This method exists to make the output of dim() for ExtImage consistent with that of Image which ExtImage inherits from, overriding the VirtualSpatialImage method.

Usage

```
## S4 method for signature 'ExtImage'
dim(x)
```

Arguments

x	A ExtImage object.

Value

An integer vector. As in EBImage, the first element indicates number of pixels in the x direction, or number of columns in the image, and the second element indicates the number of pixels in the y direction. This is unlike array indexing.

See Also

Other image methods: SFE-image, affineImg(), cropImg(), dim,BioFormatsImage-method, ext(), imgRaster(), imgSource(), mirrorImg(), rotateImg(), scaleImg(), translateImg(), transposeImg()

dimGeometries *Dimension geometry methods*

Description

"Dimension geometry" refers to Simple Feature (sf) geometries associated with rows (features, genes) or columns (cells or spots) of the gene count matrix in the SpatialFeatureExperiment object. For each dimension, the number of rows in the sf data frame specifying the geometries must match the size of the dimension of interest. For example, there must be the same number of rows in the sf data frame describing cells as there are cells in the gene count matrix. This page documents getters and setters for the dimension geometries. The getters and setters are implemented in a way similar to those of reducedDims in SingleCellExperiment.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
dimGeometries(x, MARGIN = 2, withDimnames = TRUE)

## S4 replacement method for signature 'SpatialFeatureExperiment'
dimGeometries(x, MARGIN, withDimnames = TRUE, translate = TRUE, ...) <- value
```

```
## S4 method for signature 'SpatialFeatureExperiment'
dimGeometryNames(x, MARGIN)

## S4 replacement method for signature 'SpatialFeatureExperiment,numeric,character'
dimGeometryNames(x, MARGIN) <- value

## S4 method for signature 'SpatialFeatureExperiment'
dimGeometry(x, type = 1L, MARGIN, sample_id = 1L, withDimnames = TRUE)

## S4 replacement method for signature 'SpatialFeatureExperiment'
dimGeometry(
  x,
  type = 1L,
  MARGIN,
  sample_id = 1L,
  withDimnames = TRUE,
  translate = TRUE,
  ...
) <- value
```

Arguments

x	A SpatialFeatureExperiment object.
MARGIN	As in apply. 1 stands for rows and 2 stands for columns.
withDimnames	Logical. If TRUE, then the dimnames (colnames or rownames) of the gene count matrix should correspond to row names of the sf data frames of interest.
translate	Logical. Only used if removeEmptySpace has been run of the SFE object. If that's the case, this argument indicates whether the new value to be assigned to the geometry is in the coordinates prior to removal of empty space so it should be translated to match the new coordinates after removing empty space. Default to TRUE.
...	spatialCoordsNames, spotDiameter, geometryType passed to df2sf. Defaults are the same as in df2sf. For dimGeometries<- only: geometryType can be a character vector of the geometry type of each data frame in the list of the same length as the list if the data frames specify different types of geometries.
value	Value to set. For dimGeometry, must be a sf data frame with the same number of rows as size in the dimension of interest, or an ordinary data frame that can be converted to such a sf data frame (see df2sf). For dimGeometries, must be a list of such sf or ordinary data frames.
type	An integer specifying the index or string specifying the name of the *Geometry to query or replace. If missing, then the first item in the *Geometries will be returned or replaced.
sample_id	Sample ID to get or set geometries.

Value

Getters for multiple geometries return a named list. Getters for names return a character vector of the names. Getters for single geometries return an sf data frame. Setters return an SFE object.

See Also

[colGeometries()], [rowGeometries()]

Examples

```
library(SFEData)
sfe <- McKellarMuscleData(dataset = "small")

# Get all column geometries as a named list
# Use MARGIN = 1 or rowGeometry/ies for rowGeometries
cgs <- dimGeometries(sfe, MARGIN = 2)
# Or equivalently
cgs <- colGeometries(sfe)

# Set all column geometries with a named list
dimGeometries(sfe, MARGIN = 2) <- cgs
# Or equivalently
colGeometries(sfe) <- cgs

# Get names of column geometries
cgns <- dimGeometryNames(sfe, MARGIN = 2)
cgns <- colGeometryNames(sfe)

# Set column geometry names
dimGeometryNames(sfe, MARGIN = 2) <- cgns
colGeometryNames(sfe) <- cgns

# Get a specific column geometry by name
spots <- dimGeometry(sfe, "spotPoly", MARGIN = 2)
spots <- colGeometry(sfe, "spotPoly")
# Or equivalently, the wrapper specifically for Visium spot polygons,
# for the name "spotPoly"
spots <- spotPoly(sfe)
# Other colGeometry wrappers for specific names:
# ROIPoly (for LCM and GeoMX DSP), cellSeg and nucSeg (for MERFISH; would
# query annotGeometries for Visium)
# rowGeometry wrappers for specific names: txSpots (MERFISH transcript spots)
# By index
spots <- colGeometry(sfe, 1L)

# Multiple samples, only get geometries for one sample
sfe2 <- McKellarMuscleData("small2")
sfe_combined <- cbind(sfe, sfe2)
spots1 <- colGeometry(sfe, "spotPoly", sample_id = "Vis5A")
spots2 <- spotPoly(sfe_combined, sample_id = "sample02")
# Get geometries for multiple samples
spots3 <- spotPoly(sfe_combined, sample_id = c("Vis5A", "sample02"))
# All samples
spots3 <- spotPoly(sfe_combined, sample_id = "all")

# Set specific column geometry by name
colGeometry(sfe, "foobar") <- spots
# Or use wrapper
spotPoly(sfe) <- spots
# Specify sample_id
colGeometry(sfe_combined, "foobar", sample_id = "Vis5A") <- spots1
```

```
# Only entries for the specified sample are set.
foobar <- colGeometry(sfe_combined, "foobar", sample_id = "sample02")
```

---

ext *Get and set extent of image objects*

Description

Unlike in SpatialExperiment, images in SFE have extents which are used to align them to the geometries and in geometric operations on SFE objects. These functions get or set the extent for S4 image classes inheriting from VirtualSpatialImage implemented in the SFE package.

Usage

```
## S4 method for signature 'BioFormatsImage'
ext(x)

## S4 method for signature 'ExtImage'
ext(x)

## S4 method for signature 'SpatRasterImage'
ext(x)

## S4 replacement method for signature 'BioFormatsImage,numeric'
ext(x) <- value

## S4 replacement method for signature 'ExtImage,numeric'
ext(x) <- value

## S4 replacement method for signature 'SpatRasterImage,numeric'
ext(x) <- value
```

Arguments

x	A *Image object.
value	A numeric vector with names "xmin", "xmax", "ymin", "ymax" specifying the extent to use.

Value

Getters return a numeric vector specifying the extent. Setters return a *Image object of the same class as the input.

Note

For SpatRasterImage, the image may be may not be loaded into memory. You can check if the image is loaded into memory with terra::inMemory(x), and check the original file path with imgSource. If the image is not loaded into memory, then the original file must be present at the path indicated by imgSource in order for any code using the image to work, which includes this function ext.

For BioFormatsImage, internally only the pre-transform extent is stored. The ext getter will apply the transformation on the fly. The setter sets the pre-transformation extent.

See Also

Other image methods: SFE-image, affineImg(), cropImg(), dim,BioFormatsImage-method, dim,ExtImage-method, imgRaster(), imgSource(), mirrorImg(), rotateImg(), scaleImg(), translateImg(), transposeImg()

ExtImage *Use the EBImage* Image *class in SFE objects*

Description

This is a thin wrapper around the Image class in the EBImage package so it inherits from VirtualSpatialImage to be compatible with SpatialExperiment from which SFE inherits. An ext field is added to specify the spatial extent of the image in microns to facilitate geometric operations on the SFE object (including the images) and plotting with Voyager.

Usage

```
## S4 method for signature 'ExtImage'
show(object)

ExtImage(img, ext = NULL)
```

Arguments

object	An ExtImage object.
img	An Image object or anything that inherits from Image such as AnnotatedImage in RBioFormats.
ext	Numeric vector with names "xmin", "xmax", "ymin", "ymax" in microns indicating the spatial extent covered by the image. If NULL, then the extent will be inferred from the metadata, from physical pixel size and the number of pixels.

Value

An ExtImage object.

---

findSpatialNeighbors,SpatialFeatureExperiment-method *Find spatial neighborhood graph*

Description

This function wraps all spatial neighborhood graphs implemented in the package spdep for the SpatialFeatureExperiment (SFE) class, to find spatial neighborhood graphs for the entities represented by columns or rows of the gene count matrix in the SFE object or spatial entities in the annotGeometries field of the SFE object. Results are stored as listw objects in the spatialGraphs field of the SFE object, as listw is used in many methods that facilitate the spatial neighborhood graph in the spdep, spatialreg, and adespatial. The edge weights of the graph in the listw object are by default style W (see nb2listw) and the unweighted neighbor list is in the neighbours field of the listw object.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
findSpatialNeighbors(
  x,
  sample_id = "all",
  type = "spatialCoords",
  MARGIN = 2,
 method = c("tri2nb", "knearneigh", "dnearneigh", "gabrielneigh", "relativeneigh",
    "soi.graph", "poly2nb"),
  dist_type = c("none", "idw", "exp", "dpd"),
  glist = NULL,
  style = c("raw", "W", "B", "C", "U", "minmax", "S"),
  nn_method = c("bioc", "spdep"),
  alpha = 1,
  dmax = NULL,
  BPPARAM = SerialParam(),
  BNPARAM = KmknnParam(),
  zero.policy = TRUE,
  ...
)
```

Arguments

x	A SpatialFeatureExperiment object.
sample_id	Which sample(s) in the SFE object to use for the graph. Can also be "all", which means this function will compute the graph for all samples independently.
type	Name of the geometry associated with the MARGIN of interest for which to compute the graph.
MARGIN	Just like in apply, where 1 stands for row, 2 stands for column. Here, in addition, 3 stands for annotation, to query the annotGeometries, such as nuclei segmentation in a Visium data
method	Name of function in the package spdep to use to find the spatial neighborhood graph.
dist_type	Type of distance-based weight. "none" means not using distance-based weights; the edge weights of the spatial neighborhood graph will be entirely determined by the style argument. "idw" means inverse distance weighting. "exp" means exponential decay. "dpd" means double-power distance weights. See nb2listwdist for details.
glist	list of general weights corresponding to neighbours
style	style can take values “W”, “B”, “C”, “U”, “minmax” and “S”
nn_method	Method to find k nearest neighbors and distance based neighbors. Can be either "bioc" or "spdep". For "bioc", methods from BiocNeighbors are used. For "spdep", methods from the spdep package are used. The "bioc" option is more scalable to larger datasets and supports multithreading.
alpha	Only relevant when dist_type = "dpd".
dmax	Only relevant when dist_type = "dpd".
BPPARAM	A BiocParallelParam object for multithreading. Only used for k nearest neighbor and distance based neighbor with nn_method = "bioc".

BNPARAM	A BiocNeighborParam object specifying the algorithm to find k nearest neighbors and distance based neighbors with nn_method = "bioc". For distance based neighbors, only KmknnParam and VptreeParam are applicable.
zero.policy	default NULL, use global option value; if FALSE stop with error for any empty neighbour sets, if TRUE permit the weights list to be formed with zero-length weights vectors
...	Extra arguments passed to the spdep function stated in the method argument, such as k, use_kd_tree, d1, d2, nnmult, sym, and quadsegs. Note that any arguments about using longitude and latitude, which are irrelevant, are ignored.

Value

For one sample, then a listw object representing the graph, with an attribute "method" recording the function used to build the graph, its arguments, and information about the geometry for which the graph was built. The attribute is used to reconstruct the graphs when the SFE object is subsetted since some nodes in the graph will no longer be present. If sample_id = "all" or has length > 1, then a named list of listw objects, whose names are the sample_ids. To add the list for multiple samples to a SFE object, specify the name argument in the spatialGraphs replacement method, so graph of the same name will be added to the SFE object for each sample.

Note

style = "raw" is only applicable when dist_type is not "none". If dist_type = "none" and style = "raw", then style will default to "W". Using distance based weights does not supplant finding a spatial neighborhood graph. The spatial neighborhood graph is first found and then its edges weighted based on distance in this function.

Examples

```
library(SFEData)
sfe <- McKellarMuscleData(dataset = "small")
# sample_id is optional when only one sample is present
g <- findSpatialNeighbors(sfe, sample_id = "Vis5A")
attr(g, "method")
# Returns named list for multiple samples
sfe2 <- McKellarMuscleData(dataset = "small2")
sfe_combined <- cbind(sfe, sfe2)
gs <- findSpatialNeighbors(sfe, sample_id = "all")
```

---

findVisiumGraph *Find spatial neighborhood graphs for Visium spots*

Description

Visium spots are arranged in a hexagonal grid. This function uses the known locations of the Visium barcodes to construct a neighborhood graph, so adjacent spots are connected by edges. Since the known rows and columns of the spots are used, the unit the spot centroid coordinates are in does not matter.

Usage

```
findVisiumGraph(x, sample_id = "all", style = "W", zero.policy = NULL)
```

Arguments

x	A SpatialFeatureExperiment object with Visium data. Column names of the gene count matrix must be Visium barcodes, which may have a numeric suffix to distinguish between samples (e.g. "AAACAACGAATAGTTC-1").
sample_id	Which sample(s) in the SFE object to use for the graph. Can also be "all", which means this function will compute the graph for all samples independently.
style	style can take values “W”, “B”, “C”, “U”, “minmax” and “S”
zero.policy	default NULL, use global option value; if FALSE stop with error for any empty neighbour sets, if TRUE permit the weights list to be formed with zero-length weights vectors

Value

For one sample, then a listw object representing the graph, with an attribute "method" recording the function used to build the graph, its arguments, and information about the geometry for which the graph was built. The attribute is used to reconstruct the graphs when the SFE object is subsetted since some nodes in the graph will no longer be present. If sample_id = "all" or has length > 1, then a named list of listw objects, whose names are the sample_ids. To add the list for multiple samples to a SFE object, specify the name argument in the spatialGraphs replacement method, so graph of the same name will be added to the SFE object for each sample.

