

# Package ‘NPARC’

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**Type** Package

**Title** Non-parametric analysis of response curves for thermal proteome profiling experiments

**Version** 1.21.0

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**Description** Perform non-parametric analysis of response curves as described by Childs, Bach, Franken et al. (2019): Non-parametric analysis of thermal proteome profiles reveals novel drug-binding proteins.

**License** GPL-3

**Encoding** UTF-8

**LazyData** true

**RoxygenNote** 6.1.1

**Depends** R (>= 4.0.0)

**Imports** dplyr, tidyr, BiocParallel, broom, MASS, rlang, magrittr,  
stats, methods

**Suggests** testthat, devtools, knitr, rprojroot, rmarkdown, ggplot2,  
BiocStyle

**VignetteBuilder** knitr

**biocViews** Software, Proteomics

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|                  |                          |
|------------------|--------------------------|
| fitSingleSigmoid | <i>Fit sigmoid model</i> |
|------------------|--------------------------|

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## Description

Fit sigmoid model

## Usage

```
fitSingleSigmoid(x, y, start = c(Pl = 0, a = 550, b = 10))
```

## Arguments

|       |                                                                                       |
|-------|---------------------------------------------------------------------------------------|
| x     | numeric vector of the independent variables (typically temperature)                   |
| y     | numeric vector of the dependent variables (typically relative abundance measurements) |
| start | numeric vector of start parameters for the melting curve equation                     |

## Details

Fits the following function to the data:  $y = (1 - Pl)/(1 + \exp((b - a/x))) + Pl$

## Value

model summary of type "nls"

## Examples

```
data(stauro_TPP_data_tidy)
stk4 <- dplyr::filter(stauro_TPP_data_tidy, grepl("STK4", uniqueID))
fitSingleSigmoid(stk4$temperature, stk4$relAbundance)
```

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`getParams`*Control parameters for model fitting*

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**Description**

Control parameters for model fitting

**Usage**

```
getParams(start = c(P1 = 0, a = 550, b = 10), maxAttempts = 100)
```

**Arguments**

|                          |                                                                   |
|--------------------------|-------------------------------------------------------------------|
| <code>start</code>       | Numeric vector of start parameters for the melting curve equation |
| <code>maxAttempts</code> | Number of resampling steps in case of unsuccessful model fits     |

**Value**

list of two elements: 1) "start" listing the starting parameters for melting curve fitting, 2) "maxAttempts" listing the maximal number of attempts the fit should be allowed

**Examples**

```
data(stauro_TPP_data_tidy)
df <- dplyr::filter(stauro_TPP_data_tidy, grepl("MAPK|ATP|CDK|GTP|CRK", uniqueID))
testResults <- runNPARC(x = df$temperature,
  y = df$relAbundance,
  id = df$uniqueID,
  groupsAlt = df$compoundConcentration,
  dfType = "empirical",
  control = getParams(maxAttempts = 50))
```

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`NPARC`*NPARC package*

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**Description**

Non-parametric analysis of response curves

**Details**

See the preprint on [Childs, Bach, Franken et al. \(2019\): Non-parametric analysis of thermal proteome profiles reveals novel drug-binding proteins](#)



NPARCtest

*Perform F-test***Description**

Perform F-test

**Usage**

```
NPARCtest(modelMetrics, dfType = c("empirical", "theoretical"))
```

**Arguments**

**modelMetrics** data.frame with results of the model fit in long format.

**dfType** character value indicating the method for degrees of freedom computation for the F-test. Theoretical yields the text-book solution. Empirical yields estimates derived from the distribution moments of the RSS.

**Value**

data frame with fitted model parameters and additional columns listing e.g. residuals sum of squares of null and alternative model and raw and adjusted p values retrieved from testing

**Examples**

```
data(stauro_TPP_data_tidy)
df <- dplyr::filter(stauro_TPP_data_tidy, grepl("CDK|GTP|CRK", uniqueID))
fits <- NPARCfit(x = df$temperature,
                y = df$relAbundance,
                id = df$uniqueID,
                groupsNull = NULL,
                groupsAlt = df$compoundConcentration,
                returnModels = FALSE)
modelMetrics <- fits$metrics
testRes <- NPARCtest(modelMetrics, dfType = "theoretical")
```

runNPARC

*Non-parametric analysis of response curves***Description**

Wrapper function for melting curve fitting and hypothesis testing.

**Usage**

```
runNPARC(x, y, id, groupsNull = NULL, groupsAlt,
        BPPARAM = BiocParallel::SerialParam(progressbar = TRUE),
        dfType = c("theoretical", "empirical"), control = getParams())
```

**Arguments**

|                         |                                                                                                                                                                                                                  |
|-------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <code>x</code>          | numeric vector of the independent variables (typically temperature)                                                                                                                                              |
| <code>y</code>          | numeric vector of the dependent variables (typically relative abundance measurements)                                                                                                                            |
| <code>id</code>         | character vector with the protein ID to which each data point belongs.                                                                                                                                           |
| <code>groupsNull</code> | one or more vectors with grouping variables for the null models. See details.                                                                                                                                    |
| <code>groupsAlt</code>  | one or more vectors with grouping variables for the alternative models. See details.                                                                                                                             |
| <code>BPPARAM</code>    | BiocParallel parameter object to invoke curve fitting in parallel. Default: <code>BiocParallel::SerialParam()</code>                                                                                             |
| <code>dfType</code>     | character value indicating the method for degrees of freedom computation for the F-test. Theoretical yields the text-book solution. Empirical yields estimates derived from the distribution moments of the RSS. |
| <code>control</code>    | list of parameters used to control specific parts of the analyse                                                                                                                                                 |

**Details**

`groupsNull` or `groupsAlt` can either be a single vector each, or data.frames of the same length as `x` and `y` with one column per factor

**Value**

data frame with fitted model parameters and additional columns listing e.g. residuals sum of squares of null and alterantive model

**Examples**

```
data(stauro_TPP_data_tidy)
df <- dplyr::filter(stauro_TPP_data_tidy, grepl("CDK|GTP|CRK", uniqueID))
testResults <- runNPARC(x = df$temperature,
  y = df$relAbundance,
  id = df$uniqueID,
  groupsAlt = df$compoundConcentration,
  dfType = "empirical")
```

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`stauro_TPP_data_tidy`    *TPP dataset of staurosporine treated cells.*

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**Description**

Data from a thermal proteome profiling (TPP) experiment investigating the ATP-competitive pan-kinase inhibitor staurosporine on K562 cells. The data has been downloaded the data from the supplement of the respective publication and converted into tidy format.

**Usage**

```
data(stauro_TPP_data_tidy)
```

### **Format**

An object of class "data.frame"

### **References**

Savitski et al. (2014): Tracking cancer drugs in living cells by thermal profiling of the proteome. Science 346, 1255784.

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