## Package 'vsn'

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Title Variance stabilization and calibration for microarray data

Author Wolfgang Huber, with contributions from Anja von Heydebreck. Many comments and suggestions by users are acknowledged, among them Dennis Kostka, David Kreil, Hans-Ulrich Klein, Robert Gentleman, Deepayan Sarkar and Gordon Smyth

Maintainer Wolfgang Huber <wolfgang.huber@embl.org>

**Depends** R (>= 4.0.0), methods, Biobase

Imports affy, limma, lattice, ggplot2

**Suggests** affydata, hgu95av2cdf, BiocStyle, knitr, rmarkdown, dplyr, testthat

## **Description**

The package implements a method for normalising microarray intensities from single- and multiple-color arrays. It can also be used

for data from other technologies, as long as they have similar format. The method uses a robust variant of the maximum-likelihood estimator for an additive-multiplicative error model and affine calibration. The model incorporates data calibration step (a.k.a. normalization), a model for the dependence of the variance on the mean intensity and a variance stabilizing data transformation. Differences between transformed intensities are analogous to ``normalized log-ratios". However, in contrast to the latter, their variance is independent of the mean, and they are usually more sensitive and specific in detecting differential transcription.

Reference [1] Variance stabilization applied to microarray data calibration and to the quantification of differential expression, Wolfgang Huber, Anja von Heydebreck, Holger Sueltmann, Annemarie Poustka, Martin Vingron; Bioinformatics (2002) 18 Suppl1 S96-S104. [2] Parameter estimation for the calibration and variance stabilization of microarray data, Wolfgang Huber, Anja von Heydebreck, Holger Sueltmann, Annemarie Poustka, and Martin Vingron; Statistical Applications in Genetics and Molecular Biology (2003) Vol. 2 No. 1, Article 3; http://www.bepress.com/sagmb/vol2/iss1/art3.

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URL http://www.r-project.org, http://www.ebi.ac.uk/huber

biocViews Microarray, OneChannel, TwoChannel, Preprocessing

2 vsn-package

#### VignetteBuilder knitr

Collate AllClasses.R AllGenerics.R vsn2.R vsnLogLik.R justvsn.R methods-vsnInput.R methods-vsn.R methods-vsn2.R methods-predict.R RGList\_to\_NChannelSet.R meanSdPlot-methods.R plotLikelihood.R normalize.AffyBatch.vsn.R sagmbSimulateData.R 777 R

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## **Contents**

	package <i>vsn</i>	
Index		19
	vsnInput	17
	vsn2trsf	
	vsn2	13
	vsn	12
	scalingFactorTransformation	11
	sagmbSimulateData	10
	normalize.AffyBatch.vsn	
	meanSdPlot	
	lymphoma	
	logLik-methods	
	kidney	
	justvsn	
	vsn-package	2

## **Description**

vsn

#### **Details**

The main function of the package is vsn2. Interesting for its applications are also predict and the wrapper function justvsn.

vsn2 can be applied to objects of class ExpressionSet, NChannelSet, AffyBatch (from the affy package) and RGList (from the limma package), matrix and vector. It returns an object of class vsn, which contains the results of fitting the vsn model to the data.

The most common use case is that you will want to construct a new data object with the vsn-normalized data whose class is the same as that of the input data and which preserves the metadata. This can be achieved by

justvsn 3

```
fit = vsn2(x, ...)
nx = predict(fit, newdata=x)
```

To simplify this, there exists also a simple wrapper justvsn.

#### Author(s)

Wolfgang Huber

justvsn

Wrapper functions for vsn

## **Description**

```
justvsn is equivalent to calling
  fit = vsn2(x, ...)
  nx = predict(fit, newdata=x, useDataInFit = TRUE)
vsnrma is a wrapper around vsn2 and rma.
```

#### Usage

```
justvsn(x, ...)
vsnrma(x, ...)
```

## Arguments

x For justvsn, any kind of object for which vsn2 methods exist. For vsnrma, an AffyBatch.

... Further arguments that get passed on to vsn2.

#### **Details**

vsnrma does probe-wise background correction and between-array normalization by calling vsn2 on the perfect match (PM) values only. Probeset summaries are calculated with the medianpolish algorithm of rma.