Examples

```
library(SFEData)
sfe <- McKellarMuscleData(dataset = "small")
g <- findVisiumGraph(sfe)
# For multiple samples, returns named list
sfe2 <- McKellarMuscleData(dataset = "small2")
sfe_combined <- cbind(sfe, sfe2)
gs <- findVisiumGraph(sfe, sample_id = "all")
```

---

findVisiumHDGraph *Find Visium HD spatial neighborhood graph*

Description

Visium HD spots are arranged in a square grid. This function finds either a rook or a queen spatial neighborhood graph for the spots. colData of the SFE object must have columns array_row and array_col.

Usage

```
findVisiumHDGraph(x, style = "W", queen = FALSE, zero.policy = TRUE)
```

Arguments

x	An SFE object with Visium HD data with one sample with the required information in its colData.
style	style can take values “W”, “B”, “C”, “U”, “minmax” and “S”
queen	Logical. Default is FALSE, using rook neighbors.

`zero.policy` default NULL, use global option value; if FALSE stop with error for any empty neighbour sets, if TRUE permit the weights list to be formed with zero-length weights vectors

Value

A `listw` object for the graph.

`formatTxSpots` *Read and process transcript spots geometry for SFE*

Description

The function ‘formatTxSpots‘ reads the transcript spot coordinates of smFISH-based data and formats the data. The data is not added to an SFE object. If the file specified in ‘file_out‘ already exists, then this file will be read instead of the original file in the ‘file‘ argument, so the processing is not run multiple times. The function ‘addTxSpots‘ adds the data read and processed in ‘formatTxSpots‘ to the SFE object, and reads all transcript spot data. To only read a subset of transcript spot data, first use ‘formatTxSpots‘ to write the re-formatted data to disk. Then read the specific subset and add them separately to the SFE object with the setter functions.

Usage

```
formatTxSpots(
  file,
  dest = c("rowGeometry", "colGeometry"),
  spatialCoordsNames = c("global_x", "global_y", "global_z"),
  gene_col = "gene",
  cell_col = "cell_id",
  z = "all",
  phred_col = "qv",
  min_phred = 20,
  split_col = NULL,
  not_in_cell_id = c("-1", "UNASSIGNED"),
  z_option = c("3d", "split"),
  flip = FALSE,
  file_out = NULL,
  BPPARAM = SerialParam(),
  return = TRUE
)

addTxSpots(
  sfe,
  file,
  sample_id = 1L,
  spatialCoordsNames = c("global_x", "global_y", "global_z"),
  gene_col = "gene",
  z = "all",
  phred_col = "qv",
  min_phred = 20,
  split_col = NULL,
```

```
  z_option = c("3d", "split"),
  flip = FALSE,
  file_out = NULL,
  BPPARAM = SerialParam()
)
```

Arguments

`file` File with the transcript spot coordinates. Should be one row per spot when read into R and should have columns for coordinates on each axis, gene the transcript is assigned to, and optionally cell the transcript is assigned to. Must be csv, tsv, or parquet.

`dest` Where in the SFE object to store the spot geometries. This affects how the data is processed. Options:

rowGeometry All spots for each gene will be a ‘MULTIPOINT‘ geometry, regardless of whether they are in cells or which cells they are assigned to.

colGeometry The spots for each gene assigned to a cell of interest will be a ‘MULTIPOINT‘ geometry; since the gene count matrix is sparse, the geometries are NOT returned to memory.

`spatialCoordsNames` Column names for the x, y, and optionally z coordinates of the spots. The defaults are for Vizgen.

`gene_col` Column name for genes.

`cell_col` Column name for cell IDs, ignored if ‘dest = "rowGeometry"‘. Can have length > 1 when multiple columns are needed to uniquely identify cells, in which case the contents of the columns will be concatenated, such as in CosMX data where cell ID is only unique within the same FOV. Default "cell_id" is for Vizgen MERFISH. Should be ‘c("cell_ID", "fov")‘ for CosMX.

`z` Index of z plane to read. Can be "all" to read all z-planes into MULTIPOINT geometries with XYZ coordinates. If z values are not integer, then spots with all z values will be read.

`phred_col` Column name for Phred scores of the spots.

`min_phred` Minimum Phred score to keep spot. By default 20, the conventional threshold indicating "acceptable", meaning that there’s 1 chance that the spot was decoded in error.

`split_col` Categorical column to split the geometries, such as cell compartment the spots are assigned to as in the "CellComp" column in CosMX output.

`not_in_cell_id` Value of cell ID indicating that the spot is not assigned to any cell, such as "-1" in Vizgen MERFISH and "0" in CosMX. When there’re multiple columns for ‘cell_col‘, the first column is used to identify spots that are not in cells.

`z_option` What to do with z coordinates. "3d" is to construct 3D geometries. "split" is to create a separate 2D geometry for each z-plane so geometric operations are fully supported but some data wrangling is required to perform 3D analyses. When the z coordinates are not integers, 3D geometries will always be constructed since there are no z-planes to speak of. This argument does not apply when ‘spatialCoordsNames‘ has length 2.

`flip` Logical, whether to flip the geometry to match image. Here the y coordinates are simply set to -y, so the original bounding box is not preserved. This is consistent with `readVizgen` and `readXenium`.

file_out	Name of file to save the geometry or raster to disk. Especially when the geometries are so large that it's unwieldy to load everything into memory. If this file (or directory for multiple files) already exists, then the existing file(s) will be read, skipping the processing. When writing the file, extensions supplied are ignored and extensions are determined based on 'dest'.
BPPARAM	BiocParallelParam object to specify multithreading to convert raw char in some parquet files to R objects. Not used otherwise.
return	Logical, whether to return the geometries in memory. This does not depend on whether the geometries are written to file. Always 'FALSE' when 'dest = "colGeometry"'.
sfe	A 'SpatialFeatureExperiment' object.
sample_id	Which sample in the SFE object the transcript spots should be added to.

Value

A sf data frame for vector geometries if 'file_out' is not set. 'SpatRaster' for raster. If there are multiple files written, such as when splitting by cell compartment or when 'dest = "colGeometry"', then a directory with the same name as 'file_out' will be created (but without the extension) and the files are written to that directory with informative names. 'parquet' files that can be read with 'st_read' is written for vector geometries. When 'return = FALSE', the file name or directory (when there're multiple files) is returned.

The 'sf' data frame, or path to file where geometries are written if 'return = FALSE'.

Note

When 'dest = "colGeometry"', the geometries are always written to disk and not returned in memory, because this is essentially the gene count matrix, which is sparse. This kind of reformatting is implemented so users can read in MULTIPOINT geometries with transcript spots for each gene assigned to each cell for spatial point process analyses, where not all genes are loaded at once.

Examples

```
# Default arguments are for MERFISH
fp <- tempfile()
dir_use <- SFEData::VizgenOutput(file_path = fp)
g <- formatTxSpots(file.path(dir_use, "detected_transcripts.csv"))
unlink(dir_use, recursive = TRUE)

# For CosMX, note the colnames, also dest = "colGeometry"
# Results are written to the tx_spots directory
dir_use <- SFEData::CosMXOutput(file_path = fp)
cg <- formatTxSpots(file.path(dir_use, "Run5642_S3_Quarter_tx_file.csv"),
dest = "colGeometry", z = "all",
cell_col = c("cell_ID", "fov"),
gene_col = "target", not_in_cell_id = "0",
spatialCoordsNames = c("x_global_px", "y_global_px", "z"),
file_out = file.path(dir_use, "tx_spots"))
# Cleanup
unlink(dir_use, recursive = TRUE)
```

formatTxTech *Read and process transcript spots for specific commercial technologies*

Description

To preset parameters such as spatialCoordsNames, gene_col, cell_col, and phred_col that are standard for the output of the technology.

Usage

```
formatTxTech(
  data_dir,
  tech = c("Vizgen", "Xenium", "CosMX"),
  dest = c("rowGeometry", "colGeometry"),
  z = "all",
  min_phred = 20,
  split_cell_comps = FALSE,
  z_option = c("3d", "split"),
  flip = FALSE,
  file_out = NULL,
  BPPARAM = SerialParam(),
  return = TRUE
)

addTxTech(
  sfe,
  data_dir,
  sample_id = 1L,
  tech = c("Vizgen", "Xenium", "CosMX"),
  z = "all",
  min_phred = 20,
  split_cell_comps = FALSE,
  z_option = c("3d", "split"),
  flip = FALSE,
  file_out = NULL,
  BPPARAM = SerialParam()
)
```

Arguments

data_dir Top level output directory.

tech Which technology whose output to read, must be one of "Vizgen", "Xenium", or "CosMX" though more technologies may be added later.

dest Where in the SFE object to store the spot geometries. This affects how the data is processed. Options:

rowGeometry All spots for each gene will be a ‘MULTIPOINT‘ geometry, regardless of whether they are in cells or which cells they are assigned to.

colGeometry The spots for each gene assigned to a cell of interest will be a ‘MULTIPOINT‘ geometry; since the gene count matrix is sparse, the geometries are NOT returned to memory.

z — Which z-planes to read. Always "all" for Xenium where the z coordinates are not discrete.

min_phred — Minimum Phred score to keep spot. By default 20, the conventional threshold indicating "acceptable", meaning that there's 1 chance that the spot was decoded in error.

split_cell_comps — Only relevant to CosMX whose transcript spot file assigns the spots to cell components. Setting this argument to TRUE

z_option — What to do with z coordinates. "3d" is to construct 3D geometries. "split" is to create a separate 2D geometry for each z-plane so geometric operations are fully supported but some data wrangling is required to perform 3D analyses. When the z coordinates are not integers, 3D geometries will always be constructed since there are no z-planes to speak of. This argument does not apply when 'spatialCoordsNames' has length 2.

flip — Logical, whether to flip the geometry to match image. Here the y coordinates are simply set to -y, so the original bounding box is not preserved. This is consistent with readVizgen and readXenium.

file_out — Name of file to save the geometry or raster to disk. Especially when the geometries are so large that it's unwieldy to load everything into memory. If this file (or directory for multiple files) already exists, then the existing file(s) will be read, skipping the processing. When writing the file, extensions supplied are ignored and extensions are determined based on 'dest'.

BPPARAM — BiocParallelParam object to specify multithreading to convert raw char in some parquet files to R objects. Not used otherwise.

return — Logical, whether to return the geometries in memory. This does not depend on whether the geometries are written to file. Always 'FALSE' when 'dest = "colGeometry"'.

sfe — A 'SpatialFeatureExperiment' object.

sample_id — Which sample in the SFE object the transcript spots should be added to.

Value

The 'sf' data frame, or path to file where geometries are written if 'return = FALSE'.

Examples

```
library(SFEData)
fp <- tempfile()
dir_use <- XeniumOutput("v2", file_path = fp)
fn_tx <- formatTxTech(dir_use, tech = "Xenium", flip = TRUE, return = FALSE,
                      file_out = file.path(dir_use, "tx_spots.parquet"))
```

---

gdalParquetAvailable *Check if Parquet GDAL driver is available*

Description

The GeoParquet files for geometries are typically written and read with the `sfarrow` package, but to add only a select few genes to the SFE object say for visualization purposes, the Parquet GDAL driver is required in order to use GDAL's SQL to query the GeoParquet file to only load the few genes requested. The transcript spots from a large dataset can take up a lot of memory if all loaded.

Usage

```
gdalParquetAvailable()
```

Details

The Parquet driver has been supported since GDAL 3.5.0. The `arrow` C++ library must be installed in order to make the Parquet driver available. When arrow is installed, newer versions of GDAL installed from Homebrew (Mac) should have the Parquet driver. For Linux, the binary from `apt-get`'s default repo is 3.4.1 (as of April 2024). To use the Parquet driver, GDAL may need to be installed from source. See script from the geospatial rocker. A Voyager docker container with the Parquet driver will soon be provided.

Value

Logical, indicating whether the Parquet driver is present.

Examples

```
gdalParquetAvailable()
```

`getParams` *Get parameters used in spatial methods*

Description

The `getParams` function allows users to access the parameters used to compute the results that may be stored in `colFeatureData`.

Usage

```
getParams(
  sfe,
  name,
  local = FALSE,
  colData = FALSE,
  colGeometryName = NULL,
  annotGeometryName = NULL,
  reducedDimName = NULL
)
```

Arguments

sfe	A SpatialFeatureExperiment object.
name	Name used to store the results.
local	Logical, whether the results of interest come from a local spatial method.
colData	Logical, whether the results were computed for a column of colData(sfe).
colGeometryName	To get results for a colGeometry.
annotGeometryName	To get results for an annotGeometry; colGeometry has precedence so this argument is ignored if colGeometryName is specified.
reducedDimName	Name of a dimension reduction, can be seen in reducedDimNames. colGeometryName and annotGeometryName have precedence over reducedDimName.

Value

A named list showing the parameters

Examples

```
library(SFEData)
library(scater)
library(Voyager)
sfe <- McKellarMuscleData("small")
colGraph(sfe, "visium") <- findVisiumGraph(sfe)
sfe <- colDataMoransI(sfe, "nCounts")
getParams(sfe, "moran", colData = TRUE)
```

---

getPixelSize *Get physical size of pixels*

Description

This function gets physical size of pixels in each resolution of a OME-TIFF pyramid in BioFormatsImage.

Usage

```
getPixelSize(file, resolution = 1L)
```

Arguments

file	Path to an OME-TIFF file.
resolution	Which resolution to query; 1 means the highest resolution. The pixels will be larger for the lower resolutions.

Value

Numeric vector of length 2 of pixel size in x and y. Usually they're the same.

Examples

```
library(SFEData)
fp <- tempfile()
dir_use <- XeniumOutput("v1", file_path = fp)
# RBioFormats null pointer error
try(getPixelSize(file.path(dir_use, "morphology_focus.ome.tif")))
getPixelSize(file.path(dir_use, "morphology_focus.ome.tif"))
unlink(dir_use, recursive = TRUE)
```

---

getTechTxFields — *Get relevant fields and file paths for transcript spots*

Description

Get column names for x, y, and z coordinates, gene IDs, and cell IDs from the transcript file and get file paths for transcript spot coordinates given technology.

Usage

```
getTechTxFields(tech, data_dir = NULL)
```

Arguments

`tech`	Name of the commercial technology, must be one of Vizgen, Xenium, and CosMX.
`data_dir`	Top level directory of the output.

Value

A named list with elements:

`spatialCoordsNames` A character vector for column names for the xyz coordinates of the transcript spots.

`gene_col` Column name for gene IDs.

`cell_col` Column name for cell IDs.

`fn` File path of the transcript spot file.

