

# Package ‘traviz’

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**Type** Package

**Title** Trajectory functions for visualization and interpretation.

**Version** 1.8.0

**Description** traviz provides a suite of functions to plot trajectory related objects from Bioconductor packages. It allows plotting trajectories in reduced dimension, as well as average gene expression smoothers as a function of pseudotime. Besides from general utility functions, traviz also allows plotting trajectories estimated by Slingshot, as well as smoothers estimated by tradeSeq. Furthermore, it allows for visualization of Slingshot trajectories using ggplot2.

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**Encoding** UTF-8

**LazyData** false

**Imports** ggplot2, viridis, mgcv, SingleCellExperiment, slingshot, prncurve, Biobase, methods, RColorBrewer, SummarizedExperiment, grDevices, graphics, rgl

**RoxygenNote** 7.1.1

**Collate** 'AllGenerics.R' 'data.R' 'utilsTradeSeq.R' 'plotExpression.R' 'plotGeneCount.R' 'plotSmoothers.R' 'plot\_SlingshotDataSet.R'

**Suggests** scater, dplyr, testthat (>= 3.0.0), covr, S4Vectors, rmarkdown, knitr

**Config/testthat/edition** 3

**Depends** R (>= 4.0)

**biocViews** GeneExpression, RNASeq, Sequencing, Software, SingleCell, Transcriptomics, Visualization

**VignetteBuilder** knitr

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counts	<i>A count matrix, used for testing.</i>
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Description

This object contains the gene expression counts from the data described in Paul et al. (2015).

Usage

```
data("counts", package = "traviz")
```

Format

An object of class standardGeneric of length 1.

Details

#' @references Franziska Paul, Yaara Arkin, Amir Giladi, DiegoAdhemar Jaitin, Ephraim Kenigsberg, Hadas KerenShaul, Deborah Winter, David Lara-Astiaso, Meital Gury, Assaf Weiner, Eyal David, Nadav Cohen, FeliciaKathrineBratt Lauridsen, Simon Haas, Andreas Schlitzer, Alexander Mildner, Florent Ginhoux, Steen Jung, Andreas Trumpp, BoTorben Porse, Amos Tanay, and Ido Amit. Transcriptional Heterogeneity and Lineage Commitment in Myeloid Progenitors. Cell, 163(7):1663–1677, 12 2015. ISSN 0092- 8674. doi: 10.1016/J.CELL.2015.11.013. URL <https://www.sciencedirect.com/science/article/pii/S0092867415014932?via>

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crv

*A SlingshotDataSet object, used for testing.*


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### Description

This dataset contains the Slingshot trajectory from the data described in Paul et al. (2015).

### Usage

```
data("crv", package = "traviz")
```

### Format

An object of class `SlingshotDataSet` of length 1.

### References

Franziska Paul, Yaara Arkin, Amir Giladi, DiegoAdhemar Jaitin, Ephraim Kenigsberg, Hadas KerenShaul, Deborah Winter, David Lara-Astiaso, Meital Gury, Assaf Weiner, Eyal David, Nadav Cohen, FeliciaKathrineBratt Lauridsen, Simon Haas, Andreas Schlitzer, Alexander Mildner, Florent Ginhoux, Steen Jung, Andreas Trumpp, BoTorben Porse, Amos Tanay, and Ido Amit. Transcriptional Heterogeneity and Lineage Commitment in Myeloid Progenitors. *Cell*, 163(7):1663–1677, 12 2015. ISSN 0092- 8674. doi: 10.1016/J.CELL.2015.11.013. URL <https://www.sciencedirect.com/science/article/ii/S0092867415014932?via>

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plot-SlingshotDataSet *Plot Slingshot output*


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### Description

Tools for visualizing lineages inferred by slingshot.

### Usage

```
## S4 method for signature 'SlingshotDataSet,ANY'
plot(
  x,
  type = NULL,
  linInd = NULL,
  show.constraints = FALSE,
  add = FALSE,
  dims = seq_len(2),
  asp = 1,
  cex = 2,
  lwd = 2,
  col = 1,
```

```

    ...
  )

## S4 method for signature 'SlingshotDataSet'
lines(x, type = NULL, dims = seq_len(2), ...)