## Value

justvsn returns the vsn-normalised intensities in an object generally of the same class as its first argument (see the man page of predict for details). It preserves the metadata.

vsnrma returns an ExpressionSet.

#### Author(s)

Wolfgang Huber

#### See Also

vsn2

4 kidney

#### **Examples**

#### **Description**

Intensity data for one cDNA slide with two adjacent tissue samples from a nephrectomy (kidney)

#### Usage

```
data(kidney)
```

## **Format**

kidney is an ExpressionSet containing the data from one cDNA chip. The 8704x2 matrix exprs(kidney) contains the spot intensities for the red (635 nm) and green color channels (532 nm) respectively. For each spot, a background estimate from a surrounding region was subtracted.

#### **Details**

The chip was produced in 2001 by Holger Sueltmann at the Division of Molecular Genome Analysis at the German Cancer Research Center in Heidelberg.

#### References

Huber W, Boer JM, von Heydebreck A, Gunawan B, Vingron M, Fuzesi L, Poustka A, Sueltmann H. Transcription profiling of renal cell carcinoma. Verh Dtsch Ges Pathol. 2002;86:153-64. PMID: 12647365

## **Examples**

```
data("kidney")
plot(exprs(kidney), pch = ".", log = "xy")
abline(a = 0, b = 1, col = "blue")
```

logLik-methods 5

logLik-methods Calculate the log likelihood and its gradient for the vsn model	logLik-methods	Calculate the log likelihood and its gradient for the vsn model	
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## Description

logLik calculates the log likelihood and its gradient for the vsn model. plotVsnLogLik makes a false color plot for a 2D section of the likelihood landscape.

## Usage

## **Arguments**

object	A vsnInput object.
p	For plotVsnLogLik, a vector or a 3D array with the point in parameter space around which to plot the likelihood. For logLik, a matrix whose columns are the set of parameters at which the likelihoods are to be evaluated.
mu	Numeric vector of length 0 or nrow(object). If the length is 0, there is no reference and sigsq must be NA (the default value). See vsn2.
sigsq	Numeric scalar.
calib	as in vsn2.
whichp	Numeric vector of length 2, with the indices of those two parameters in p along which the section is to be taken.
expand	Numeric vector of length 1 or 2 with expansion factors for the plot range. The range is auto-calculated using a heuristic, but manual adjustment can be useful; see example.
ngrid	Integer scalar, the grid size.
fun	Function to use for log-likelihood calculation. This parameter is exposed only for testing purposes.
main	This parameter is passed on levelplot.
	Arguments that get passed on to fun, use this for mu, sigsq, calib.

## **Details**

logLik is an R interface to the likelihood computations in vsn (which are done in C).

6 lymphoma

#### Value

For logLik, a numeric matrix of size nrow(p)+1 by ncol(p). Its columns correspond to the columns of p. Its first row are the likelihood values, its rows 2...nrow(p)+1 contain the gradients. If mu and sigsq are specified, the ordinary negative log likelihood is calculated using these parameters as given. If they are not specified, the profile negative log likelihood is calculated.

For plotVsnLogLik, a dataframe with the 2D grid coordinates and log likelihood values.

## Author(s)

Wolfgang Huber

#### See Also

vsn2

#### **Examples**

```
data("kidney")
v = new("vsnInput", x=exprs(kidney),
    pstart=array(as.numeric(NA), dim=c(1, ncol(kidney), 2)))
fit = vsn2(kidney)
print(coef(fit))

p = sapply(seq(-1, 1, length=31), function(f) coef(fit)+c(0,0,f,0))

1l = logLik(v, p)

plot(p[3, ], ll[1, ], type="l", xlab=expression(b[1]), ylab=expression(-log(L)))
abline(v=coef(fit)[3], col="red")

plotVsnLogLik(v, coef(fit), whichp=c(1,3), expand=0.2)
```

lymphoma

Intensity data for 8 cDNA slides with CLL and DLBL samples from the Alizadeh et al. paper in Nature 2000