---

imageIDs — *Show all image_ids in the SFE object*

Description

The title is self-explanatory. Some functions require `image_id` to get or set images.

Usage

```
imageIDs(sfe)
```

Arguments

sfe	A SpatialFeatureExperiment object.

Value

A character vector of image_ids.

Examples

```
fp <- system.file(file.path("extdata", "sample01"),
package = "SpatialFeatureExperiment")
sfe <- read10xVisiumSFE(fp, type = "sparse")
imageIDs(sfe)
```

Img<-,SpatialExperiment-method

Image setter

Description

Modify or replace images stored in a SpatialExperiment object. This is different from addImg which adds the image from files and can't replace existing images, which is there to be consistent with SpatialExperiment. This setter here can replace existing images with another object that inherits from VirtualSpatialImage, including SpatRasterImage, BioFormatsImage, and ExtImage.

Usage

```
## S4 replacement method for signature 'SpatialExperiment'
Img(x, sample_id = 1L, image_id, scale_fct = 1) <- value
```

Arguments

x	A SpatialExperiment object, which includes SFE.
sample_id	Which sample the image is associated with. Use sampleIDs to get sample IDs present in the SFE object.
image_id	Image ID, such as "lowres" and "hires" for Visium data and "DAPI" and "PolyT" for Vizgen MERFISH data.
scale_fct	Scale factor to convert pixels in lower resolution to those in the full resolution. Only relevant to image classes implemented in SpatialExperiment but not SpatialFeatureExperiment because the spatial extent of images in SFE takes precedence.
value	New version of image to add, must inherit from VirtualSpatialImage.

Value

SFE object with the new image added.

Examples

```
library(EBImage)
library(SFEData)
library(RBioFormats)
fp <- tempfile()
fn <- XeniumOutput("v2", file_path = fp)
# Weirdly the first time I get the null pointer error
try(sfe <- readXenium(fn))
sfe <- readXenium(fn)
img <- getImg(sfe) |> toExtImage(resolution = 1L)
img <- img[,,1] > 500
Img(sfe, image_id = "mask") <- img
imageIDs(sfe)
unlink(fn, recursive = TRUE)
```

---

imgRaster *Get the image from *Image class*

Description

In SFE, S4 classes inheriting from `VirtualSpatialImage` have been implemented to make these image classes compatible with `SpatialExperiment`.

Usage

```
## S4 method for signature 'SpatRasterImage'
imgRaster(x, maxcell = 1e+07, col = terra::map.pal("viridis", 100))

## S4 method for signature 'BioFormatsImage'
imgRaster(x, resolution = 4L)

## S4 method for signature 'ExtImage'
imgRaster(x)
```

Arguments

`x`	An object of class `*Image` as implemented in this package.
`maxcell`	positive integer. Maximum number of cells to use for the plot
`col`	vector of colors. The default is `map.pal("viridis", 100)`
`resolution`	Resolution to read in from OME-TIFF, defaults to 4, which is a medium resolution in Xenium.

Value

Since version 1.9.8, `imgRaster` will return an array of hex colors, or the raster object, as required by `SpatialExperiment`. This will break older SFE code calling `imgRaster`.

See Also

Other image methods: `SFE-image`, `affineImg()`, `cropImg()`, `dim,BioFormatsImage-method`, `dim,ExtImage-method`, `ext()`, `imgSource()`, `mirrorImg()`, `rotateImg()`, `scaleImg()`, `translateImg()`, `transposeImg()`

imgSource — *Source of images that are on disk*

Description

Get the file path of images that are on disk and not read into memory. Only applies to SpatRasterImage and BioFormatsImage.

Usage

```
## S4 method for signature 'SpatRasterImage'
imgSource(x)

## S4 method for signature 'BioFormatsImage'
imgSource(x)

## S4 method for signature 'ExtImage'
imgSource(x)
```

Arguments

x	An object of class *Image as implemented in this package.

Value

String, file path to the original image on disk. For SpatRasterImage, if the image is loaded into memory, then NULL.

See Also

Other image methods: SFE-image, affineImg(), cropImg(), dim,BioFormatsImage-method, dim,ExtImage-method, ext(), imgRaster(), mirrorImg(), rotateImg(), scaleImg(), translateImg(), transposeImg()

internal-Voyager — *Internal functions also used in Voyager*

Description

Not meant for the user, but exporting to be used internally in Voyager. But one day I may clean these up and remove the internal note for people building on top of SFE.

Usage

```
.value2df(value, use_geometry, feature = NULL)

.check_features(x, features, colGeometryName = NULL, swap_rownames = NULL)

.warn_symbol_duplicate(x, symbols, swap_rownames = "symbol")
```

```
.symbol2id(x, features, swap_rownames)

.check_sample_id(x, sample_id, one = TRUE, mustWork = TRUE)

.rm_empty_geometries(g, MARGIN)

.check_rg(type, x, sample_id)

.ext_(x)
```

Value

Internal

`listw2sparse` *Convert listw into sparse adjacency matrix*

Description

Edge weights are used in the adjacency matrix. Because most elements of the matrix are 0, using sparse matrix greatly reduces memory use.

Usage

```
listw2sparse(listw)
```

Arguments

`listw`	A `listw` object for spatial neighborhood graph.

Value

A sparse `dgCMatrix`, whose row represents each cell or spot and whose columns represent the neighbors. The matrix does not have to be symmetric. If `region.id` is present in the `listw` object, then it will be the row and column names of the output matrix.

Examples

```
library(SFEData)
sfe <- McKellarMuscleData("small")
g <- findVisiumGraph(sfe)
mat <- listw2sparse(g)
```

`localResults` *Get and set results from local spatial statistics*

Description

Local spatial statics like local Moran's I, local Geary's C, Getis-Ord Gi*, and geographically weighted summary statistics return values at each spatial location. Just like dimension reductions, these results are clearly associated with the broader SFE object, so they should have a place within the object. However, a separate field is needed because these analyses are conceptually distinct from dimension reduction. Also, each feature (e.g. gene) can have its own results with values at each location. The `localResults` field in the SFE object stores these results that has a value for each spatial location.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
localResults(
  x,
  sample_id = "all",
  name = "all",
  features = NULL,
  colGeometryName = NULL,
  annotGeometryName = NULL,
  withDimnames = TRUE,
  swap_rownames = NULL,
  ...
)

## S4 replacement method for signature 'SpatialFeatureExperiment'
localResults(
  x,
  sample_id = "all",
  name = "all",
  features = NULL,
  colGeometryName = NULL,
  annotGeometryName = NULL,
  withDimnames = TRUE,
  swap_rownames = NULL,
  ...
) <- value

## S4 method for signature 'SpatialFeatureExperiment'
localResultNames(x)

## S4 replacement method for signature 'SpatialFeatureExperiment,character'
localResultNames(x) <- value

## S4 method for signature 'SpatialFeatureExperiment'
localResultFeatures(
  x,
  type = 1L,
```

```
  colGeometryName = NULL,
  annotGeometryName = NULL,
  swap_rownames = NULL
)

## S4 method for signature 'SpatialFeatureExperiment'
localResultAttrs(
  x,
  type = 1L,
  feature,
  colGeometryName = NULL,
  annotGeometryName = NULL,
  swap_rownames = NULL
)

## S4 method for signature 'SpatialFeatureExperiment'
localResult(
  x,
  type = 1L,
  feature,
  colGeometryName = NULL,
  annotGeometryName = NULL,
  sample_id = 1L,
  withDimnames = TRUE,
  simplify = TRUE,
  swap_rownames = NULL
)

## S4 replacement method for signature 'SpatialFeatureExperiment'
localResult(
  x,
  type = 1L,
  feature,
  colGeometryName = NULL,
  annotGeometryName = NULL,
  sample_id = 1L,
  withDimnames = TRUE
) <- value
```

Arguments

`x`	A `SpatialFeatureExperiment` object.
`sample_id`	Sample ID to get or set geometries.
`name`	Name of the spatial method used, such as "localmoran".
`features`	Features whose local results to get or set, for `localResults` getter and setter for multiple features at a time.
`colGeometryName`	Which `colGeometry` to get or set local results.
`annotGeometryName`	Which `annotGeometry` to get or set local results.
`withDimnames`	Logical. If `TRUE`, then the dimnames (colnames or rownames) of the gene count matrix should correspond to row names of the `sf` data frames of interest.

swap_rownames	Name of a column in rowData to identify features instead of the row names of the SFE object. For example, if the row names of the SFE object are Ensembl IDs and gene symbols are in the "symbol" column in rowData, then putting "symbol" for this argument will use the gene symbols to identify which gene's local results to get or set.
...	Ignored
value	Values to set, should be either a matrix or a data frame.
type	Name or index of the spatial method used, such as "localmoran".
feature	Feature whose local results to get or set, for localResult getter and setter for one feature at a time.
simplify	Basically whether to return the content of the list rather than a list when the list only has one element, such as results for one type and one feature.

Value

localResults returns a named list each element of which is a set of local results of interest. localResult returns a matrix or a data frame, whichever the original is when it's set. localResultNames returns a character vector. Setters return an SFE object with the desired field set. For genes and colData columns, the local results are stored in the localResults field in int_colData, whereas for colGeometries and annotGeometries, the local results are stored as columns in the same sf data frames. localResultFeatures returns a character vector of names of features for which local results are available. localResultAttrs returns a character vector of the column names of the local results of one type for one feature. It returns NULL if the results are a vector.

Examples

```
# Toy example
sfe <- readRDS(system.file("extdata/sfe_toy.rds",
    package = "SpatialFeatureExperiment"
))
# localResults functions are written for organizing results from local
# spatial statistics (see the Voyager package). But for the examples here,
# random toy matrices are used. The real results are often matrices, with a
# matrix for each feature.
library(S4Vectors)
set.seed(29)
toy_res1 <- matrix(rnorm(10),
    nrow = 5, ncol = 2,
    dimnames = list(colnames(sfe), c("meow", "purr"))
)
toy_res1b <- matrix(rgamma(10, shape = 2),
    nrow = 5, ncol = 2,
    dimnames = list(colnames(sfe), c("meow", "purr"))
)
toy_df1 <- DataFrame(gene1 = I(toy_res1), gene2 = I(toy_res1b))

toy_res2 <- matrix(rpois(10, lambda = 2),
    nrow = 5, ncol = 2,
    dimnames = list(colnames(sfe), c("sassy", "tortitude"))
)
toy_df2 <- DataFrame(gene1 = I(toy_res2))
# Set all local results
localResults(sfe) <- list(localmoran = toy_df1, Gistar = toy_df2)
# Get all local results
```

```
lrs <- localResults(sfe)

# Set results of the same type for multiple genes
localResults(sfe, name = "localmoran") <- toy_df1
# Can also use a list
localResults(sfe, name = "localmoran") <- as.list(toy_df1)
# Get results of the same type for multiple genes
lrs <- localResults(sfe, name = "localmoran", features = c("gene1", "gene2"))

# Set results for one type and one gene
localResult(sfe, "localmoran", feature = "gene1") <- toy_res1
# Get results for one type and one gene
lr <- localResult(sfe, "localmoran", feature = "gene1")

# Set results for a feature in colGeometries
cg_toy <- readRDS(system.file("extdata/cg_toy.rds",
    package = "SpatialFeatureExperiment"
))
colGeometry(sfe, "cg") <- cg_toy
localResult(sfe, "localmoran",
    feature = "gene1",
    colGeometryName = "cg"
) <- toy_res1
# Get results for a feature in colGeometries
lr <- localResult(sfe, "localmoran", "gene1", colGeometryName = "cg")
```

---

mirrorImg *Mirror/flip images*

Description

Flip images along the middle horizontal or vertical axis.

Usage

```
## S4 method for signature 'SpatRasterImage'
mirrorImg(
  x,
  direction = c("vertical", "horizontal"),
  filename = "",
  maxcell = NULL,
  ...
)

## S4 method for signature 'BioFormatsImage'
mirrorImg(x, direction = c("vertical", "horizontal"), ...)

## S4 method for signature 'ExtImage'
mirrorImg(x, direction = c("vertical", "horizontal"), ...)
```

Arguments

x	SpatRaster or SpatVector

direction	character. Should (partially) match "vertical" to flip by rows, or "horizontal" to flip by columns
filename	character. Output filename
maxcell	Max number of pixels to load SpatRasterImage into memory. The default 1e7 is chosen because this is the approximate number of pixels in the medium resolution image at resolution = 4L in Xenium OME-TIFF to make different methods of this function consistent.
...	additional arguments for writing files as in writeRaster

Value

*Image object of the same class.

See Also

Other image methods: SFE-image, affineImg(), cropImg(), dim,BioFormatsImage-method, dim,ExtImage-method, ext(), imgRaster(), imgSource(), rotateImg(), scaleImg(), translateImg(), transposeImg()

---

multi_listw2sparse *Convert multiple listw graphs into a single sparse adjacency matrix*

Description

Each sample in the SFE object has a separate spatial neighborhood graph. Spatial analyses performed jointly on multiple samples require a combined spatial neighborhood graph from the different samples, where the different samples would be disconnected components of the graph. This combined adjacency matrix can be used in MULTISPATI PCA.

Usage

```
multi_listw2sparse(listws)
```

Arguments

listws	A list of listw objects.

Value

A sparse dgCMatrix of the combined spatial neighborhood graph, with the original spatial neighborhood graphs of the samples on the diagonal. When the input is an SFE object, the rows and columns will match the column names of the SFE object.

Examples

```
# example code
```

`read10xVisiumSFE` *Read 10X Visium data as SpatialFeatureExperiment*

Description

Read Space Ranger output from Visium v1 (not HD) as a SpatialFeatureExperiment object, where spots are represented with polygons in the colGeometry called "spotPoly". Other geometries can be added later after the dataset is read. If `data = "filtered"`, then spatial neighborhood graphs of the spots are also computed and stored in the colGraph called "visium" in all samples for downstream spatial analyses.

Usage

