

# Package: concordexR (via r-universe)

July 24, 2024

**Title** Identify Spatial Homogeneous Regions with concordex

**Version** 1.5.5

**Description** Spatial homogeneous regions (SHRs) in tissues are domains that are homogenous with respect to cell type composition. We present a method for identifying SHRs using spatial transcriptomics data, and demonstrate that it is efficient and effective at finding SHRs for a wide variety of tissue types. concordex relies on analysis of k-nearest-neighbor (kNN) graphs. The tool is also useful for analysis of non-spatial transcriptomics data, and can elucidate the extent of concordance between partitions of cells derived from clustering algorithms, and transcriptomic similarity as represented in kNN graphs.

**License** Artistic-2.0

**URL** <https://github.com/pachterlab/concordexR>,  
<https://pachterlab.github.io/concordexR/>

**BugReports** <https://github.com/pachterlab/concordexR/issues>

**Depends** R (>= 4.3.0)

**Imports** BiocGenerics, BiocNeighbors, BiocParallel, bluster, cli,  
DelayedArray, Matrix, methods, purrr, rlang,  
SingleCellExperiment, sparseMatrixStats, SpatialExperiment,  
SummarizedExperiment

**Suggests** BiocManager, BiocStyle, ggplot2, glue, knitr, mbkmeans,  
patchwork, rmarkdown, scater, SFData,  
SpatialFeatureExperiment, TENxPBMCDData, testthat (>= 3.0.0)

**VignetteBuilder** knitr

**biocViews** SingleCell, Clustering, Spatial, Transcriptomics

**Config/testthat/edition** 3

**Encoding** UTF-8

**Roxygen** list(markdown = TRUE)

**RoxygenNote** 7.3.2

**Repository** <https://bioc.r-universe.dev>

**RemoteUrl** <https://github.com/bioc/concordexR>

**RemoteRef** HEAD

**RemoteSha** d14e3aa11f19adbae394046e016d3ebfbd23e4b

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## Description

Compute the neighborhood consolidation matrix and identify spatial homogeneous regions.

## Usage

```
calculateConcordex(x, ...)

## S4 method for signature 'ANY'
calculateConcordex(
  x,
  labels,
  ...,
  n_neighbors = 30,
  compute_similarity = FALSE,
  BLUSPARAM,
  BNINDEX,
  BNPARAM = KmknParam(),
  BPPARAM = SerialParam()
)

## S4 method for signature 'SummarizedExperiment'
calculateConcordex(x, labels, ..., assay.type = "logcounts")

## S4 method for signature 'SingleCellExperiment'
calculateConcordex(x, labels, ..., use.dimred = NULL)

## S4 method for signature 'SpatialExperiment'
calculateConcordex(x, labels, ..., use.spatial = TRUE)
```

**Arguments**

x	A <a href="#">SpatialExperiment</a> , <a href="#">SpatialFeatureExperiment</a> , <a href="#">SingleCellExperiment</a> , or <a href="#">SummarizedExperiment</a> object containing a count matrix. Otherwise, a numeric matrix-like object containing counts for observations (e.g. cells/spots) on the rows and features on the columns.
...	Other parameters passed to default method
labels	Observation labels used to compute the neighborhood consolidation matrix. Continuous or discrete labels are allowed, and typically, integer labels are assumed to be discrete. Labels can be specified as follows: <ul style="list-style-type: none"> <li>• A vector or matrix-like object with one entry per observation. If a matrix, the observations should be on the rows and the label identifiers on the columns.</li> <li>• If x inherits from <a href="#">SummarizedExperiment-class</a>, a string or character vector specifying the names of columns in <code>colData(x)</code> or the name of a dimensionality reduction result (see <a href="#">reducedDimNames</a>)</li> </ul>
n_neighbors	Number of neighbors to expect for each observation. Defaults to 30.
compute_similarity	Logical. Whether to return the label similarity matrix. Only useful if discrete labels are provided.
BLUSPARAM	A <a href="#">BlusterParam-class</a> object specifying the clustering algorithm to use to identify spatial homogeneous regions. If this parameter is not specified, then regions are not returned. By default, this parameter is missing.
BNINDEX	A <a href="#">BiocNeighborIndex</a> object containing the precomputed index information, see <a href="#">findKNN</a> .
BNPARAM	A <a href="#">BiocNeighborParam</a> object specifying the algorithm to use. This can be missing if BNINDEX is supplied, see <a href="#">findKNN</a> .
BPPARAM	A <a href="#">BiocParallelParam</a> object specifying whether and how computing the metric for numerous observations shall be parallelized (see <a href="#">bpparam</a> ).
assay.type	String or integer scalar indicating the assay of x containing the counts.
use.dimred	Integer or string specifying the reduced dimensions to use for construction of the k-nearest neighbor graph. Note that if this is not NULL, reduced dimensions can not be used as labels to compute the neighborhood consolidation matrix.
use.spatial	Logical, should the spatial coordinates be used to compute the k-nearest neighbor graph?

**Value**

A sparse matrix

**Examples**

```
example(read10xVisium, "SpatialExperiment")
library(bluster)

## Setting BLUSPARAM clusters the consolidation
```

```
## matrix into SHRs
cdx <- calculateConcordex(
  spe, "in_tissue",
  n_neighbors=10,
  BLUSPARAM=KmeansParam(3)
)

## SHRs are an attribute of the result
shr <- attr(cdx, "shr")

## The label similarity matrix can be computed
## with `compute_similarity=TRUE`
cdx <- calculateConcordex(
  spe, "in_tissue",
  n_neighbors=10,
  compute_similarity=TRUE,
  BLUSPARAM=KmeansParam(3)
)
```

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