#### **Description**

8 cDNA chips from Alizadeh lymphoma paper

#### Usage

```
data(lymphoma)
```

## **Format**

lymphoma is an ExpressionSet containing the data from 8 chips from the lymphoma data set by Alizadeh et al. (see references). Each chip represents two samples: on color channel 1 (CH1, Cy3, green) the common reference sample, and on color channel 2 (CH2, Cy5, red) the various disease samples. See pData(lymphoma). The 9216x16 matrix exprs(lymphoma) contains the background-subtracted spot intensities (CH1I-CH1B and CH2I-CH2B, respectively).

meanSdPlot 7

#### **Details**

The chip intensity files were downloaded from the Stanford microarray database. Starting from the link below, this was done by following the links *Published Data -> Alizadeh AA*, et al. (2000) *Nature 403(6769):503-11 -> Data in SMD -> Display Data*, and selecting the following 8 slides:

lc7b019 lc7b047 lc7b048 lc7b056 lc7b057 lc7b058 lc7b069 lc7b070

Then, the script makedata. R from the scripts subdirectory of this package was run to generate the R data object.

#### Source

http://genome-www5.stanford.edu/MicroArray/SMD

#### References

A. Alizadeh et al., Distinct types of diffuse large B-cell lymphoma identified by gene expression profiling. Nature 403(6769):503-11, Feb 3, 2000.

## **Examples**

```
data("lymphoma")
lymphoma
pData(lymphoma)
```

meanSdPlot

Plot row standard deviations versus row means

## **Description**

Methods for objects of classes matrix, ExpressionSet, vsn and MAList to plot row standard deviations versus row means.

## Usage

8 meanSdPlot

#### **Arguments**

X	An object of class matrix, ExpressionSet, vsn or MAList.
ranks	Logical, indicating whether the x-axis (means) should be plotted on the original scale (FALSE) or on the rank scale (TRUE). The latter distributes the data more evenly along the x-axis and allows a better visual assessment of the standard deviation as a function of the mean.
xlab	Character, label for the x-axis.
ylab	Character, label for the y-axis.
pch	Ignored - exists for backward compatibility.
plot	Logical. If TRUE (default), a plot is produced. Calling the function with plot=FALSE can be useful if only its return value is of interest.
bins	Gets passed on to geom_hex.
	Further arguments that get passed on to geom_hex.

#### **Details**

Standard deviation and mean are calculated row-wise from the expression matrix (in) x. The scatterplot of these versus each other allows you to visually verify whether there is a dependence of the standard deviation (or variance) on the mean. The red line depicts the running median estimator (window-width 10%). If there is no variance-mean dependence, then the line should be approximately horizontal.

## Value

A named list with five components: its elements px and py are the x- and y-coordinates of the individual data points in the plot; its first and second element are the x-coordinates and values of the running median estimator (the red line in the plot). Its element gg is the plot object (see examples). Depending on the value of plot, the method can (and by default does) have a side effect, which is to print gg on the active graphics device.

## Author(s)

Wolfgang Huber

#### **Examples**

```
data("kidney")
log.na <- function(x) log(ifelse(x>0, x, NA))

exprs(kidney) <- log.na(exprs(kidney))
msd <- meanSdPlot(kidney)

## The `ggplot` object is returned in list element `gg`, here is an example of how to modify the plot
library("ggplot2")
msd$gg + ggtitle("Hello world") + scale_fill_gradient(low = "yellow", high = "darkred") + scale_y_continuous(
## Try this out with not log-transformed data, vsn2-transformed data, the lymphoma data, your data ...</pre>
```

```
normalize.AffyBatch.vsn
```

Wrapper for vsn to be used as a normalization method with expresso

#### **Description**

Wrapper for vsn2 to be used as a normalization method with the expresso function of the package affy. The expresso function is deprecated, consider using justvsn instead. The normalize.AffyBatch.vsn can still be useful on its own, as it provides some additional control of the normalization process (fitting on subsets, alternate transform parameters).