```
read10xVisiumSFE(
  samples = "",
  dirs = file.path(samples, "outs"),
  sample_id = paste0("sample", sprintf("%02d", seq_along(samples))),
  type = c("HDF5", "sparse"),
  data = c("filtered", "raw"),
  images = c("lowres", "hires"),
  unit = c("full_res_image_pixel", "micron"),
  style = "W",
  zero.policy = NULL,
  load = deprecated(),
  row.names = c("id", "symbol"),
  flip = c("geometry", "image", "none")
)
```

Arguments

`samples` A character vector containing one or more directory names, each corresponding to a 10X sample. Each directory should contain a matrix file, a gene/feature annotation file, and a barcode annotation file.

Alternatively, each string may contain a path to a HDF5 file in the sparse matrix format generated by 10X. These can be mixed with directory names when `type="auto"`.

Alternatively, each string may contain a prefix of names for the three-file system described above, where the rest of the name of each file follows the standard 10X output.

`dirs` Directory for each sample that contains the `spatial` and `raw/filtered_featues_bc_matrix` directories. By default, the `outs` directory under the directory specified in the `samples` argument, as in Space Ranger output. Change the `dirs` argument if you have moved or renamed the output directory.

`sample_id` Which sample(s) in the SFE object to use for the graph. Can also be "all", which means this function will compute the graph for all samples independently.

`type` Either "HDF5", and the matrix will be represented as `TENxMatrix`, or "sparse", and the matrix will be read as a `dgCMatrix`.

`data` character string specifying whether to read in filtered (spots mapped to tissue) or raw data (all spots).

`images`	character vector specifying which images to include. Valid values are `"lowres"`, `"hires"`, `"fullres"`, `"detected"`, `"aligned"`
`unit`	Whether to use pixels in full resolution image or microns as the unit. If using microns, then spacing between spots in pixels will be used to convert the coordinates into microns, as the spacing is known to be 100 microns. This is used to plot scale bar.
`style`	`style` can take values "W", "B", "C", "U", "minmax" and "S"
`zero.policy`	default NULL, use global option value; if FALSE stop with error for any empty neighbour sets, if TRUE permit the weights list to be formed with zero-length weights vectors
`load`	Deprecated. Not used, kept for backward compatibility for now.
`row.names`	String specifying whether to use Ensembl IDs ("ID") or gene symbols ("Symbol") as row names. If using symbols, the Ensembl ID will be appended to disambiguate in case the same symbol corresponds to multiple Ensembl IDs.
`flip`	Whether to flip the geometries or the images, because in `sf` and `terra`, the geometries use the Cartesian coordinates greater y coordinates going up, while in images, greater y values go down. Originally the Visium spots are in pixels in full res image. Either the image or the geometry needs to be flipped for them match in the Cartesian coordinate system.
`sample`	To be consistent with `SpatialExperiment::read10xVisium`, one or more directories with Space Ranger output for a Visium sample. It is assumed to have the `outs` directory in it but this can be overridden with the `dirs` argument.

Value

A SpatialFeatureExperiment object. The images might need to be manually transposed and/or mirrored to match the spots in this version of this package.

Note

It is assumed that the images have not been cropped. Otherwise the images might not align with the spots.

Examples

```
dir <- system.file("extdata", package = "SpatialFeatureExperiment")

sample_ids <- c("sample01", "sample02")
samples <- file.path(dir, sample_ids)

list.files(samples[1])
list.files(file.path(samples[1], "spatial"))
(sfe <- read10xVisiumSFE(samples, sample_id = sample_ids,
    type = "sparse", data = "filtered",
    load = FALSE
))
```

readCosMX *Read CosMX data into SFE*

Description

This function reads the standard CosMX output into an SFE object, as in "Basic Data Files" on the Nanostring website. For new version of CosMX, these files are the flat files in the AtoMX output.

Usage

```
readCosMX(
  data_dir,
  z = "all",
  sample_id = "sample01",
  min_area = NULL,
  add_molecules = FALSE,
  split_cell_comps = FALSE,
  BPPARAM = SerialParam(),
  file_out = file.path(data_dir, "tx_spots.parquet"),
  z_option = c("3d", "split")
)
```

Arguments

`data_dir` Top level output directory.

`z` Integer z index or "all" to indicate which z-planes to read for the transcript spots.

`sample_id` A `character` sample identifier, which matches the `sample_id` in `imgData`. The `sample_id` will also be stored in a new column in `colData`, if not already present. Default = `sample01`.

`min_area` Minimum cell area in square microns or pixel units (eg for CosMX). Anything smaller will be considered artifact or debris and removed. Default to 'NULL', ie no filtering of polygons.

`add_molecules` Logical, whether to add transcripts coordinates to an object.

`split_cell_comps` Logical, whether to split transcript spot geometries by cell compartment. Only relevant when 'add_molecules = TRUE'.

`BPPARAM` A `BiocParallelParam` object specifying parallel processing backend and number of threads to use for parallelizable tasks:

1. To load cell segmentation from HDF5 files from different fields of view (FOVs) with multiple cores. A progress bar can be configured in the `BiocParallelParam` object. When there are numerous FOVs, reading in the geometries can be time consuming, so we recommend using a server and larger number of threads. This argument is not used if `use_cellpose = TRUE` and the parquet file is present.
2. To get the largest piece and see if it's larger than `min_area` when there are multiple pieces in the cell segmentation for one cell.

`file_out` Name of file to save the geometry or raster to disk. Especially when the geometries are so large that it's unwieldy to load everything into memory. If this

file (or directory for multiple files) already exists, then the existing file(s) will be read, skipping the processing. When writing the file, extensions supplied are ignored and extensions are determined based on ‘dest‘.

`z_option` What to do with z coordinates. "3d" is to construct 3D geometries. "split" is to create a separate 2D geometry for each z-plane so geometric operations are fully supported but some data wrangling is required to perform 3D analyses. When the z coordinates are not integers, 3D geometries will always be constructed since there are no z-planes to speak of. This argument does not apply when ‘spatialCoordsNames‘ has length 2.

Value

An SFE object. Cell polygons are written to ‘cell_boundaries_sf.parquet‘ in ‘data_dir‘. If reading transcript spots (‘add_molecules = TRUE‘), then the reformatted transcript spots are saved to file specified in the ‘file_out‘ argument, which is by default ‘tx_spots.parquet‘ in the same directory as the rest of the data.

Examples

```
fp <- tempfile()
dir_use <- SFEData::CosMXOutput(file_path = fp)
sfe <- readCosMX(dir_use, z = "all", add_molecules = TRUE)
# Clean up
unlink(dir_use, recursive = TRUE)
```

---

readSelectTx *Read transcript spots of select genes*

Description

I speculate that in practice, the most common use of the transcript spots is visualization, and only a few genes can be visualized at a time or the spots will overcrowd. Then it doesn’t make sense to load the transcript spots of all genes into memory as they can take up a lot of memory. The function `readSelectTx` reads transcript spots of select genes into R, and the function `addSelectTx` adds them to `rowGeometries` of the SFE object.

Usage

```
readSelectTx(file, gene_select, z = "all", z_option = c("3d", "split"))

addSelectTx(
  sfe,
  file,
  gene_select,
  sample_id = 1L,
  z = "all",
  z_option = c("3d", "split"),
  swap_rownames = NULL
)
```

Arguments

file	File path of a GeoParquet file (e.g. already reformatted with the formatTxSpots or addTxSpots function, should have already flipped to match image if necessary).
gene_select	Character vector of a subset of genes. If NULL, then all genes that have transcript spots are added. Only relevant when reading data from formatted files on disk. If specified, then return = TRUE.
z	Index of z plane to read. Can be "all" to read all z-planes into MULTIPOINT geometries with XYZ coordinates. If z values are not integer, then spots with all z values will be read.
z_option	What to do with z coordinates. "3d" is to construct 3D geometries. "split" is to create a separate 2D geometry for each z-plane so geometric operations are fully supported but some data wrangling is required to perform 3D analyses. When the z coordinates are not integers, 3D geometries will always be constructed since there are no z-planes to speak of. This argument does not apply when ‘spatialCoordsNames‘ has length 2.
sfe	A ‘SpatialFeatureExperiment‘ object.
sample_id	Which sample in the SFE object the transcript spots should be added to.
swap_rownames	Name of a column in rowData(sfe) to use as gene identifiers in place of the actual row names. In some cases this may be needed to match each transcript spot MULTIPOINT geometry to rows of sfe.

Value

When there are multipel parquet files to be read, a list of sf data frames with MULTIPOINT geometry for genes selected. When there is only one file, then one sf data frame. For addSelectTx, an SFE object with the transcript spots of the selected genes added.

Note

The GDAL Parquet driver is required for this function, though not for other functions that work with GeoParquet files. GDAL Parquet driver has been supported since GDAL 3.5.0, but is not part of the default installation. The z and z_option arguments are there since the file names contain z-plane information when relevant. See the GDAL documentation page for the Parquet driver.

Examples

```
library(SFEData)
if (gdalParquetAvailable()) {
    fp <- tempfile()
    dir_use <- XeniumOutput("v2", file_path = fp)
    fn_tx <- formatTxTech(dir_use, tech = "Xenium", flip = TRUE, return = FALSE,
                          file_out = file.path(dir_use, "tx_spots.parquet"))
    gene_select <- c("ACE2", "BMX")
    df <- readSelectTx(fn_tx, gene_select)
    # RBioFormats null pointer error the first time
    try(sfe <- readXenium(dir_use))
    sfe <- readXenium(dir_use)
    sfe <- addSelectTx(sfe, fn_tx, head(rownames(sfe), 5), swap_rownames = "Symbol")
    unlink(dir_use, recursive = TRUE)
}
```

readVisiumHD — *Read Visium HD data*

Description

This function reads Visium HD Space Ranger output into R.

Usage

```
readVisiumHD(
  data_dir,
  bin_size = c(2L, 8L, 16L),
  sample_id = NULL,
  type = c("HDF5", "sparse"),
  data = c("filtered", "raw"),
  images = c("lowres", "hires"),
  unit = c("full_res_image_pixel", "micron"),
  style = "W",
  zero.policy = NULL,
  row.names = c("id", "symbol"),
  flip = c("geometry", "image"),
  add_graph = FALSE,
  rotate = FALSE
)
```

Arguments

`data_dir`	Directory
`bin_size`	One or more resolutions to load, must be 2, 8, or 16. Can be either integer or character.
`sample_id`	Which sample(s) in the SFE object to use for the graph. Can also be "all", which means this function will compute the graph for all samples independently.
`type`	Either "HDF5", and the matrix will be represented as `TENxMatrix`, or "sparse", and the matrix will be read as a `dgCMatrix`.
`data`	character string specifying whether to read in filtered (spots mapped to tissue) or raw data (all spots).
`images`	character vector specifying which images to include. Valid values are `"lowres"`, `"hires"`, `"fullres"`, `"detected"`, `"aligned"`
`unit`	Whether to use pixels in full resolution image or microns as the unit. If using microns, then spacing between spots in pixels will be used to convert the coordinates into microns, as the spacing is known to be 100 microns. This is used to plot scale bar.
`style`	`style` can take values “W”, “B”, “C”, “U”, “minmax” and “S”
`zero.policy`	default NULL, use global option value; if FALSE stop with error for any empty neighbour sets, if TRUE permit the weights list to be formed with zero-length weights vectors
`row.names`	String specifying whether to use Ensembl IDs ("ID") or gene symbols ("Symbol") as row names. If using symbols, the Ensembl ID will be appended to disambiguate in case the same symbol corresponds to multiple Ensembl IDs.

flip — Whether to flip the geometries or the images, because in sf and terra, the geometries use the Cartesian coordinates greater y coordinates going up, while in images, greater y values go down. Originally the Visium spots are in pixels in full res image. Either the image or the geometry needs to be flipped for them match in the Cartesian coordinate system.

add_graph — c(local), if to add spatial neighborhood graph for spots and only if c(data = "filtered"). Default is c(TRUE). This is optional because for larger datasets, the graph can take a while to compute.

rotate — Logical, whether to rotate the geometry, because usually the grid of spots is slightly, but just very slightly, rotated from the perfect horizontal line. The spots can be rotated so the square polygons are more accurate, because for computational efficiency, st_buffer is used to create the polygons which are not rotated. This is optional because the rotation is very slight.

Value

An SFE object if 'length(bin_size) == 1L', otherwise a list of SFE objects each element of which is for one bin size. They're not concatenated since it might not make sense to perform joint analyses on the different resolutions that benefit from having them in the same SFE object, unlike different biological replica. Here unlike in read10xVisiumSFE, the centroids geometry is also added because it will greatly facilitate plotting when there are many spots when not zooming in. See the scattermore argument in plotSpatialFeature.

Examples

```
#
```

---

readVizgen — *Read Vizgen MERFISH output as SpatialFeatureExperiment*

Description

This function reads the standard Vizgen MERFISH output into an SFE object. The coordinates are in microns. Cell centroids are read into colGeometry "centroids", and cell segmentations are read into colGeometry "cellSeg". The image(s) (polyT, DAPI, and cell boundaries) are also read as SpatRaster objects so they are not loaded into memory unless necessary. Because the image's origin is the top left while the geometry's origin is bottom left, either the image or the geometry needs to be flipped. Because the image accompanying MERFISH datasets are usually very large, the coordinates will be flipped so the flipping operation won't load the entire image into memory. Large datasets with hundreds of thousands of cells can take a while to read if reading transcript spots as it takes a while to convert the spots to MULTIPOINT geometries.

Usage

```
readVizgen(
  data_dir,
  z = "all",
  sample_id = "sample01",
  min_area = NULL,
  image = c("DAPI", "PolyT", "Cellbound"),
  flip = c("geometry", "image", "none"),
```