```

### Arguments

<code>x</code>	a <code>SlingshotDataSet</code> with results to be plotted.
<code>type</code>	character, the type of output to be plotted, can be one of "lineages", "curves", or "both" (by partial matching), see Details for more.
<code>linInd</code>	integer, an index indicating which lineages should be plotted (default is to plot all lineages). If <code>col</code> is a vector, it will be subsetting by <code>linInd</code> .
<code>show.constraints</code>	logical, whether or not the user-specified initial and terminal clusters should be specially denoted by green and red dots, respectively.
<code>add</code>	logical, indicates whether the output should be added to an existing plot.
<code>dims</code>	numeric, which dimensions to plot (default is 1:2).
<code>asp</code>	numeric, the y/x aspect ratio, see <a href="#">plot.window</a> .
<code>cex</code>	numeric, amount by which points should be magnified, see <a href="#">par</a> .
<code>lwd</code>	numeric, the line width, see <a href="#">par</a> .
<code>col</code>	character or numeric, color(s) for lines, see <a href="#">par</a> .
<code>...</code>	additional parameters to be passed to <a href="#">lines</a> .

### Details

If `type == 'lineages'`, straight line connectors between cluster centers will be plotted. If `type == 'curves'`, simultaneous principal curves will be plotted.

When `type` is not specified, the function will first check the curves slot and plot the curves, if present. Otherwise, lineages will be plotted, if present.

### Value

returns `NULL`.

### Examples

```

library(slingshot)
data("slingshotExample", package="slingshot")
rd <- slingshotExample$rd
cl <- slingshotExample$cl
pto <- slingshot(rd, cl, start.clus = "1")
plot(SlingshotDataSet(pto), type = 'b')

# add to existing plot
sds <- as.SlingshotDataSet(pto)
plot(rd, col = 'grey50', asp = 1)
lines(sds, lwd = 3)

```

---

plot3d-SlingshotDataSet

*Plot Slingshot output in 3D*


---

## Description

Tools for visualizing lineages inferred by slingshot.

## Usage

```
## S3 method for class 'SlingshotDataSet'
plot3d(
  x,
  type = NULL,
  linInd = NULL,
  add = FALSE,
  dims = seq_len(3),
  aspect = "iso",
  size = 10,
  col = 1,
  ...
)
```

## Arguments

<code>x</code>	a <code>SlingshotDataSet</code> with results to be plotted.
<code>type</code>	character, the type of output to be plotted, can be one of "lineages", curves, or both (by partial matching), see Details for more.
<code>linInd</code>	integer, an index indicating which lineages should be plotted (default is to plot all lineages). If <code>col</code> is a vector, it will be subsetted by <code>linInd</code> .
<code>add</code>	logical, indicates whether the output should be added to an existing plot.
<code>dims</code>	numeric, which dimensions to plot (default is 1:3).
<code>aspect</code>	either a logical indicating whether to adjust the aspect ratio or a new ratio, see <a href="#">plot3d</a> .
<code>size</code>	numeric, size of points for MST (default is 10), see <a href="#">plot3d</a> .
<code>col</code>	character or numeric, color(s) for lines, see <a href="#">par</a> .
<code>...</code>	additional parameters to be passed to <code>lines3d</code> .

## Details

If `type == 'lineages'`, straight line connectors between cluster centers will be plotted. If `type == 'curves'`, simultaneous principal curves will be plotted.

When `type` is not specified, the function will first check the curves slot and plot the curves, if present. Otherwise, lineages will be plotted, if present.

**Value**

returns NULL.

**Examples**

```
library(rgl)
library(slingshot)
data("crv", package="traviz")
rd <- slingReducedDim(crv)
rd <- cbind(rd, rnorm(nrow(rd), sd=.1))
cl <- apply(slingClusterLabels(crv), 1, which.max)
sds <- slingshot::slingshot(rd, clusterLabels=cl, start.clus=1)
slingshot::plot3d.SlingshotDataSet(as.SlingshotDataSet(sds), type = 'b')

# add to existing plot
plot3d(rd, col = cl, aspect = 'iso')
slingshot::plot3d.SlingshotDataSet(as.SlingshotDataSet(sds), lwd = 3, add = TRUE)
```

---

plotExpression	<i>Plot gene expression along pseudotime.</i>
----------------	---

---

**Description**

Plot gene expression along pseudotime.