## Usage

```
normalize.AffyBatch.vsn(
   abatch,
   reference,
   strata = NULL,
   subsample = if (nrow(exprs(abatch))>30000L) 30000L else 0L,
   subset,
   log2scale = TRUE,
   log2asymp=FALSE,
   ...)
```

#### **Arguments**

abatch An object of type AffyBatch.

reference Optional, a 'vsn' object from a previous fit. If this argument is specified, the

data in 'x' are normalized "towards" an existing set of reference arrays whose parameters are stored in the object 'reference'. If this argument is not specified, then the data in 'x' are normalized "among themselves". See vsn2 for details.

strata The 'strata' functionality is not supported, the parameter is ignored.

subsample Is passed on to vsn2.

subset This allows the specification of a subset of expression measurements to be used

for the vsn fit. The transformation with the parameters of this fit is then, however, applied to the whole dataset. This is useful for excluding expression measurements that are known to be differentially expressed or control probes that may not match the vsn model, thus avoiding that they influence the normalization process. This operates at the level of probesets, not probes. Both 'subset'

and 'subsample' can be used together.

log2scale If TRUE, this will perform a global affine transform on the data to put them

on a similar scale as the original non-transformed data. Many users prefer this. Fold-change estimates are not affected by this transform. In some situations, however, it may be helpful to turn this off, e.g., when comparing independently

normalized subsets of the data.

log2asymp If TRUE, this will perform a global affine transform on the data to make the gen-

eralized log (asinh) transform be asymptotically identical to a log base 2 transform. Some people find this helpful. Only **one** of 'log2scale' or 'log2asymp' can be set to TRUE. Fold-change estimates are not affected by this transform.

... Further parameters for vsn2.

10 sagmbSimulateData

#### **Details**

Please refer to the *Details* and *References* sections of the man page for vsn2 for more details about this method.

**Important note**: after calling vsn2, the function normalize. AffyBatch.vsn **exponentiates** the data (base 2). This is done in order to make the behavior of this function similar to the other normalization methods in affy. That packages uses the convention of taking the logarithm to base in subsequent analysis steps (e.g. in medpolish).

## Value

An object of class AffyBatch. The vsn object returned, which can be used as reference for subsequent fits, is provided by description(abatch)@preprocessing\$vsnReference.

#### Author(s)

```
D. P. Kreil http://bioinf.boku.ac.at/, Wolfgang Huber
```

#### See Also

vsn2

#### **Examples**

## Please see vignette.

sagmbSimulateData

Simulate data and assess vsn's parameter estimation

## **Description**

Functions to validate and assess the performance of vsn through simulation of data.

## Usage

```
sagmbSimulateData(n=8064, d=2, de=0, up=0.5, nrstrata=1, miss=0, log2scale=FALSE) \\ sagmbAssess(h1, sim)
```

#### **Arguments**

n	Numeric. Number of probes (rows).
d	Numeric. Number of arrays (columns).
de	Numeric. Fraction of differentially expressed genes.
up	Numeric. Fraction of up-regulated genes among the differentially expressed genes.
nrstrata	Numeric. Number of probe strata.
miss	Numeric. Fraction of data points that is randomly sampled and set to NA.
log2scale	Logical. If TRUE, glog on base 2 is used, if FALSE, (the default), then base e.
h1	Matrix. Calibrated and transformed data, according, e.g., to vsn
sim	List. The output of a previous call to sagmbSimulateData, see Value

#### **Details**

Please see the vignette.

#### Value

For sagmbSimulateData, a list with four components: hy, an  $n \times d$  matrix with the true (=simulated) calibrated, transformed data; y, an  $n \times d$  matrix with the simulated uncalibrated raw data this is intended to be fed into vsn2; is.de, a logical vector of length n, specifying which probes are simulated to be differentially expressed. strata, a factor of length n.

For sagmbSimulateData, a number: the root mean squared difference between true and estimated transformed data.