```
  max_flip = "50 MB",
  filter_counts = FALSE,
  add_molecules = FALSE,
  use_bboxes = FALSE,
  use_cellpose = TRUE,
  BPPARAM = SerialParam(),
  file_out = file.path(data_dir, "detected_transcripts.parquet"),
  z_option = c("3d", "split")
)
```

Arguments

`data_dir` Top level output directory.

`z` Integer, z index to read, or "all", indicating z-planes of the images and transcript spots to read. While cell segmentation seems to have multiple z-planes, the segmentation in all z-planes are the same so in effect the cell segmentatio is only in 2D.

`sample_id` A `character` sample identifier, which matches the `sample_id` in `imgData`. The `sample_id` will also be stored in a new column in `colData`, if not already present. Default = `sample01`.

`min_area` Minimum cell area in square microns or pixel units (eg for CosMX). Anything smaller will be considered artifact or debris and removed. Default to 'NULL', ie no filtering of polygons.

`image` Which image(s) to load, can be "DAPI", "PolyT", "Cellbound" or any combination of them.

`flip` To flip the image, geometry coordinates, or none. Because the image has the origin at the top left while the geometry has origin at the bottom left, one of them needs to be flipped for them to match. If one of them is already flipped, then use "none". The image will not be flipped if it's GeoTIFF.

`max_flip` Maximum size of the image allowed to flip the image. Because the image will be loaded into memory to be flipped. If the image is larger than this size then the coordinates will be flipped instead.

`filter_counts` Logical, whether to keep cells with counts > `0`.

`add_molecules` Logical, whether to add transcripts coordinates to an object.

`use_bboxes` If no segmentation output is present, use `cell_metadata` to make bounding boxes instead.

`use_cellpose` Whether to read the parquet files from CellPose cell segmentation. If `FALSE`, cell segmentation will be read from the HDF5 files. Note that reading HDF5 files for numerous FOVs is very slow.

`BPPARAM` A `BiocParallelParam` object specifying parallel processing backend and number of threads to use for parallelizable tasks:

1. To load cell segmentation from HDF5 files from different fields of view (FOVs) with multiple cores. A progress bar can be configured in the `BiocParallelParam` object. When there are numerous FOVs, reading in the geometries can be time consuming, so we recommend using a server and larger number of threads. This argument is not used if `use_cellpose = TRUE` and the parquet file is present.
2. To get the largest piece and see if it's larger than `min_area` when there are multiple pieces in the cell segmentation for one cell.

`file_out` Name of file to save the geometry or raster to disk. Especially when the geometries are so large that it's unwieldy to load everything into memory. If this file (or directory for multiple files) already exists, then the existing file(s) will be read, skipping the processing. When writing the file, extensions supplied are ignored and extensions are determined based on 'dest'.

`z_option` What to do with z coordinates. "3d" is to construct 3D geometries. "split" is to create a separate 2D geometry for each z-plane so geometric operations are fully supported but some data wrangling is required to perform 3D analyses. When the z coordinates are not integers, 3D geometries will always be constructed since there are no z-planes to speak of. This argument does not apply when 'spatialCoordsNames' has length 2.

Value

A `SpatialFeatureExperiment` object.

Note

Since the transcript spots file is often very large, we recommend only using `add_molecules = TRUE` on servers with a lot of memory. If reading all z-planes, conversion of transcript spot geometry to parquet file might fail due to arrow data length limit. In a future version, when the transcript spot geometry is large, it will be written to multiple separate parquet files which are then concatenated with DuckDB. Also, in a future version, the transcript spot processing function might be rewritten in C++ to stream the original CSV file so it's not entirely loaded into memory.

Examples

```
fp <- tempfile()
dir_use <- SFEData::VizgenOutput(file_path = fp)
sfe <- readVizgen(dir_use, z = 3L, image = "PolyT",
flip = "geometry")

## Filtering of counts, and addition of molecule coordinates..
sfe <- readVizgen(dir_use, z = 3L, image = "PolyT", filter_counts = TRUE,
add_molecules = TRUE, flip = "geometry")

unlink(dir_use, recursive = TRUE)
```

---

readXenium *Read 10X Xenium output as SpatialFeatureExperiment*

Description

This function reads the standard 10X Xenium output into an SFE object.

Usage

```
readXenium(
  data_dir,
  sample_id = "sample01",
  min_area = NULL,
  image = c("morphology_focus", "morphology_mip"),
```

```
  segmentations = c("cell", "nucleus"),
  row.names = c("id", "symbol"),
  flip = c("geometry", "image", "none"),
  max_flip = "50 MB",
  filter_counts = FALSE,
  add_molecules = FALSE,
  min_phred = 20,
  BPPARAM = SerialParam(),
  file_out = file.path(data_dir, "tx_spots.parquet")
)
```

Arguments

`data_dir` Top level output directory.

`sample_id` A `character` sample identifier, which matches the `sample_id` in `imgData`. The `sample_id` will also be stored in a new column in `colData`, if not already present. Default = `sample01`.

`min_area` Minimum cell area in square microns or pixel units (eg for CosMX). Anything smaller will be considered artifact or debris and removed. Default to 'NULL', ie no filtering of polygons.

`image` Which image(s) to load, can be "morphology_mip", "morphology_focus" or both. Note that in Xenium Onboarding Analysis (XOA) v2, there is no longer "morphology_mip" and "morphology_focus" is a directory with 4 images corresponding to 4 channels: DAPI, "Cadherin", 18S, and Vimentin. So this argument is ignored for XOA v2.

`segmentations` Which segmentation outputs to read, can be "cell", "nucleus", or both.

`row.names` String specifying whether to use Ensembl IDs ("id") or gene symbols ("symbol") as row names. If using symbols, the Ensembl ID will be appended to disambiguate in case the same symbol corresponds to multiple Ensembl IDs. Always "symbol" if 'add_molecules = TRUE' because only gene symbols are used in the transcript spot files.

`flip` To flip the image, geometry coordinates, or none. Because the image has the origin at the top left while the geometry has origin at the bottom left, one of them needs to be flipped for them to match. If one of them is already flipped, then use "none". The image will not be flipped if it's GeoTIFF.

`max_flip` Maximum size of the image allowed to flip the image. Because the image will be loaded into memory to be flipped. If the image is larger than this size then the coordinates will be flipped instead.

`filter_counts` Logical, whether to keep cells with counts > `0`.

`add_molecules` Logical, whether to add transcripts coordinates to an object.

`min_phred` Minimum Phred score to keep spot. By default 20, the conventional threshold indicating "acceptable", meaning that there's 1 chance that the spot was decoded in error.

`BPPARAM` A `BiocParallelParam` object specifying parallel processing backend and number of threads to use for parallelizable tasks:

1. To load cell segmentation from HDF5 files from different fields of view (FOVs) with multiple cores. A progress bar can be configured in the `BiocParallelParam` object. When there are numerous FOVs, reading in the geometries can be time consuming, so we recommend using a server and larger number of

threads. This argument is not used if `use_cellpose = TRUE` and the parquet file is present.

2. To get the largest piece and see if it's larger than `min_area` when there are multiple pieces in the cell segmentation for one cell.

`file_out` Name of file to save the geometry or raster to disk. Especially when the geometries are so large that it's unwieldy to load everything into memory. If this file (or directory for multiple files) already exists, then the existing file(s) will be read, skipping the processing. When writing the file, extensions supplied are ignored and extensions are determined based on 'dest'.

Value

An SFE object. If reading segmentations, the cell or nuclei segmentation will be saved to 'cell_boundaries_sf.parquet' and 'nucleus_boundaries_sf.parquet' respectively in 'data.dir' so next time the boundaries can be read much more quickly. If reading transcript spots ('add_molecules = TRUE'), then the reformatted transcript spots are saved to file specified in the 'file_out' argument, which is by default 'tx_spots.parquet' in the same directory as the rest of the data. If images are present, then the images will be of the `BioFormatsImage` class and not loaded into memory until necessary in later operations.

Note

Sometimes when reading images, you will see this error the first time: 'java.lang.NullPointerException: Cannot invoke "loci.formats.DimensionSwapper.setMetadataFiltered(boolean)" because "RBioFormats.reader" is null'. See this issue https://github.com/aoles/RBioFormats/issues/42 Rerun the code and it should work the second time.

Examples

```
library(SFEData)
library(RBioFormats)
fp <- tempfile()
dir_use <- XeniumOutput("v2", file_path = fp)
# RBioFormats issue
try(sfe <- readXenium(dir_use, add_molecules = TRUE))
sfe <- readXenium(dir_use, add_molecules = TRUE)
unlink(dir_use, recursive = TRUE)
```

`reexports` *Functions re-exported from other packages*

Description

These are some commonly used getters and setters of classes that SFE inherits so you don't have to separately attach those packages to use these functions.

Usage

```
colData(x, ...)

rowData(x, use.names = TRUE, ...)
```

```
colData(x, ...) <- value

spatialCoords(x, ...)

spatialCoords(x) <- value

spatialCoordsNames(x)

getImg(x, ...)

imgData(x)

rmvImg(x, ...)

counts(object, ...)

logcounts(object, ...)

reducedDim(x, type, ...)
```

Arguments

x	A SummarizedExperiment object or derivative.
...	For assay, arguments in ... are forwarded to assays. For rbind, cbind, ... contains SummarizedExperiment objects (or derivatives) to be combined. For other accessors, ignored.
use.names	For rowData: Like mcols(x), by default rowData(x) propagates the rownames of x to the returned DataFrame object (note that for a SummarizedExperiment object or derivative, the rownames are also the names i.e. rownames(x) is always the same as names(x)). Setting use.names=FALSE suppresses this propagation i.e. it returns a DataFrame object with no rownames. Use this when rowData(x) fails, which can happen when the rownames contain NAs (because the rownames of a SummarizedExperiment object or derivative can contain NAs, but the rownames of a DataFrame object cannot). For combineRows and combineCols: See Combining section below.
value	An object of a class specified in the S4 method signature or as outlined in 'Details'.
object	A SingleCellExperiment object, which includes SFE.
type	Name or numeric index to indicate which reducedDim to get, such as "PCA". By default the first item in reducedDims.

---

removeEmptySpace *Remove empty space*

Description

For each sample independently, all geometries and spatialCoords are translated so the origin is at the minimum coordinates of the bounding box of all geometries of the sample. This way coordinates of different samples will be more comparable. This removes empty space in the images if present.

Usage

```
removeEmptySpace(sfe, sample_id = "all")
```

Arguments

sfe	An SFE object.
sample_id	Sample to remove empty space.

Value

An SFE object with empty space removed.

Note

Unlike other functions in this package, this function operates on all samples by default.

Examples

```
library(SFEData)
library(SingleCellExperiment)
sfe <- McKellarMuscleData("full")
# Only keep spots on tissue
sfe <- sfe[, colData(sfe)$in_tissue]
# Move the coordinates of the tissue
sfe <- removeEmptySpace(sfe)
```

---

rotateImg *Rotate image*

Description

As in `SpatialExperiment`, rotation here must be a multiple of 90 degrees.

Usage

```
## S4 method for signature 'SpatRasterImage'
rotateImg(x, degrees, maxcell = 1e+07, ...)

## S4 method for signature 'BioFormatsImage'
rotateImg(x, degrees, ...)

## S4 method for signature 'ExtImage'
rotateImg(x, degrees, ...)
```

Arguments

x	An object of class `*Image` as implemented in this package.
degrees	How many degrees to rotate. Positive number means clockwise and negative number means counterclockwise.

maxcell	Max number of pixels to load SpatRasterImage into memory. The default 1e7 is chosen because this is the approximate number of pixels in the medium resolution image at resolution = 4L in Xenium OME-TIFF to make different methods of this function consistent.
...	Ignored. It's there so different methods can all be passed to the same lapply in the method for SFE objects. Some methods have extra arguments.

Value

SpatRasterImage will be loaded into memory and converted to ExtImage. Otherwise *Image object of the same class.

See Also

Other image methods: SFE-image, affineImg(), cropImg(), dim,BioFormatsImage-method, dim,ExtImage-method, ext(), imgRaster(), imgSource(), mirrorImg(), scaleImg(), translateImg(), transposeImg()

---

rowGeometries *Row geometry getters and setters*

Description

rowGeometries are geometries that corresponding to rows of the gene count matrix, such as sm-FISH transcript spots. The txSpots() function is a convenience wrapper for transcript spots, although this entirely depends on the rowGeometry being named txSpots.

Usage

```
rowGeometry(x, type = 1L, sample_id = 1L, withDimnames = TRUE)

rowGeometry(
  x,
  type = 1L,
  sample_id = 1L,
  withDimnames = TRUE,
  partial = FALSE,
  translate = TRUE
) <- value

rowGeometries(x, sample_id = "all", withDimnames = TRUE)

rowGeometries(
  x,
  sample_id = "all",
  withDimnames = TRUE,
  partial = FALSE,
  translate = TRUE
) <- value

rowGeometryNames(x)
```

```
rowGeometryNames(x) <- value

txSpots(x, sample_id = 1L, withDimnames = TRUE)

txSpots(
  x,
  sample_id = 1L,
  withDimnames = TRUE,
  partial = FALSE,
  translate = TRUE
) <- value
```

Arguments

x	A SpatialFeatureExperiment object.
type	An integer specifying the index or string specifying the name of the *Geometry to query or replace. If missing, then the first item in the *Geometries will be returned or replaced.
sample_id	Sample ID to get or set geometries.
withDimnames	Logical. If TRUE, then the dimnames (colnames or rownames) of the gene count matrix should correspond to row names of the sf data frames of interest.
partial	In setters, if a rowGeometry of the same name exists, whether to only replace the rows present in value.
translate	Logical. Only used if removeEmptySpace has been run of the SFE object. If that's the case, this argument indicates whether the new value to be assigned to the geometry is in the coordinates prior to removal of empty space so it should be translated to match the new coordinates after removing empty space. Default to TRUE.
value	Value to set. For dimGeometry, must be a sf data frame with the same number of rows as size in the dimension of interest, or an ordinary data frame that can be converted to such a sf data frame (see df2sf). For dimGeometries, must be a list of such sf or ordinary data frames.

Details

When there are multiple samples in the SFE object, rowGeometries for each sample has the sample_id appended to the name of the geometry. For example, if the name is txSpots and the sample ID is sample01, then the actual name of the rowGeometry is txSpots_sample01. In the getter, one can still specify rowGeometry(sfe, "txSpots", sample_id = "sample01").

Appending the sample_id is unnecessary when there is only one sample, but sample_id will be appended when to SFE objects are combined with cbind. It is necessary to distinguish bewteen different samples because they can have overlapping coordinate values.

See Also

[dimGeometries()], [colGeometries()]

Examples

```
library(SFEData)
library(RBioFormats)
fp <- tempfile()
dir_use <- XeniumOutput("v2", file_path = fp)
# RBioFormats issue
try(sfe <- readXenium(dir_use, add_molecules = TRUE))
sfe <- readXenium(dir_use, add_molecules = TRUE)
rowGeometries(sfe)
rowGeometryNames(sfe)
tx <- rowGeometry(sfe, "txSpots")
txSpots(sfe)
unlink(dir_use, recursive = TRUE)
```

---

sampleIDs *Get all unique sample IDs*

Description

The title is self-explanatory.

Usage

```
sampleIDs(sfe)
```

Arguments

sfe	A SpatialFeatureExperiment object.

Value

A character vector of all unique entries of the sample_id column in colData(x).

Examples

```
library(SFEData)
sfe <- McKellarMuscleData(dataset = "small")
sampleIDs(sfe)
```

---

saveRDS,SpatialFeatureExperiment-method *Save SpatialFeatureExperiment as RDS file*

Description

Saving SFE objects as RDS files is complicated by the SpatRaster class of the images. If present, the images need to be wrapped with the wrap function in terra before serializing the SFE object. Otherwise the images will be invalid pointers when the RDS is reloaded. If the image does not fit in memory and its file source is unknown, then it will be written to a temporary file, which is reloaded when the RDS file is loaded. When an SFE object with images is read from an RDS file, the images will not be unwrapped until necessary.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
saveRDS(
  object,
  file = "",
  ascii = FALSE,
  version = NULL,
  compress = TRUE,
  refhook = NULL
)
```

Arguments

object	A SpatialFeatureExperiment object.
file	a connection or the name of the file where the R object is saved to or read from.
ascii	a logical. If TRUE or NA, an ASCII representation is written; otherwise (default), a binary one is used. See the comments in the help for save.
version	the workspace format version to use. NULL specifies the current default version (3). The only other supported value is 2, the default from R 1.4.0 to R 3.5.0.
compress	a logical specifying whether saving to a named file is to use "gzip" compression, or one of "gzip", "bzip2" or "xz" to indicate the type of compression to be used. Ignored if file is a connection.
refhook	a hook function for handling reference objects.

