

# Package: CySA (via r-universe)

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**Title** Interactive Cluster Selector for Cytometry Data

**Version** 0.99.7

**Author** Bernd Jagla [aut, cre]  
(<https://orcid.org/0000-0002-7696-0484>)

**Maintainer** Bernd Jagla <bernd.jagla@pasteur.fr>

**Description** Provides an interactive Shiny application for selecting and visualizing clusters from flow cytometry data stored in SingleCellExperiment objects.

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**URL** <https://github.com/baj12/CySA>

**BugReports** <https://github.com/baj12/CySA/issues>

**Depends** R (>= 4.5.0)

**Imports** shiny, shinydashboard, shinydashboardPlus, shinyjs, shinyjqui, htmltools, SingleCellExperiment, S4Vectors, SummarizedExperiment, Matrix, RColorBrewer, CATALYST, dplyr, tidy, tibble, data.table, stringr, purrr, reshape2, rlang, raster, ggplot2, plotly, viridis, dendextend, ComplexHeatmap, cowplot, Rtsne, umap, DT, collapsibleTree, withr

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## Contents

clusterSelector . . . . .	2
CySA_default_cluster_cols . . . . .	4
CySA_example_sce . . . . .	5
plotScatterBJ . . . . .	5
plotSOMScatter . . . . .	6
prepClusterSelectorData . . . . .	8
<b>Index</b>	<b>9</b>

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clusterSelector	<i>Cluster Selector Shiny Application</i>
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## Description

Creates an interactive Shiny dashboard for selecting and visualizing clusters from a SingleCellExperiment object.

## Usage

```
clusterSelector(
  sce,
  sce_subsampled,
  outputList = list(),
  colTree = NULL,
  dList,
  dend,
  dendTable,
  clusterPatientTable,
  somCodesName = "SOM_codes",
  nPlots = 6,
  somRasterData,
  somRasterObj,
  env = environment()
)
```

**Arguments**

sce	A <a href="#">SingleCellExperiment</a> containing the full dataset.
sce_subsampled	A subsampled <a href="#">SingleCellExperiment</a> for performance-sensitive plots.
outputList	A named list of cluster groupings.
colTree	Optional collapsible tree object.
dList	List of marker pairs for 2D plots.
dend	Dendrogram object.
dendTable	Data frame for dendrogram navigation.
clusterPatientTable	Table of sample by cluster counts.
somCodesName	Name of the SOM codes metadata slot.
nPlots	Number of 2D SOM plots to display.
somRasterData	Data frame for SOM raster visualization.
somRasterObj	Raster object for SOM visualization.
env	Environment used to store mutable state (legacy argument).

**Value**

A [shinyApp](#) object. Launch the app with `shiny::runApp(app)`.

**Examples**

```
sce <- CySA_example_sce()
prepped <- prepClusterSelectorData(sce, total_cells_to_sample = 100)
som_codes <- S4Vectors::metadata(sce)$SOM_codes
dend <- stats::as.dendrogram(stats::hclust(stats::dist(som_codes)))
dendTable <- data.frame(
  id = seq_len(nrow(som_codes)),
  label = rownames(som_codes),
  stringsAsFactors = FALSE
)
clusterPatientTable <- table(
  sample_id = sce$sample_id,
  cluster_id = sce$cluster_id
)
markers <- S4Vectors::metadata(sce)$map$colsUsed
somRasterData <- data.frame(
  x = rep(seq_len(5), length.out = nrow(som_codes)),
  y = rep(seq_len(2), each = nrow(som_codes) / 2),
  id = seq_len(nrow(som_codes))
)
for (m in markers) {
  somRasterData[[m]] <- seq_len(nrow(som_codes)) / nrow(som_codes)
}
arr <- array(
  data = seq_len(10 * 10 * length(markers)),
  dim = c(10, 10, length(markers))
)
```

```
)  
somRasterObj <- raster::brick(arr)  
names(somRasterObj) <- markers  
  
app <- clusterSelector(  
  sce = prepped$sce,  
  sce_subsampled = prepped$sce_subsampled,  
  dList = prepped$dList,  
  dend = dend,  
  dendTable = dendTable,  
  clusterPatientTable = clusterPatientTable,  
  somRasterData = somRasterData,  
  somRasterObj = somRasterObj  
)  
if (interactive()) {  
  shiny::runApp(app)  
}
```

---

CySA\_default\_cluster\_cols

*Default cluster color palette*

---

## Description

Returns the default 20-color palette used by CySA for cluster visualizations.

## Usage

```
CySA_default_cluster_cols()
```

## Value

A character vector of hex colors.

## Examples

```
CySA_default_cluster_cols()
```

---

CySA\_example\_sce      *Minimal example SingleCellExperiment for CySA*

---

### Description

Creates a small, deterministic `SingleCellExperiment` object that contains the metadata and column data expected by CySA functions.