Plots a fast loess smoother of gene expression for each lineage.

**Usage**

```
plotExpression(counts, sds, gene, ...)

## S4 method for signature 'matrix,SlingshotDataSet,character'
plotExpression(
  counts,
  sds,
  gene,
  type = "loess",
  span = 0.75,
  alpha = 1,
  lwd = 1,
  size = 2/3
)

## S4 method for signature 'matrix,PseudotimeOrdering,character'
plotExpression(
  counts,
  sds,
```

```

    gene,
    type = "loess",
    span = 0.75,
    alpha = 1,
    lwd = 1,
    size = 2/3
  )

```

### Arguments

counts	The matrix of gene expression counts.
sds	A SlingshotDataSet or PseudotimeOrdering object, typically obtained after trajectory inference using Slingshot.
gene	Gene name of gene to plot.
...	parameters including:
type	The type of smoother. Defaults to "loess".
span	If type is "loess", the span of the smoother. See loess documentation.
alpha	Numeric between 0 and 1, determines the transparency of data points, see scale_color_viridis_d.
lwd	Line width of the smoother. Passed to <a href="#">geom_line</a> .
size	Character expansion of the data points. Passed to <a href="#">geom_point</a> .

### Value

A ggplot object.

### Examples

```

library(ggplot2)
data(crv, package="traviz")
data(counts, package="traviz")
plotExpression(counts = counts, sds=crv, gene=rownames(counts)[1])

```

---

plotGeneCount	<i>Plot gene expression in reduced dimension.</i>
---------------	---

---

### Description

Plot the gene in reduced dimensional space.

**Usage**

```
plotGeneCount(curve, ...)

## S4 method for signature 'SlingshotDataSet'
plotGeneCount(
  curve,
  counts = NULL,
  gene = NULL,
  clusters = NULL,
  models = NULL,
  title = NULL
)

## S4 method for signature 'SingleCellExperiment'
plotGeneCount(
  curve,
  counts = NULL,
  gene = NULL,
  clusters = NULL,
  models = NULL,
  title = NULL
)
```

**Arguments**

curve	One of two <ul style="list-style-type: none"> <li>• A <a href="#">SlingshotDataSet</a> object. The output from trajectory inference using Slingshot.</li> <li>• A <a href="#">SingleCellExperiment</a> object. The output from trajectory inference using Slingshot.</li> </ul>
...	parameters including:
counts	The count matrix, genes in rows and cells in columns. Only needed if the input is of the type <a href="#">SlingshotDataSet</a> and the gene argument is not NULL.
gene	The name of gene for which you want to plot the count or the row number of that gene in the count matrix. Alternatively, one can specify the <code>clusters</code> argument.
clusters	The assignment of each cell to a cluster. Used to color the plot. Either <code>clusters</code> or <code>gene</code> and <code>counts</code> must be supplied.
models	The fitted GAMs, typically the output from <code>fitGAM</code> . Used to display the knots.
title	Title for the plot.

**Details**

If both `gene` and `clusters` arguments are supplied, the plot will be colored according to gene count level. If none are provided, the function will fail.



**Value**

A `ggplot` object

**Examples**

```
set.seed(97)
library(slingshot)
data(crv, package="traviz")
data(counts, package="traviz")
plotGeneCount(crv, counts, gene = "Mpo")
```

---

plotSmootherers	<i>Plot the log-transformed counts and the fitted values for a particular gene along all lineages</i>
-----------------	---

---

**Description**

Plot the smoothers estimated by tradeSeq.