Value

Invisibly NULL.

Examples

```
outdir <- system.file("extdata", package = "SpatialFeatureExperiment")
samples <- file.path(outdir, paste0("sample0", 1:2))
sfe <- read10xVisiumSFE(samples, type = "sparse", data = "filtered")
saveRDS(sfe, "foo.rds")
# Clean up
file.remove("foo.rds")
```

---

scaleImg *Scale image*

Description

This function scales the image about its center. After scaling, the center of the image is not shifted.

Usage

```
## S4 method for signature 'AlignedSpatialImage'
scaleImg(x, factor, ...)

## S4 method for signature 'BioFormatsImage'
scaleImg(x, factor, ...)
```

Arguments

x	An object of class *Image as implemented in this package.
factor	Numeric, scaling factor.
...	Ignored. It's there so different methods can all be passed to the same lapply in the method for SFE objects. Some methods have extra arguments.

Value

A *Image object of the same class that has been scaled. Behind the scene, it's only the extent that has been changed and the images are not changed. The center of the image is unchanged.

See Also

Other image methods: SFE-image, affineImg(), cropImg(), dim,BioFormatsImage-method, dim,ExtImage-method, ext(), imgRaster(), imgSource(), mirrorImg(), rotateImg(), translateImg(), transposeImg()

SFE-image *Methods for handling image-related data*

Description

Generics of these functions are defined in SpatialExperiment, except for transposeImg. These SFE methods cater to the new image-related classes in SFE. The SPE method for getImg, rmvImg, and imgRaster don't need to be modified for SFE and are hence not implemented here, but are simply re-exported.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
addImg(x, imageSource, sample_id = 1L, image_id, extent = NULL, scale_fct = 1)

## S4 method for signature 'SpatialFeatureExperiment'
transposeImg(
  x,
  sample_id = 1L,
  image_id = NULL,
  maxcell = 1e+07,
  filename = ""
)

## S4 method for signature 'SpatialFeatureExperiment'
mirrorImg(
  x,
  sample_id = 1L,
  image_id = NULL,
  direction = "vertical",
  maxcell = 1e+07,
  filename = ""
)
```

```
## S4 method for signature 'SpatialFeatureExperiment'
rotateImg(x, sample_id = 1L, image_id = NULL, degrees, maxcell = 1e+07)

## S4 method for signature 'SpatialFeatureExperiment'
translateImg(x, sample_id = 1L, image_id = NULL, v)

## S4 method for signature 'SpatialFeatureExperiment'
scaleImg(x, sample_id = 1L, image_id = NULL, factor)

## S4 method for signature 'SpatialFeatureExperiment'
affineImg(x, sample_id = 1L, image_id = NULL, M, v)
```

Arguments

x	A SFE object.
imageSource	a character string specifying an image file name (.png, .jpg or .tif) or URL to source the image from
sample_id	Which sample the image is associated with. Use sampleIDs to get sample IDs present in the SFE object.
image_id	Image ID, such as "lowres" and "hires" for Visium data and "DAPI" and "PolyT" for Vizgen MERFISH data.
extent	A numeric vector of length 4 with names of the set xmin, ymin, xmax, and ymax, specifying the extent of the image.
scale_fct	Scale factor – multiply pixel coordinates in full resolution image by this scale factor should yield pixel coordinates in a different resolution. extent takes precedence over scale_fct.
maxcell	Max number of pixels to load SpatRasterImage into memory. The default 1e7 is chosen because this is the approximate number of pixels in the medium resolution image at resolution = 4L in Xenium OME-TIFF to make different methods of this function consistent.
filename	character. Output filename
direction	character. Should (partially) match "vertical" to flip by rows, or "horizontal" to flip by columns
degrees	How many degrees to rotate. Positive number means clockwise and negative number means counterclockwise.
v	A numeric vector of length 2 specifying the vector in the xy plane to translate the SFE object.
factor	Numeric, scaling factor.
M	A 2x2 numeric matrix for the linear transformation in the xy plane.

Details

Method of transposeImg, mirrorImg, and rotateImg perform the method on all images within the SFE object that are specified with sample_id and image_id. For images that are not loaded into memory, rotateImg will load SpatRasterImage into memory and all image operations except translate will load BioFormatsImage into memory.

Note

If the image is already a GeoTIFF file that already has an extent, then the extent associated with the file will be honored and the `extent` and `scale_fct` arguments are ignored. Transposing the image is just like transposing a matrix. It's flipped about the line going from the top left to the bottom right.

See Also

Other image methods: `affineImg()`, `cropImg()`, `dim,BioFormatsImage-method`, `dim,ExtImage-method`, `ext()`, `imgRaster()`, `imgSource()`, `mirrorImg()`, `rotateImg()`, `scaleImg()`, `translateImg()`, `transposeImg()`

Examples

```
library(SFEData)
sfe <- McKellarMuscleData("small")
img_path <- system.file(file.path("extdata", "sample01", "outs", "spatial",
"tissue_lowres_image.png"), package = "SpatialFeatureExperiment")
sfe <- addImg(sfe, img_path, sample_id = "Vis5A", image_id = "lowres", scale_fct =
0.023)
img <- getImg(sfe)
# SpatRasterImage method
img_t <- transposeImg(img)
# SFE method
sfe <- transposeImg(sfe, sample_id = "Vis5A", image_id = "lowres")
```

SFE-transform — *Affine transfortaion of SFE object in histological space*

Description

These functions perform affine transformations on SFE objects, including all geometries and images. The transformation is performed on each sample as a whole. This differs from functions such as `mirrorImg` in that `mirrorImg` and `rotateImg` transform the image with the center at the center of the image itself. In contrast, the center of transformation here is the center of the bounding box of the entire sample, including images.

Usage

```
transpose(sfe, sample_id = "all", maxcell = NULL, filename = "")

mirror(
  sfe,
  sample_id = "all",
  direction = c("vertical", "horizontal"),
  maxcell = NULL,
  filename = ""
)

rotate(sfe, sample_id = "all", degrees, maxcell = 1e+07)

translate(sfe, sample_id = "all", v)
```

```
scale(sfe, sample_id = "all", factor)

affine(sfe, sample_id = "all", M, v, maxcell = 1e+07)
```

Arguments

sfe	An SFE object.
sample_id	Sample(s) to transform.
maxcell	Rotating SpatRasterImage will convert it into ExtImage, loading the image into memory. This argument specifies the maximum number of pixels in the image loaded into memory. The image will be down sampled to approximately this number of pixels.
filename	character. Output filename
direction	character. Should (partially) match "vertical" to flip by rows, or "horizontal" to flip by columns
degrees	How many degrees to rotate. Positive number means clockwise and negative number means counterclockwise.
v	Vector to spatially translate the SFE object.
factor	Numeric, scaling factor.
M	A 2x2 numeric matrix for the linear transformation in the xy plane.

Details

For images that are not loaded into memory, rotateImg will load SpatRasterImage into memory and all image operations except translate will load BioFormatsImage into memory.

Value

An SFE object with the sample(s) transformed.

Examples

```
library(SFEData)
sfe <- McKellarMuscleData("small")
sfe2 <- transpose(sfe)
sfe3 <- mirror(sfe)
```

---

show,SpatialFeatureExperiment-method

Print method for SpatialFeatureExperiment

Description

Printing summaries of colGeometries, rowGeometries, and annotGeometries in addition to what's shown for SpatialExperiment. Geometry names and types are printed.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
show(object)
```

Arguments

object	A SpatialFeatureExperiment object.

Value

None (invisible NULL).

Examples

```
library(SFEData)
sfe <- McKellarMuscleData(dataset = "small")
sfe # The show method is implicitly called
```

SpatialFeatureExperiment

Constructor of SpatialFeatureExperiment object

Description

Create a SpatialFeatureExperiment object.

Usage

```
SpatialFeatureExperiment(
  assays,
  colData = DataFrame(),
  rowData = NULL,
  sample_id = "sample01",
  spatialCoordsNames = c("x", "y"),
  spatialCoords = NULL,
  colGeometries = NULL,
  rowGeometries = NULL,
  annotGeometries = NULL,
  spotDiameter = NA_real_,
  annotGeometryType = "POLYGON",
  spatialGraphs = NULL,
  unit = c("full_res_image_pixel", "micron"),
  ...
)
```

Arguments

`assays` A `list` or `SimpleList` of matrix-like elements, or a matrix-like object (e.g. an ordinary matrix, a data frame, a DataFrame object from the **S4Vectors** package, a SparseMatrix derivative from the **SparseArray** package, a sparseMatrix derivative from the **Matrix** package, a DelayedMatrix object from the **DelayedArray** package, etc...). All elements of the list must have the same dimensions, and dimension names (if present) must be consistent across elements and with the row names of `rowRanges` and `colData`.

`colData` An optional DataFrame describing the samples. Row names on `colData`, if present, become the column names of the returned object.

`rowData` `NULL` (the default) or a DataFrame object describing the rows. Row names, if present, become the row names of the constructed SummarizedExperiment object. The number of rows of the DataFrame must equal the number of rows of the matrices in `assays`.

`sample_id` A `character` sample identifier, which matches the `sample_id` in `imgData`. The `sample_id` will also be stored in a new column in `colData`, if not already present. Default = `sample01`.

`spatialCoordsNames` A `character` vector of column names if `*Geometries` arguments have ordinary data frames, to identify the columns in the ordinary data frames that specify the spatial coordinates. If `colGeometries` is not specified, then this argument will behave as in `SpatialExperiment`, but `colGeometries` will be given precedence if provided.

`spatialCoords` A numeric matrix containing columns of spatial coordinates, as in `SpatialExperiment`. The coordinates are centroids of the entities represented by the columns of the gene count matrix. If `colGeometries` is also specified, then it will be given priority and a warning is issued. Otherwise, the `sf` representation of the centroids will be stored in the `colGeometry` called `centroids`.

`colGeometries` Geometry of the entities that correspond to the columns of the gene count matrix, such as cells and Visium spots. It must be a named list of one of the following:

An `sf` data frame The geometry column specifies the geometry of the entities.

An ordinary data frame specifying centroids Column names for the coordinates are specified in the `spatialCoordsNames` argument. For Visium and ST, in addition to the centroid coordinate data frame, the spot diameter in the same unit as the coordinates can be specified in the `spotDiamter` argument.

An ordinary data frame specifying polygons Also use `spatialCoordsNames`. There should an additional column "ID" to specify which vertices belong to which polygon. The coordinates should not be in list columns. Rather, the data frame should look like it is passed to `ggplot2::geom_polygon`. If there are holes, then there must also be a column "subID" that differentiates between the outer polygon and the holes.

In all cases, the data frame should specify the same number of geometries as the number of columns in the gene count matrix. If the column "barcode" is present, then it will be matched to column names of the gene count matrix. Otherwise, the geometries are assumed to be in the same order as columns in the gene count matrix. If the geometries are specified in an ordinary data frame, then it will be converted into `sf` internally. Named list of data frames because each entity can have multiple geometries, such as whole cell and nuclei segmentations. The

geometries are assumed to be POINTs for centroids and POLYGONs for segmentations. If polygons are specified in an ordinary data frame, then anything with fewer than 3 vertices will be removed. For anything other than POINTs, attributes of the geometry will be ignored.

`rowGeometries` Geometry associated with genes or features, which correspond to rows of the gene count matrix.

`annotGeometries`
Geometry of entities that do not correspond to columns or rows of the gene count matrix, such as tissue boundary and pathologist annotations of histological regions, and nuclei segmentation in a Visium dataset. Also a named list as in `colGeometries`. The ordinary data frame may specify POINTs, POLYGONs, or LINESTRINGs, or their MULTI versions. Each data frame can only specify one type of geometry. For MULTI versions, there must be a column "group" to identify each MULTI geometry.

`spotDiameter` Spot diameter for technologies with arrays of spots of fixed diameter per slide, such as Visium, ST, DBiT-seq, and slide-seq. The diameter must be in the same unit as the coordinates in the *Geometry arguments. Ignored for geometries that are not POINT or MULTIPOINT.

`annotGeometryType`
Character vector specifying geometry type of each element of the list if `annotGeometry` is specified. Each element of the vector must be one of POINT, LINESTRING, POLYGON, MULTIPOINT, MULTILINESTRING, and MULTIPOLYGON. Must be either length 1 (same for all elements of the list) or the same length as the list. Ignored if the corresponding element is an `sf` object.

`spatialGraphs` A named list of `listw` objects (see `spdep`) for spatial neighborhood graphs.

`unit` Unit the coordinates are in, either microns or pixels in full resolution image.

`...` Additional arguments passed to the `SpatialExperiment` and `SingleCellExperiment` constructors.

Value

A SFE object. If neither `colGeometries` nor `spotDiameter` is specified, then a `colGeometry` called "centroids" will be made, which is essentially the spatial coordinates as sf POINTs. If `spotDiameter` is specified, but not `colGeometries`, then the spatial coordinates will be buffered by half the diameter to get spots with the desired diameter, and the resulting `colGeometry` will be called "spotPoly", for which there's a convenience getter and setter, `spotPoly`.