## Author(s)

Wolfgang Huber

#### References

Wolfgang Huber, Anja von Heydebreck, Holger Sueltmann, Annemarie Poustka, and Martin Vingron (2003) "Parameter estimation for the calibration and variance stabilization of microarray data", Statistical Applications in Genetics and Molecular Biology: Vol. 2: No. 1, Article 3. http://www.bepress.com/sagmb/vol2/iss1/art3

## **Examples**

```
sim <- sagmbSimulateData(nrstrata = 4)
ny <- vsn2(sim$y, strata = sim$strata)
res <- sagmbAssess(exprs(ny), sim)
res</pre>
```

scalingFactorTransformation

The transformation that is applied to the scaling parameter of the vsn model

#### **Description**

The transformation that is applied to the scaling parameter of the vsn model

#### Usage

```
scalingFactorTransformation(b)
```

## **Arguments**

b Real vector.

## Value

A real vector of same length as b, with transformation f applied (see vignette *Likelihood Calculations for vsn*).

12 vsn

#### Author(s)

Wolfgang Huber

#### **Examples**

```
b = seq(-3, 2, length=20)
fb = scalingFactorTransformation(b)
if(interactive())
  plot(b, fb, type="b", pch=16)
```

vsn

Class to contain result of a vsn fit

#### **Description**

Class to contain result of a vsn fit

## **Creating Objects**

```
new("vsn") vsn2(x) with x being an ExpressionSet.
```

## **Slots**

coefficients: A 3D array of size (number of strata) x (number of columns of the data matrix) x 2. It contains the fitted normalization parameters (see vignette).

strata: A factor of length 0 or n. If its length is n, then its levels correspond to different normalization strata (see vignette).

mu: A numeric vector of length n with the fitted parameters  $\hat{\mu}_k$ , for k=1,...,n.

sigsq: A numeric scalar,  $\hat{\sigma}^2$ .

hx: A numeric matrix with 0 or n rows. If the number of rows is n, then hx contains the transformed data matrix.

1bfgsb: An integer scalar containing the return code from the L-BFGS-B optimizer.

hoffset: Numeric scalar, the overall offset c- see manual page of vsn2.

calib: Character of length 1, see manual page of vsn2.

## Methods

Γ Subset

dim Get dimensions of data matrix.

nrow Get number of rows of data matrix.

ncol Get number of columns of data matrix.

show Print a summary of the object

exprs Accessor to slot hx.

coef, coefficients Accessors to slot coefficients.

#### Author(s)

Wolfgang Huber

vsn2 13

#### See Also

vsn2

#### **Examples**

```
data("kidney")
v = vsn2(kidney)
show(v)
dim(v)
v[1:10, ]
```

vsn2

Fit the vsn model

#### **Description**

vsn2 fits the vsn model to the data in x and returns a vsn object with the fit parameters and the transformed data matrix. The data are, typically, feature intensity readings from a microarray, but this function may also be useful for other kinds of intensity data that obey an additive-multiplicative error model. To obtain an object of the same class as x, containing the normalised data and the same metdata as x, use

```
fit = vsn2(x, ...)
nx = predict(fit, newdata=x)
```

or the wrapper justvsn. Please see the vignette Introduction to vsn.

#### Usage

```
vsnMatrix(x,
          reference,
          strata,
          lts.quantile = 0.9,
          subsample
                    = 0L,
          verbose
                       = interactive(),
          returnData = TRUE,
                      = "affine",
          calib
          pstart,
          minDataPointsPerStratum = 42L,
          optimpar
                      = list(),
          defaultpar = list(factr=5e7, pgtol=2e-4, maxit=60000L,
                              trace=0L, cvg.niter=7L, cvg.eps=0))
## S4 method for signature 'ExpressionSet'
vsn2(x, reference, strata, ...)
## S4 method for signature 'AffyBatch'
vsn2(x, reference, strata, subsample, ...)
## S4 method for signature 'NChannelSet'
```

14 vsn2

#### **Arguments**

x An object containing the data to which the model is fitted.

reference Optional, a vsn object from a previous fit. If this argument is specified, the

data in x are normalized "towards" an existing set of reference arrays whose parameters are stored in the object reference. If this argument is not specified, then the data in x are normalized "among themselves". See Details for a more

precise explanation.

strata Optional, a factor or integer whose length is nrow(x). It can be used for

stratified normalization (i.e. separate offsets a and factors b for each level of strata). If missing, all rows of x are assumed to come from one stratum. If strata is an integer, its values must cover the range  $1, \ldots, n$ , where n is the

number of strata.