**Usage**

```
plotSmootherers(models, ...)

## S4 method for signature 'gam'
plotSmootherers(
  models,
  nPoints = 100,
  lwd = 2,
  size = 2/3,
  xlab = "Pseudotime",
  ylab = "Log(expression + 1)",
  border = TRUE,
  alpha = 1,
  sample = 1
)

## S4 method for signature 'SingleCellExperiment'
plotSmootherers(
  models,
  counts,
  gene,
  nPoints = 100,
  lwd = 2,
  size = 2/3,
  xlab = "Pseudotime",
  ylab = "Log(expression + 1)",
  border = TRUE,
```

```

    alpha = 1,
    sample = 1,
    pointCol = NULL,
    curvesCols = NULL,
    plotLineages = TRUE
  )

```

## Arguments

<code>models</code>	Either the <code>SingleCellExperiment</code> object obtained after running <code>fitGAM</code> , or the specific GAM model for the corresponding gene, if working with the list output of <code>tradeSeq</code> .
<code>...</code>	parameters including:
<code>nPoints</code>	The number of points used to extrapolate the fit. Defaults to 100.
<code>lwd</code>	Line width of the smoother. Passed to <code>geom_line</code> .
<code>size</code>	Character expansion of the data points. Passed to <code>geom_point</code> .
<code>xlab</code>	x-axis label. Passed to <code>labs</code> .
<code>ylab</code>	y-axis label. Passed to <code>labs</code> .
<code>border</code>	Logical: should a white border be drawn around the mean smoother.
<code>alpha</code>	Numeric between 0 and 1, determines the transparency of data points, see <code>scale_color_viridis_d</code> .
<code>sample</code>	Numeric between 0 and 1, use to subsample the cells when there are too many so that it can plot faster.
<code>counts</code>	The matrix of gene expression counts.
<code>gene</code>	Gene name or row in count matrix of gene to plot.
<code>pointCol</code>	Plotting colors for each cell. Can be either character vector of length 1, denoting a variable in the <code>colData(models)</code> to color cells by, or a vector of length equal to the number of cells.
<code>curvesCols</code>	Plotting colors for each curve Should be a list of colors of the exact same length as the number of curves, i.e. the number of lineages (if there is no conditions) or the number of lineages by the number of conditions. In the second case, the colors are grouped by condition (lineage 1 - condition 1, lineage 1 - condition 2,...).
<code>plotLineages</code>	Logical, should the mean smoothers for each lineage be plotted?

## Value

A `ggplot` object

## Examples

```

set.seed(82)
library(ggplot2)
data(crv, package="traviz")
data(counts, package="traviz")
data(sce, package="traviz")

```

```

plotSmoothers(sce, counts, rownames(counts)[1])
# show only one curve
curvesCols <- c("#440154FF", "transparent")
plotSmoothers(sce, counts, rownames(counts)[1], curvesCols = curvesCols,
border = FALSE)
# Show only first curve and cells assigned to first lineage
plotSmoothers(sce, counts, rownames(counts)[1], curvesCols = curvesCols,
border = FALSE) +
  ggplot2::scale_color_manual(values = curvesCols)

```

---

sce

*A SingleCellExperiment object, used for testing.*


---

## Description

This object contains fitted smoothers using tradeSeq.

## Usage

```
data("sce", package = "traviz")
```

## Format

An object of class SingleCellExperiment with 240 rows and 2660 columns.

## Details

#\* @references Franziska Paul, Yaara Arkin, Amir Giladi, Diego Adhemar Jaitin, Ephraim Kenigsberg, Hadas KerenShaul, Deborah Winter, David Lara-Astiaso, Meital Gury, Assaf Weiner, Eyal David, Nadav Cohen, Felicia Kathrine Bratt Lauridsen, Simon Haas, Andreas Schlitzer, Alexander Mildner, Florent Ginhoux, Steen Jung, Andreas Trumpp, Bo Torben Porse, Amos Tanay, and Ido Amit. Transcriptional Heterogeneity and Lineage Commitment in Myeloid Progenitors. *Cell*, 163(7):1663–1677, 12 2015. ISSN 0092- 8674. doi: 10.1016/J.CELL.2015.11.013. URL <https://www.sciencedirect.com/science/article/pii/S0092867415014932?via=ih>

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