Examples

```
library(Matrix)
data("visium_row_col")
coords1 <- visium_row_col[visium_row_col$col < 6 & visium_row_col$row < 6, ]
coords1$row <- coords1$row * sqrt(3)
cg <- df2sf(coords1[, c("col", "row")], c("col", "row"), spotDiameter = 0.7)

set.seed(29)
col_inds <- sample(seq_len(13), 13)
row_inds <- sample(seq_len(5), 13, replace = TRUE)
values <- sample(seq_len(5), 13, replace = TRUE)
mat <- sparseMatrix(i = row_inds, j = col_inds, x = values)
colnames(mat) <- coords1$barcode
rownames(mat) <- sample(LETTERS, 5)
```

```
rownames(cg) <- colnames(mat)

sfe <- SpatialFeatureExperiment(list(counts = mat),
    colData = coords1,
    spatialCoordsNames = c("col", "row"),
    spotDiameter = 0.7
)
sfe2 <- SpatialFeatureExperiment(list(counts = mat),
    colGeometries = list(foo = cg)
)
```

SpatialFeatureExperiment-class

The SpatialFeatureExperiment class

Description

This class inherits from the SpatialExperiment (SPE) class, which in turn inherits from SingleCellExperiment (SCE). SpatialFeatureExperiment stores geometries of spots or cells in sf objects which form columns of a DataFrame which is in turn a column of the int_colData DataFrame of the underlying SCE object, just like reducedDim in SCE. Geometries of the tissue outline, pathologist annotations, and objects (e.g. nuclei segmentation in a Visium dataset) are stored in sf objects in a named list called annotGeometries in int_metadata.

SpatialFeatureExperiment-coercion

SpatialFeatureExperiment coercion methods

Description

The SpatialFeatureExperiment class inherits from SpatialExperiment, which in turn inherits from SingleCellExperiment. A SpatialExperiment object with geometries in colGeometries in the int_colData, rowGeometries in the int_elementMetadata, or annotGeometries in the int_metadata can be directly converted to SpatialFeatureExperiment with as(spe,"SpatialFeatureExperiment" A SpatialExperiment object without the geometries can also be converted; the coordinates in the spatialCoords field will be used to make POINT geometries named "centroids" to add to colGeometries. The geometries can also be supplied separately when using toSpatialFeatureExperiment. Images are converted to SpatRaster.

Usage

```
## S4 method for signature 'SpatialExperiment'
toSpatialFeatureExperiment(
  x,
  colGeometries = NULL,
  rowGeometries = NULL,
  annotGeometries = NULL,
  spatialCoordsNames = c("x", "y"),
  annotGeometryType = "POLYGON",
  spatialGraphs = NULL,
```

```
  spotDiameter = NA,
  unit = NULL
)

## S4 method for signature 'SingleCellExperiment'
toSpatialFeatureExperiment(
  x,
  sample_id = "sample01",
  spatialCoordsNames = c("x", "y"),
  spatialCoords = NULL,
  colGeometries = NULL,
  rowGeometries = NULL,
  annotGeometries = NULL,
  annotGeometryType = "POLYGON",
  spatialGraphs = NULL,
  spotDiameter = NA,
  scaleFactors = 1,
  imageSources = NULL,
  image_id = NULL,
  loadImage = TRUE,
  imgData = NULL,
  unit = NULL
)

## S4 method for signature 'Seurat'
toSpatialFeatureExperiment(
  x,
  add_molecules = TRUE,
  flip = c("geometry", "image", "none"),
  image_scalefactors = c("lowres", "hires"),
  unit = NULL,
  BPPARAM = SerialParam()
)
```

Arguments

`x` A `SpatialExperiment` or `Seurat` object to be coerced to a `SpatialFeatureExperiment` object.

`colGeometries` Geometry of the entities that correspond to the columns of the gene count matrix, such as cells and Visium spots. It must be a named list of one of the following:

An `sf` data frame The geometry column specifies the geometry of the entities.

An ordinary data frame specifying centroids Column names for the coordinates are specified in the `spatialCoordsNames` argument. For Visium and ST, in addition to the centroid coordinate data frame, the spot diameter in the same unit as the coordinates can be specified in the `spotDiamter` argument.

An ordinary data frame specifying polygons Also use `spatialCoordsNames`. There should an additional column "ID" to specify which vertices belong to which polygon. The coordinates should not be in list columns. Rather, the data frame should look like it is passed to `ggplot2::geom_polygon`. If there are holes, then there must also be a column "subID" that differentiates between the outer polygon and the holes.

In all cases, the data frame should specify the same number of geometries as the number of columns in the gene count matrix. If the column "barcode" is present, then it will be matched to column names of the gene count matrix. Otherwise, the geometries are assumed to be in the same order as columns in the gene count matrix. If the geometries are specified in an ordinary data frame, then it will be converted into `sf` internally. Named list of data frames because each entity can have multiple geometries, such as whole cell and nuclei segmentations. The geometries are assumed to be POINTs for centroids and POLYGONs for segmentations. If polygons are specified in an ordinary data frame, then anything with fewer than 3 vertices will be removed. For anything other than POINTs, attributes of the geometry will be ignored.

`rowGeometries` Geometry associated with genes or features, which correspond to rows of the gene count matrix.

`annotGeometries`
Geometry of entities that do not correspond to columns or rows of the gene count matrix, such as tissue boundary and pathologist annotations of histological regions, and nuclei segmentation in a Visium dataset. Also a named list as in `colGeometries`. The ordinary data frame may specify POINTs, POLYGONs, or LINESTRINGs, or their MULTI versions. Each data frame can only specify one type of geometry. For MULTI versions, there must be a column "group" to identify each MULTI geometry.

`spatialCoordsNames`
A `character` vector of column names if `*Geometries` arguments have ordinary data frames, to identify the columns in the ordinary data frames that specify the spatial coordinates. If `colGeometries` is not specified, then this argument will behave as in `SpatialExperiment`, but `colGeometries` will be given precedence if provided.

`annotGeometryType`
Character vector specifying geometry type of each element of the list if `annotGeometry` is specified. Each element of the vector must be one of POINT, LINESTRING, POLYGON, MULTIPOINT, MULTILINESTRING, and MULTIPOLYGON. Must be either length 1 (same for all elements of the list) or the same length as the list. Ignored if the corresponding element is an `sf` object.

`spatialGraphs` A named list of `listw` objects (see `spdep`) for spatial neighborhood graphs.

`spotDiameter` Spot diameter for technologies with arrays of spots of fixed diameter per slide, such as Visium, ST, DBiT-seq, and slide-seq. The diameter must be in the same unit as the coordinates in the *Geometry arguments. Ignored for geometries that are not POINT or MULTIPOINT.

`unit` # Default unit is `"micron"`. However for Visium one can choose between `"micron"` or `"full_res_image_pixel"`.

`sample_id` A `character` sample identifier, which matches the `sample_id` in `imgData`. The `sample_id` will also be stored in a new column in `colData`, if not already present. Default = `sample01`.

`spatialCoords` A numeric matrix containing columns of spatial coordinates, as in `SpatialExperiment`. The coordinates are centroids of the entities represented by the columns of the gene count matrix. If `colGeometries` is also specified, then it will be given priority and a warning is issued. Otherwise, the `sf` representation of the centroids will be stored in the `colGeometry` called `centroids`.

`scaleFactors` Optional scale factors associated with the image(s). This can be provided as a numeric value, numeric vector, list, or file path to a JSON file for the 10x

Genomics Visium platform. For 10x Genomics Visium, the correct scale factor will automatically be selected depending on the resolution of the image from `imageSources`. Default = `1`.

`imageSources` Optional file path(s) or URL(s) for one or more image sources.

`image_id` Optional character vector (same length as `imageSources`) containing unique image identifiers.

`loadImage` Logical indicating whether to load image into memory. Default = `FALSE`.

`imgData` Optional `DataFrame` containing the image data. Alternatively, this can be built from the arguments `imageSources` and `image_id` (see Details).

`add_molecules` Logical, whether to add transcripts coordinates to an object.

`flip` To flip the image, geometry coordinates, or none. Because the image has the origin at the top left while the geometry has origin at the bottom left, one of them needs to be flipped for them to match. If one of them is already flipped, then use "none". The image will not be flipped if it's GeoTIFF.

`image_scalefactors` # A `character`, choose between "lowres" or "hires". Only for 10X Visium, image scaling factors are from 'scalefactors_json.json' file.

`BPPARAM` Deprecated when coercing from `SpatialExperiment`, but is used when coercing from `Seurat` object.

Value

A `SpatialFeatureExperiment` object

Examples

```
library(SpatialExperiment)
example(read10xVisium)
# There can't be duplicate barcodes
colnames(spe) <- make.unique(colnames(spe), sep = "-")
rownames(spatialCoords(spe)) <- colnames(spe)
sfe <- toSpatialFeatureExperiment(spe)
# For coercing Seurat to SFE see this -> ./vignettes/seurat_sfe_coerce.Rmd
```

SpatialFeatureExperiment-subset

Subsetting SpatialFeatureExperiment objects

Description

The SFE method has special treatment for the spatial graphs. In `listw`, the neighbors are indicated by indices, which will change after subsetting. The `SFE_graph_subset` option determines whether the graphs are subsetted or reconstructed. In the default (`options(SFE_graph_subset = TRUE)`), the graphs are subsetted, in which case singletons may be produced. For `options(SFE_graph_subset = FALSE)`, which is the behavior of versions earlier than Bioc 3.20, the graphs are reconstructed with the parameters recorded in an attribute of the graphs. This option can result into different graphs. For example, suppose we start with a k nearest neighbor graph. After subsetting, cells at the boundary of the region used to subset the SFE object may lose some of their neighbors. In contrast, when the graph is reconstructed, these same edge cells will gain other cells that remain after subsetting as neighbors in the new KNN graph.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment,ANY,ANY,ANY'
x[i, j, ..., drop = FALSE]
```

Arguments

x	A SpatialFeatureExperiment object.
i	Row indices for subsetting.
j	column indices for subsetting.
...	Passed to the SingleCellExperiment method of [.
drop	Only used if graphs are reconstructed (options(SFE_graph_subset = FALSE)). If TRUE then colGraphs are dropped but annotGraphs are kept.

Details

The option SFE_graph_subset was introduced because subsetting is usually faster than reconstructing and in some cases such as distance-based neighbors and Visium spot adjacency give the same results. It was introduced also because of the development of alabster.sfe for a language-agnostic on-disk serialization of SFE objects and some parameters used to construct graphs have special classes whose alabaster methods have not been implemented, such as BPPARAM and BNPARAM, so when reconstructing, the defaults for those arguments will be used.

The edge weights will be recomputed from the binary neighborhood indicator with the same normalization style as the original graph, such as "W" for row normalization. When distance-based edge weights are used instead of the binary indicator, the edge weights will be re-normalized, which is mostly some rescaling. This should give the same results as recomputing the distance based edge weights for styles "raw", "W", and "B" since the distances themselves don't change, but the effects of other more complicated styles of re-normalization on spatial statistics should be further investigated.

By default, upon subsetting, the images are cropped to the bounding box of the remaining cells. However, when the image is large and the bounding box contains most of the original image, cropping is slow. Cropping can be disabled by options(SFE_subset_crop = FALSE). Also, when the remaining part of the image is larger than a threshold, the image will not be cropped; the threshold can be set with the SFE_subset_crop_max option, such as options(SFE_subset_crop_max = "100MB").

Value

A subsetted SpatialFeatureExperiment object.

Examples

```
# Just like subsetting matrices and SingleCellExperiment
library(SFEData)
sfe <- McKellarMuscleData(dataset = "small")
sfe_subset <- sfe[seq_len(10), seq_len(10), drop = TRUE]
# Gives warning as graph reconstruction fails

sfe_subset <- sfe[seq_len(10), seq_len(10)]
```

spatialGraphs *Spatial graph methods*

Description

Spatial neighborhood graphs as spdep's `listw` objects are stored in the `int_metadata` of the SFE object. The `listw` class is used because spdep has many useful methods that rely on the neighborhood graph as `listw`.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
spatialGraphs(x, MARGIN = NULL, sample_id = "all", name = "all")

colGraphs(x, sample_id = "all", name = "all")

rowGraphs(x, sample_id = "all", name = "all")

annotGraphs(x, sample_id = "all", name = "all")

## S4 replacement method for signature 'SpatialFeatureExperiment'
spatialGraphs(x, MARGIN = NULL, sample_id = "all", name = "all") <- value

colGraphs(x, sample_id = "all", name = "all") <- value

rowGraphs(x, sample_id = "all", name = "all") <- value

annotGraphs(x, sample_id = "all", name = "all") <- value

## S4 method for signature 'SpatialFeatureExperiment,numeric'
spatialGraphNames(x, MARGIN, sample_id = 1L)

## S4 replacement method for signature 'SpatialFeatureExperiment,numeric,ANY,character'
spatialGraphNames(x, MARGIN, sample_id = 1L) <- value

colGraphNames(x, sample_id = 1L)

rowGraphNames(x, sample_id = 1L)

annotGraphNames(x, sample_id = 1L)

colGraphNames(x, sample_id = 1L) <- value

rowGraphNames(x, sample_id = 1L) <- value

annotGraphNames(x, sample_id = 1L) <- value

## S4 method for signature 'SpatialFeatureExperiment'
spatialGraph(x, type = 1L, MARGIN, sample_id = 1L)

colGraph(x, type = 1L, sample_id = 1L)
```

```
rowGraph(x, type = 1L, sample_id = 1L)

annotGraph(x, type = 1L, sample_id = 1L)

## S4 replacement method for signature 'SpatialFeatureExperiment'
spatialGraph(x, type = 1L, MARGIN, sample_id = NULL) <- value

colGraph(x, type = 1L, sample_id = 1L) <- value

rowGraph(x, type = 1L, sample_id = 1L) <- value

annotGraph(x, type = 1L, sample_id = 1L) <- value
```

Arguments

x	A SpatialFeatureExperiment object.
MARGIN	As in apply. 1 stands for rows and 2 stands for columns. In addition, 3 stands for spatial neighborhood graphs that correspond to annotGeometries.
sample_id	Name of the sample the graph is associated with. This is useful when multiple pieces of tissues are in the same SFE object (say for a joint dimension reduction and clustering) and the spatial neighborhood is only meaningful within the same piece of tissue. See the sample_id argument in SpatialExperiment.
name	Name of the graphs to add to each sample_id; used in the spatialGraphs replacement method as it must be character while type can be either an integer index or a name.
value	A listw object (*Graph), or a named list of list of listw objects (*Graphs) where the names of the top level list are sample_ids when adding graphs for all samples in the margin of interest, or a list of listw objects when adding graphs for one sample in one margin.
type	An integer specifying the index or string specifying the name of the *Graph to query or replace. If missing, then the first item in the *Graph will be returned or replaced.

