

# Package ‘affypdnn’

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**Title** Probe Dependent Nearest Neighbours (PDNN) for the affy package

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**Depends** R (>= 2.13.0), affy (>= 1.5)

**Suggests** affydata, hgu95av2probe

**Description** The package contains functions to perform the PDNN method described by Li Zhang et al.

**License** LGPL

**biocViews** OneChannel, Microarray, Preprocessing

**NeedsCompilation** no

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expressopdnn

*Position Dependant Nearest Neighbors model for affy*

## Description

A wrapper to perform the PDNN method.

## Usage

```
pdnn.scalevalue.exprSet(eset, scale.to=500)
expressopdnn(abatch,
             # background correction
             bg.correct = FALSE,
             bgcorrect.method = NULL,
             bgcorrect.param = list(),
             # normalize
             normalize = FALSE,
             normalize.method = NULL,
             normalize.param = list(),
             pmcorrect.method = c("pdnn", "pdnnpredict"),

             # pdnn
             findparams.param = list(),
             # expression values
             summary.subset = NULL,
             # PDNN expression values scaling
             eset.normalize = TRUE,
             scale.to = 500,
             # misc.
             verbose = TRUE)
```

## Arguments

<code>abatch</code>	object of <a href="#">AffyBatch-class</a> .
<code>bg.correct</code>	a boolean to express whether background correction is wanted or not.
<code>bgcorrect.method</code>	the name of the background adjustment method.
<code>bgcorrect.param</code>	a list of parameters for <code>bgcorrect.method</code> (if needed/wanted).
<code>eset</code>	an object of <a href="#">ExpressionSet-class</a> .
<code>normalize</code>	normalization step wished or not.
<code>normalize.method</code>	the normalization method to use.
<code>normalize.param</code>	a list of parameters to be passed to the normalization method (if wanted).

```

pmcorrect.method
    the name of the PM adjustement method (only two choices here, default to
    'pdnn').

findparams.param
    a list of parameters to be passed to find.params.pdnn.

eset.normalize
    is any normalization step on expression values to be performed.

scale.to
    a value to scale against.

summary.subset
    a list of 'affyids'. If NULL, then an expression summary value is computed for
    everything on the chip.

verbose
    logical value. If TRUE it writes out some messages.

```

## Details

`expressopdnn` is very similar to [expresso](#). It is mainly a wrapper around the pre-processing steps ‘background correction’, ‘normalization’, ‘perfect match correction’ and the PDNN method to compute expression values (see the first reference for more details about the preprocessing steps and and the second reference for further details about the PDNN method).

The wrapper `expresso` has no way to handle easily the computation of chip-wide results that have to be used during the `computeExprSet` step. An easy way to overcome this was to write this simple wrapper.

`pdnn.scalevalue` is performed after the expression values have computed to somehow ‘normalize’ the values between different chips. When setting `normalize` to TRUE this step might be considered unnecessary (and the `eset.normalize` set to FALSE).

## Value

An object of [ExpressionSet-class](#), with an attribute `pps.warnings` as returned by the method `computeExprSet`.

## See Also

[expresso](#) and `generateExprVal.method.pdnn`

## Examples

```

## load pre-computed parameters
data(hgu95av2.pdnn.params)

library(affydata)
data(Dilution)

## one CEL to go faster
afbatch <- Dilution[, 1]

## Take only few IDs (the 10 first)
ids <- ls(getCdfInfo(afbatch))[1:10]
eset <- expressopdnn(afbatch, bg.correct=FALSE,
                      normalize=FALSE,
                      findparams.param=list(params.chiptype=hgu95av2.pdnn.params,

```

```
give.warnings=FALSE),
summary.subset=ids)
```

**find.params.pdnn**      *A function to find the experiment specific PDNN parameters*

## Description

A function to find the paramaters specific to the chips in an AffyBatch object.

## Usage

```
find.params.pdnn(abatch, params.chiptype, optim.method = "BFGS", verbose = TRUE, give.warnings=TRUE)
```

## Arguments

abatch	an instance of <a href="#">AffyBatch-class</a> .
params.chiptype	chip-type specific parameters (see details)
optim.method	method for the optimization function <a href="#">optim</a> . When FALSE, a steepest-descent method of our own is used.
verbose	verbosity (TRUE or FALSE)
give.warnings	report probeset IDs in the abatch that could not be found in the params.chiptype

## Details

This function fits PDNN parameters that are specific to experimental values. The parameters common to all the chips of a certain type are returned by the function [pdnn.params.chiptype](#). If NULL, the parameter files included in the package will be used whenever possible...