lts.quantile Numeric of length 1. The quantile that is used for the resistant least trimmed

sum of squares regression. Allowed values are between 0.5 and 1. A value of 1

corresponds to ordinary least sum of squares regression.

subsample Integer of length 1. If its value is greater than 0, the model parameters are esti-

mated from a subsample of the data of size subsample only, yet the fitted transformation is then applied to all data. For large datasets, this can substantially reduce the CPU time and memory consumption at a negligible loss of precision. Note that the AffyBatch method of vsn2 sets a value of 30000 for this parameter if it is missing from the function call - which is different from the behaviour

of the other methods.

backgroundsubtract

Logical of length 1: should local background estimates be subtracted before fitting vsn?

foreground, background

Aligned character vectors of the same length, naming the channels of x that

should be used as foreground and background values.

verbose Logical. If TRUE, some messages are printed.

returnData Logical. If TRUE, the transformed data are returned in a slot of the resulting

vsn object. Setting this option to FALSE allows saving memory if the data are

not needed.

calib Character of length 1. Allowed values are affine and none. The default,

affine, corresponds to the behaviour in package versions <= 3.9, and to what is described in references [1] and [2]. The option none is an experimental new feature, in which no affine calibration is performed and only two global variance stabilisation transformation parameters a and b are fitted. This functionality might be useful in conjunction with other calibration methods, such as quantile

normalisation - see the vignette Introduction to vsn.

Optional, a three-dimensional numeric array that specifies start values for the iterative parameter estimation algorithm. If not specified, the function tries

to guess useful start values. The first dimension corresponds to the levels of  $\mathsf{strata}$ , the second dimension to the columns of  $\mathsf{x}$  and the third dimension must

be 2, corresponding to offsets and factors.

vsn2 15

minDataPointsPerStratum

The minimum number of data points per stratum. Normally there is no need for the user to change this; refer to the vignette for further documentation.

optimpar Optional, a list with parameters for the likelihood optimisation algorithm. De-

fault parameters are taken from defaultpar. See details.

defaultpar The default parameters for the likelihood optimisation algorithm. Values in

optimpar take precedence over those in defaultpar. The purpose of this argument is to expose the default values in this manual page - it is not intended to be

changed, please use optimpar for that.

... Arguments that get passed on to vsnMatrix.

#### Value

An object of class vsn.

#### Note on overall scale and location of the glog transformation

The data are returned on a glog scale to base 2. More precisely, the transformed data are subject to the transformation  $glog_2(f(b)*x+a)+c$ , where the function  $glog_2(u)=log_2(u+\sqrt{u*u+1})=asinh(u)/\log(2)$  is called the generalised logarithm, the offset a and the scaling parameter b are the fitted model parameters (see references), and  $f(x)=\exp(x)$  is a parameter transformation that allows ensuring positivity of the factor in front of x while using an unconstrained optimisation over b [4]. The overall offset c is computed from the b's such that for large x the transformation approximately corresponds to the  $\log_2$  function. This is done separately for each stratum, but with the same value across arrays. More precisely, if the element b[s,i] of the array b is the scaling parameter for the s-th stratum and the i-th array, then c[s] is computed as log2(2\*f(mean(b[,i]))). The offset c is inconsequential for all differential expression calculations, but many users like to see the data in a range that they are familiar with.

#### Specific behaviour of the different methods

vsn2 methods exist for ExpressionSet, NChannelSet, AffyBatch (from the affy package), RGList (from the limma package), matrix and numeric. If x is an NChannelSet, then vsn2 is applied to the matrix that is obtained by horizontally concatenating the color channels. Optionally, available background estimates can be subtracted before. If x is an RGList, it is converted into an NChannelSet using a copy of Martin Morgan's code for RGList to NChannelSet coercion, then the NChannelSet method is called.

#### Standalone versus reference normalisation

If the reference argument is *not* specified, then the model parameters  $\mu_k$  and  $\sigma$  are fit from the data in x. This is the mode of operation described in [1] and that was the only option in versions 1.X of this package. If reference is specified, the model parameters  $\mu_k$  and  $\sigma$  are taken from it. This allows for 'incremental' normalization [4].