Value

Getters for multiple graphs return a named list. Getters for names return a character vector of the names. Getters for single graphs return a listw object. Setters return an SFE object.

Examples

```
library(SFEData)
sfe <- McKellarMuscleData(dataset = "small")
g1 <- findVisiumGraph(sfe)
g2 <- findSpatialNeighbors(sfe)

# Set all graphs of a margin by a named list
spatialGraphs(sfe, MARGIN = 2L, sample_id = "Vis5A") <-
    list(tri2nb = g2, visium = g1)
# Or equivalently
colGraphs(sfe, sample_id = "Vis5A") <- list(tri2nb = g2, visium = g1)

# Get all graphs of a margin, returning a named list
```

```
gs <- spatialGraphs(sfe, MARGIN = 2L)
# Or equivalently
gs <- colGraphs(sfe)

# Set graph of the same name and same margin for multiple samples
# Each sample has a separate graph
sfe2 <- McKellarMuscleData("small2")
sfe_combined <- cbind(sfe, sfe2)
colGraphs(sfe_combined, name = "visium", sample_id = "all") <-
    findVisiumGraph(sfe_combined, sample_id = "all")

# Get graph names
spatialGraphNames(sfe, MARGIN = 2L, sample_id = "Vis5A")
# Or equivalently (sample_id optional as only one sample is present)
colGraphNames(sfe)

# Set graph names
spatialGraphNames(sfe, MARGIN = 2L) <- c("foo", "bar")
colGraphNames(sfe) <- c("tri2nb", "visium")

# MARGIN = 1 means rowGraphs; MARGIN = 3 means annotation graphs (annotGraphs)
# for both getters and setters

# Set single graph by
# Spatial graph for myofibers
g_myofiber <- findSpatialNeighbors(sfe,
    type = "myofiber_simplified",
    MARGIN = 3L
)
spatialGraph(sfe, type = "myofiber", MARGIN = 3L) <- g_myofiber
# Or equivalently
annotGraph(sfe, "myofiber") <- g_myofiber

# Get a specific graph by name
g <- spatialGraph(sfe, "myofiber", MARGIN = 3L)
g2 <- spatialGraph(sfe, "visium", MARGIN = 2L)
# Or equivalently
g <- annotGraph(sfe, "myofiber")
g2 <- colGraph(sfe, "visium")
```

SpatRasterImage *SpatRaster representation of images in SFE objects*

Description

SpatialFeatureExperiment and the Voyager package work with images differently from SpatialExperiment. In SFE and Voyager's, plotting functions for SFE objects, the images can be read with rast and represented as SpatRaster, so the image is not entirely loaded into memory unless necessary. Plotting will not load a large image into memory; rather the image will be downsampled and the downsampled version is plotted. A SpatRasterImage object (as of Bioc 3.19 or SFE version 1.6 and above) is a SpatRaster object but also inheriting from VirtualSpatialImage as required by SpatialExperiment.

Usage

```
SpatRasterImage(img)

## S4 method for signature 'SpatRasterImage'
show(object)
```

Arguments

img	A SpatRaster or PackedSpatRaster object.
object	A SpatRasterImage object.

Value

A SpatRasterImage object.

Examples

```
# Example code
```

---

splitByCol — *Split SFE object with categorical vector or geometry*

Description

The split methods for SFE split an SFE object into multiple SFE objects by geometries (all cells/spots intersecting with each geometry will become a separate SFE object). The splitSamples function splits the SFE object by sample_id so each sample will become a separate SFE object. The splitContiguity function splits the SFE object by contiguity of an annotGeometry, which by default is "tissueBoundary".

Usage

```
## S4 method for signature 'SpatialFeatureExperiment,sf'
splitByCol(x, f, sample_id = "all", colGeometryName = 1L, cover = FALSE)

## S4 method for signature 'SpatialFeatureExperiment,sfc'
splitByCol(x, f, sample_id = 1L, colGeometryName = 1L, cover = FALSE)

## S4 method for signature 'SpatialFeatureExperiment,list'
splitByCol(x, f, sample_id = "all", colGeometryName = 1L, cover = FALSE)

splitSamples(x)

splitContiguity(
  x,
  colGeometryName = 1L,
  annotGeometryName = "tissueBoundary",
  min_area = 0,
  cover = FALSE
)
```

Arguments

- `x` — An SFE object
- `f` — It can be a `sf` data frame or `sfc` to split by geometry. Each row of the `sf` data frame or each element in the `sfc` will correspond to a new SFE object. The `sf` data frame must have a column `sample_id` when splitting multiple samples. Can also be a list of `sfc` whose names correspond to `sample_ids` to split.
- `sample_id` — Which samples to split.
- `colGeometryName` — Which `colGeometry` to use to determine which cells or spots should belong to which new SFE object when splitting by `sf` or `sfc`. Default to the first one.
- `cover` — Logical, whether the geometries in `x` must be entirely covered by `y` if `op = st_intersection` or whether `x` must be entirely outside `y` if `op = st_difference`. Only relevant when `keep_whole != "none"`.
- `annotGeometryName` — Name of `annotGeometry` to use to split by contiguity.
- `min_area` — Minimum area in the same unit as the geometry coordinates (squared) for each piece to be considered a separate piece when splitting by contiguity. Only pieces that are large enough are considered.

Value

A list of SFE objects.

Examples

```
# example code
```

---

st_any_pred — *Simple geometry predicates*

Description

Unlike functions in `sf` like `st_intersects`, this function simply returns a logical vector indicating whether each geometry in `x` intersects (or returns `TRUE` from other predicates) anything in `y`, preferably when `y` only contains a small number of geometries or is one single MULTI geometry. This is useful when cropping or subsetting an SFE object with a geometry, such as tissue boundary or histological region polygons or a bounding box.

Usage

```
st_any_pred(x, y, pred, yx = FALSE, sparse = FALSE, ...)

st_any_intersects(x, y, yx = FALSE, sparse = FALSE)

st_n_pred(x, y, pred, ...)

st_n_intersects(x, y)
```

Arguments

x	An object of class sf, sfc, or sfg.
y	Another object of class sf, sfc, or sfg.
pred	A geometric binary predicate function, such as st_intersects. It should return an object of class sgbp, for sparse predicates.
yx	Whether to do pred(y, x) instead of pred(x, y). For symmetric predicates, the results should be the same. When x has a large number of geometries and y has few, pred(y, x) is much faster than pred(x, y) for st_intersects, st_disjoint, and st_is_within_distance.
sparse	If TRUE, returns numeric indices rather than logical vector. Defaults to FALSE for backward compatibility, though the default in st_intersects is TRUE.
...	Arguments passed to pred.

Value

For st_any_*, a logical vector indicating whether each geometry in x intersects (or other predicates such as is covered by) anything in y or a numeric vector of indices of TRUE when sparse = TRUE. Simplified from the sgbp results which indicate which item in y each item in x intersects, which might not always be relevant. For st_n_*, an integer vector indicating the number of geometries in y returns TRUE for each geometry in x.

Examples

```
library(sf)
pts <- st_sfc(
    st_point(c(.5, .5)), st_point(c(1.5, 1.5)),
    st_point(c(2.5, 2.5))
)
pol <- st_polygon(list(rbind(c(0, 0), c(2, 0), c(2, 2), c(0, 2), c(0, 0))))
st_any_pred(pts, pol, pred = st_disjoint)
st_any_intersects(pts, pol)
st_n_pred(pts, pol, pred = st_disjoint)
st_n_intersects(pts, pol)
```

---

toExtImage *Convert images to ExtImage*

Description

The ExtImage class is a thin wrapper around the Image class in ExtImage so it inherits from VirtualSpatialImage as required by SpatialExperiment and has extent as used in Voyager's plotting functions. This function converts SpatRasterImage (thin wrapper around the class in terra) and BioFormatsImage into ExtImage for image operations as implemented in the ExtImage package.

Usage

```
## S4 method for signature 'BioFormatsImage'
toExtImage(x, resolution = 4L, channel = NULL)

## S4 method for signature 'SpatRasterImage'
toExtImage(x, maxcell = 1e+07, channel = NULL)
```

Arguments

x	Either a BioFormatsImage or SpatRasterImage object.
resolution	Integer, which resolution in the BioFormatsImage to read and convert. Defaults to 4, which is a lower resolution. Ignored if only 1 resolution is present.
channel	Integer vector to indicate channel(s) to read. If NULL, then all channels will be read.
maxcell	Maximum number of pixels when SpatRasterImage is read into memory.

Value

A ExtImage object. The image is loaded into memory.

See Also

toSpatRasterImage

---

toSpatRasterImage *Convert images to SpatRasterImage*

Description

The resolution specified from the OME-TIFF file will be read into memory and written to disk as a GeoTIFF file that has the extent. The output file will have the same file name as the input file except without the ome in the extension.

Usage

```
## S4 method for signature 'ExtImage'
toSpatRasterImage(
  x,
  save_geotiff = TRUE,
  file_out = "img.tiff",
  overwrite = FALSE
)

## S4 method for signature 'BioFormatsImage'
toSpatRasterImage(
  x,
  save_geotiff = TRUE,
  resolution = 4L,
  channel = NULL,
  overwrite = FALSE
)
```

Arguments

x	Either a BioFormatsImage or EBIImage object.
save_geotiff	Logical, whether to save the image to GeoTIFF file.
file_out	File to save the non-OME TIFF file for SpatRaster.

overwrite	Logical, whether to overwrite existing file of the same name.
resolution	Integer, which resolution in the BioFormatsImage to read and convert. Defaults to 4, which is a lower resolution. Ignored if only 1 resolution is present.
channel	Integer vector to indicate channel(s) to read. If NULL, then all channels will be read.

Value

A SpatRasterImage object

See Also

toExtImage

---

translateImg *Translate/shift image in space*

Description

This function shifts the spatial extent of the image in the x-y plane.

Usage

```
## S4 method for signature 'SpatRasterImage'
translateImg(x, v, ...)

## S4 method for signature 'BioFormatsImage'
translateImg(x, v, ...)

## S4 method for signature 'ExtImage'
translateImg(x, v, ...)
```

Arguments

x	An object of class *Image as implemented in this package.
v	Numeric vector of length 2 to shift the image in the x-y plane.
...	Ignored. It's there so different methods can all be passed to the same lapply in the method for SFE objects. Some methods have extra arguments.

Value

A *Image object of the same class that has been shifted in space.

See Also

Other image methods: SFE-image, affineImg(), cropImg(), dim,BioFormatsImage-method, dim,ExtImage-method, ext(), imgRaster(), imgSource(), mirrorImg(), rotateImg(), scaleImg(), transposeImg()

`transposeImg` *Transpose images*

Description

Swap rows and columns of images. In effect, this will flip the image around the diagonal running from top left to bottom right.

Usage

```
## S4 method for signature 'SpatRasterImage'
transposeImg(x, filename = "", maxcell = NULL, ...)

## S4 method for signature 'BioFormatsImage'
transposeImg(x, ...)

## S4 method for signature 'ExtImage'
transposeImg(x, ...)
```

Arguments

`x`	An object of class `*Image` as implemented in this package.
`filename`	Output file name for transformed SpatRaster.
`maxcell`	Max number of pixels to load `SpatRasterImage` into memory. The default 1e7 is chosen because this is the approximate number of pixels in the medium resolution image at `resolution = 4L` in Xenium OME-TIFF to make different methods of this function consistent.
`...`	Ignored. It's there so different methods can all be passed to the same `lapply` in the method for SFE objects. Some methods have extra arguments.

Value

For `SpatRasterImage` and `ExtImage`, object of the same class. For `BioFormatsImage`, the image of the specified resolution is read into memory and then the `ExtImage` method is called, returning `ExtImage`. For the extent: xmin and xmax are switched with ymin and ymax.

See Also

Other image methods: `SFE-image`, `affineImg()`, `cropImg()`, `dim,BioFormatsImage-method`, `dim,ExtImage-method`, `ext()`, `imgRaster()`, `imgSource()`, `mirrorImg()`, `rotateImg()`, `scaleImg()`, `translateImg()`

unit,SpatialFeatureExperiment-method

Get unit of a SpatialFeatureExperiment

Description

Length units can be microns or pixels in full resolution image in SFE objects.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
unit(x)
```

Arguments

x	A SpatialFeatureExperiment object.

Value

A string for the name of the unit. At present it's merely a string and udunits is not used.

Examples

```
library(SFEData)
sfe <- McKellarMuscleData(dataset = "small")
SpatialFeatureExperiment::unit(sfe)
```

updateObject

Update a SpatialFeatureExperiment object

Description

Update a SpatialFeatureExperiment to the latest version of object structure. This is usually called by internal functions.

Usage

```
## S4 method for signature 'SpatialFeatureExperiment'
updateObject(object, ..., verbose = FALSE)

SFEVersion(object)
```

Arguments

object	An old SpatialFeatureExperiment object.
...	Additional arguments that are ignored.
verbose	Logical scalar indicating whether a message should be emitted as the object is updated.

Details

Version 1.1.4 adds package version to the SFE object. We are considering an overhaul of the `spatialGraphs` slot in a future version using the `sfdep` package to decouple the adjacency graph from the edge weights.

Value

An updated version of `object`.

See Also

`objectVersion`, which is used to determine if the object is up-to-date.

Examples

```
library(SFEData)
sfe <- McKellarMuscleData("small")
# First version of SFE object doesn't log SFE package version, so should be NULL
SFEVersion(sfe)
sfe <- updateObject(sfe)
# See current version
SFEVersion(sfe)
```

`visium_row_col` *Row and columns of Visium barcodes on the slide*

Description

From Space Ranger 1.3.1.

Usage

```
visium_row_col
```

Format

A data frame with 4992 rows with columns barcode, col, and row.

Source

Space Ranger 1.3.1

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