### Usage

```
CySA_example_sce(n_cells = 1000, n_nodes = 50, n_markers = 12)
```

### Arguments

<code>n_cells</code>	Number of cells (columns).
<code>n_nodes</code>	Number of SOM nodes.
<code>n_markers</code>	Number of markers (rows).

### Value

A `SingleCellExperiment` with `SOM_codes`, `SOM_stats`, `sample_id`, and `cluster_id`.

### Examples

```
sce <- CySA_example_sce()
head(S4Vectors::metadata(sce)$SOM_codes)
```

---

plotScatterBJ      *Plot Scatter for Bioconductor Experiment*

---

### Description

This function generates a scatter plot for a Bioconductor Experiment object.

### Usage

```
plotScatterBJ(
  x,
  chs,
  gate = NULL,
  color_by = NULL,
  facet_by = NULL,
  bins = 100,
  assay = "exprs",
```

```

  label = c("target", "channel", "both"),
  zeros = FALSE,
  k_pal = NULL,
  rowNames = NULL
)

```

### Arguments

x	A Bioconductor Experiment object.
chs	A character vector specifying the variables to be plotted on the x and y axes.
gate	Optional gate object.
color_by	A character specifying the variable used for color coding points (optional).
facet_by	A character specifying the variable used for faceting the plot (optional).
bins	Number of bins for the 2D histogram (default is 100).
assay	The assay to use for plotting (default is "exprs").
label	Label for points on the plot ("target", "channel", or "both").
zeros	Logical, whether to exclude rows with all zero values (default is FALSE).
k_pal	Color palette for plotting.
rowNames	Optional character vector to specify row names for the plot.

### Value

A ggplot2 scatter plot.

### Examples

```

sce <- CySA_example_sce()
plotScatterBJ(sce, chs = c("marker1", "marker2"))

```

---

plotSOMScatter

*Scatter plot*

---

### Description

Bivariate scatter plots including visualization of (group-specific) gates, their boundaries and percentage of selected cells.

**Usage**

```
plotSOMScatter(  
  x,  
  chs,  
  metaSlot = "SOM_codes",  
  pointSize = "n",  
  color_by = "n",  
  bins = 100,  
  assay = "exprs",  
  statsSlot = "SOM_stats",  
  label = c("target", "channel", "both"),  
  zeros = FALSE,  
  k_pal = NULL,  
  xRN = NULL,  
  xCN = NULL  
)
```

**Arguments**

x	a <a href="#">SingleCellExperiment</a> .
chs	character vector specifying which channels to plot.
metaSlot	name of the metadata slot containing SOM codes.
pointSize	column in SOM stats used to size points.
color_by	column to color points by.
bins	number of bins when coloring by density.
assay	name of the assay to use.
statsSlot	name of the metadata slot containing SOM stats.
label	how axis labels should be constructed.
zeros	logical specifying whether to include 0 values.
k_pal	optional cluster color palette.
xRN	optional row names.
xCN	optional channel names.

**Value**

a ggplot object.

**Examples**

```
sce <- CySA_example_sce()  
plotSOMScatter(sce, chs = c("marker1", "marker2"))
```

---

`prepClusterSelectorData`*Prepare Data for the Cluster Selector Shiny App*

---

## Description

Subsamples a `SingleCellExperiment` object and builds the inputs required by `clusterSelector`.

## Usage

```
prepClusterSelectorData(  
  sce,  
  somFile = NULL,  
  dList = NULL,  
  total_cells_to_sample = 1e+05,  
  somCodesName = "SOM_codes",  
  assay = "exprs",  
  seed = 123  
)
```

## Arguments

<code>sce</code>	A <code>SingleCellExperiment</code> containing the full dataset.
<code>somFile</code>	Path to the SOM object. Used to cache/restore the subsampled version.
<code>dList</code>	Optional list of marker pairs for 2D plots. If NULL, defaults to the first 12 row names of <code>sce</code> .
<code>total_cells_to_sample</code>	Number of cells to subsample in total.
<code>somCodesName</code>	Name of the SOM codes metadata slot.
<code>assay</code>	Name of the assay to use.
<code>seed</code>	Random seed for reproducibility.

## Value

A list with `sce`, `sce_subsampled`, and `dList`.

## Examples

```
sce <- CySA_example_sce(n_cells = 200, n_nodes = 10)  
prepped <- prepClusterSelectorData(sce, total_cells_to_sample = 100)  
names(prepped)
```

# Index

clusterSelector, [2](#), [8](#)  
CySA\_default\_cluster\_cols, [4](#)  
CySA\_example\_sce, [5](#)  
  
plotScatterBJ, [5](#)  
plotSOMScatter, [6](#)  
prepClusterSelectorData, [8](#)  
  
shinyApp, [3](#)  
SingleCellExperiment, [3](#), [5](#), [7](#), [8](#)