#### Convergence of the iterative likelihood optimisation

L-BFGS-B uses three termination criteria:

- 1.  $(f_k f_{k+1}) / \max(|f_k|, |f_{k+1}|, 1) \le factr * epsmch where epsmch is the machine precision.$
- 2. |gradient| < pgtol
- 3. iterations > maxit

16 vsn2trsf

These are set by the elements factr, pgtol and maxit of optimpar. The remaining elements are

trace An integer between 0 and 6, indicating the verbosity level of L-BFGS-B, higher values create more output.

cvg.niter The number of iterations to be used in the least trimmed sum of squares regression.

cvg.eps Numeric. A convergence threshold for the least trimmed sum of squares regression.

## Author(s)

Wolfgang Huber

#### References

- [1] Variance stabilization applied to microarray data calibration and to the quantification of differential expression, Wolfgang Huber, Anja von Heydebreck, Holger Sueltmann, Annemarie Poustka, Martin Vingron; Bioinformatics (2002) 18 Suppl.1 S96-S104.
- [2] Parameter estimation for the calibration and variance stabilization of microarray data, Wolfgang Huber, Anja von Heydebreck, Holger Sueltmann, Annemarie Poustka, and Martin Vingron; Statistical Applications in Genetics and Molecular Biology (2003) Vol. 2 No. 1, Article 3. http://www.bepress.com/sagmb/vol2/
- [3] L-BFGS-B: Fortran Subroutines for Large-Scale Bound Constrained Optimization, C. Zhu, R.H. Byrd, P. Lu and J. Nocedal, Technical Report, Northwestern University (1996).
- [4] Package vignette: Likelihood Calculations for vsn

#### See Also

justvsn, predict

#### **Examples**

```
data("kidney")

fit = vsn2(kidney)  ## fit
nkid = predict(fit, newdata=kidney) ## apply fit

plot(exprs(nkid), pch=".")
abline(a=0, b=1, col="red")
```

vsn2trsf

Apply the vsn transformation to data

#### **Description**

Apply the vsn transformation to data.

## Usage

```
## S4 method for signature 'vsn'
predict(object, newdata, strata=object@strata, log2scale=TRUE, useDataInFit=FALSE)
```

vsnInput 17

#### **Arguments**

object An object of class vsn that contains transformation parameters and strata infor-

mation, typically this is the result of a previous call to vsn2.

newdata Object of class ExpressionSet, NChannelSet, AffyBatch (from the affy pack-

age),  $\operatorname{\mathsf{RGList}}$  (from the limma package),  $\operatorname{\mathsf{matrix}}$  or numeric, with the data to

which the fit is to be applied to.

Strata Optional, a factor or integer that aligns with the rows of newdata; see the

strata argument of vsn2.

log2scale If TRUE, the data are returned on the glog scale to base 2, and an overall offset

c is added (see  $\mathit{Value}$  section of the  $\mathsf{vsn2}$  manual page). If FALSE, the data are

returned on the glog scale to base e, and no offset is added.

transferred appropriately into resulting object, which otherwise preserves the class and metadata of newdata. This option exists to increase performance in

constructs like

```
fit = vsn2(x, ...)
nx = predict(fit, newdata=x)
```

and is used, for example, in the justvsn function.

#### Value

An object typically of the same class as newdata. There are two exceptions: if newdata is an RGList, the return value is an NChannelSet, and if newdata is numeric, the return value is a matrix with 1 column.

## Author(s)

Wolfgang Huber

#### **Examples**

```
data("kidney")
## nb: for random subsampling, the 'subsample' argument of vsn
## provides an easier way to do this
fit = vsn2(kidney[sample(nrow(kidney), 500), ])
tn = predict(fit, newdata=exprs(kidney))
```

vsnInput

Class to contain input data and parameters for vsn functions

## Description

Class to contain input data and parameters for vsn functions

#### **Creating Objects**

```
new("vsnInput")
```

18 vsnInput

#### **Slots**

x: A numeric matrix with the input data.

reference: An object of vsn, typically this would have been obtained from a previous fit to a set of reference arrays (data).

strata: A factor of length 0 or n. If its length is n, then its levels correspond to different normalization strata (see vsn2).

ordered: Logical scalar; are the rows reordered so that the strata are contiguous.

lts.quantile: Numeric scalar, seevsn2.

subsample: Integer scalar, seevsn2.

verbose: Logical scalar, seevsn2.

calib Character of length 1, see manual page of vsn2.

pstart: A 3D array of size (number of strata) x (number of columns of the data matrix) x 2. It contains the start parameters.

optimpar: List with parameters for the numerical optimiser L-BFGS-B; see the manual page of vsn2.

#### Methods

[ Subset

dim Get dimensions of data matrix.

nrow Get number of rows of data matrix.

ncol Get number of columns of data matrix.

show Print a summary of the object

#### Author(s)

Wolfgang Huber

#### See Also

vsn2

# Index

* classes	<pre>meanSdPlot,MAList-method(meanSdPlot),^</pre>
vsn, 12	<pre>meanSdPlot,matrix-method(meanSdPlot),^</pre>
vsnInput, 17	<pre>meanSdPlot, vsn-method (meanSdPlot), 7</pre>
* datasets	<pre>meanSdPlot-methods (meanSdPlot), 7</pre>
kidney, 4	medpolish, <i>10</i>
lymphoma, 6	•
* hplot	NChannelSet, 2, 15, 17
meanSdPlot, 7	<pre>ncol, vsn-method (vsn), 12</pre>
* methods	<pre>ncol,vsnInput-method(vsnInput), 17</pre>
meanSdPlot, 7	normalize.AffyBatch.vsn,9
* package	nrow, vsn-method (vsn), 12
vsn-package, 2	nrow, vsnInput-method (vsnInput), 17
[, vsn-method (vsn), 12	, , , , , , , , , , , , , , , , , , , ,
[,vsnInput-method (vsnInput), 17	plotVsnLogLik (logLik-methods), 5
	predict, 3, 16
AffyBatch, 2, 3, 9, 10, 15, 17	predict, vsn-method (vsn2trsf), 16
class:vsn (vsn), 12	RGList, 2, 15, 17
class:vsnInput(vsnInput), 17	rma, <i>3</i>
coef, vsn-method (vsn), 12	
coefficients, vsn-method (vsn), 12	<pre>sagmbAssess (sagmbSimulateData), 10</pre>
coerce,RGList,NChannelSet-method	sagmbSimulateData, 10
(vsn2), 13	scalingFactorTransformation, 11
( ( ( ) ) , 10	show, vsn-method (vsn), 12
dim, vsn-method (vsn), 12	<pre>show, vsnInput-method (vsnInput), 17</pre>
dim, vsnInput-method (vsnInput), 17	
	vsn, 2, 7, 8, 12, 13–15, 17, 18
ExpressionSet, 2-4, 6-8, 12, 15, 17	vsn-class (vsn), 12
exprs, vsn-method (vsn), 12	vsn-package, 2
	vsn2, 2, 3, 5, 6, 9–13, 13, 17, 18
geom_hex, 8	vsn2, AffyBatch-method (vsn2), 13
justvsn, 2, 3, 3, 9, 13, 16, 17	vsn2, ExpressionSet-method (vsn2), 13
Justvsii, 2, 3, 3, 9, 13, 10, 17	vsn2, matrix-method (vsn2), 13
kidney, 4	vsn2, NChannelSet-method (vsn2), 13
,	vsn2, numeric-method (vsn2), 13
logLik,vsnInput-method	vsn2, RGList-method (vsn2), 13
(logLik-methods), 5	vsn2-methods (vsn2), 13
logLik-methods, 5	vsn2trsf, 16
lymphoma, 6	vsnInput, 5, 17
	vsnInput-class (vsnInput), 17
MAList, <i>7</i> , <i>8</i>	vsnMatrix (vsn2), 13
matrix, <i>7</i> , <i>8</i>	vsnrma (justvsn), 3
meanSdPlot, 7	· · · · · · · · · · · · · · · · · · ·
meanSdPlot,ExpressionSet-method	
(meanSdPlot), 7	