## Value

A list of

lambda	The lambda's
Bs	The B's
Ns	The N's
Fs	The F's

## References

Li Zhang, Michael F. Miles and Kenneth D. Aldape - A model of molecular interactions on short oligonucleotide arrays, 2003, Nature Biotech., vol. 21, n.7

## See Also

[pdnn.params.chiptype](#), [generateExprVal.method.pdnn](#)

## Examples

```
## load a chip-specific parameter file
## (as returned by the function pdnn.params.chiptype)
data(hgu95av2.pdnn.params)

## load experimental data
library(affydata)
data(Dilution)

## one CEL to go faster
afbatch <- Dilution[, 1]
params <- find.params.pdnn(afbatch, hgu95av2.pdnn.params, optim.method =
FALSE, give.warnings=FALSE)
```

generateExprVal.method.pdnn

*Compute PM correction and summary expression value*

## Description

Computes PM correction and summary expression value with PDNN method.

## Usage

```
pmcorrect.pdnn(object, params, gene=NULL, gene.i=NULL,
                params.chiptype=NULL, outlierlim=3, callingFromExpresso=FALSE)
pmcorrect.pdnnpredict(object, params, gene=NULL, gene.i=NULL,
                       params.chiptype=NULL, outlierlim=3, callingFromExpresso=FALSE)
generateExprVal.method.pdnn(probes, params)
```

## Arguments

object	object of <a href="#">ProbeSet</a> .
probes	matrix of PM-corrected signals (should be coming out of <code>pmcorrect.pdnn</code> ).
params	experiments specific parameters.
gene	gene (probe set) ID (from which the <code>gene.i</code> would be derived).
gene.i	gene index (see details).
params.chiptype	chip-specific parameters.
outlierlim	threshold for tagging a probe as an outlier.
callingFromExpresso	is the function called through expresso. DO NOT play with that.

## Details

Only one of gene, gene.i should be specified. For most the users, this is gene. [pmcorrect.pdnn](#) and [pmcorrect.pdnnpredict](#) return what is called GSB and GSB + NSB + B in the paper by Zhang Li and collaborators.

## Value

[pmcorrect.pdnn](#) and [pmcorrect.pdnnpredict](#) return a matrix (one row per probe, one column per chip) with attributes attached. [generateExprVal](#) returns a list:

exprs	expression values
se.exprs	se expr. val.

## See Also

[pdnn.params.chiptype](#)

## Examples

```
data(hgu95av2.pdnn.params)
library(affydata)
data(Dilution)

## only one CEL to go faster
abatch <- Dilution[, 1]

## get the chip specific parameters
params <- find.params.pdnn(abatch, hgu95av2.pdnn.params)

## The thrill part: do we get like in the Figure 1-a of the reference ?
par(mfrow=c(2,2))
##ppset.name <- sample(featureNames(abatch), 2)
ppset.name <- c("41206_r_at", "31620_at")
ppset <- probeset(abatch, ppset.name)
for (i in 1:2) {
  ##ppset[[i]] <- transform(ppset[[i]], fun=log) # take the log as they do
  probes.pdnn <- pmcorrect.pdnnpredict(ppset[[i]], params,
                                         params.chiptype=hgu95av2.pdnn.params)
  ##probes.pdnn <- log(probes.pdnn)
  plot(ppset[[i]], main=paste(ppset.name[i], "\n(raw intensities)"))
  matplotProbesPDNN(probes.pdnn, main=paste(ppset.name[i], "\n(predicted intensities)"))
}

## pick the 50 first probeset IDs
## (to go faster)
ids <- featureNames(abatch)[1:100]

## compute the expression set (object of class 'ExpressionSet')
eset <- computeExprSet(abatch, pmcorrect.method="pdnn",
                       summary.method="pdnn", ids=ids,
                       summary.param = list(params, params.chiptype=hgu95av2.pdnn.params))
```

---

hgu133a.pdnn.params      *Chip-type specific data*

---

## Description

Chip-type specific data structure.

## Usage

```
data(hgu95av2.pdnn.params)
```

## Format

The format is a list:

**Eg** environment (one entry per dinucleotide)

**Wg** numerical vector.

**En** environment (one entry per dinucleotide)

**Wn** numerical vector.

**gene.Sn** list (one entry per affyID)

**gene.Sg** list (one entry per affyID)

**gene.xy** list (one entry per affyID)

**params.gene** environment

## Details

These chip-specific data structures are generated from the data files made available by the author of the PDNN paper (see the section source). They are stored as data to save some computation time. The data structures were made using the function `pdnn.params.chiptype`. The data files are included in the the directory ‘data’ of the package.

## Note

To lower the size of the package, the only chip-specific data structures included in the package is the one for HG-U95Av2.

## Source

Li Zhang, Michael F. Miles and Kenneth D. Aldape - A model of molecular interactions on short oligonucleotide arrays, 2003, Nature Biotech., vol. 21, n.7

## Examples

```
## give the path the original energy parameter files included in the package
list.files(system.file("exampleData", package="affypdnn"),
          "^pdnn-energy-parameter_", full.names=TRUE)
```

`matplotlibProbesPDNN`      *Plot the PDNN computed probe intensities*

## Description

Plot the probe intensities as computed by '`pmcorrect.pdnn`' or '`pmcorrect.pdnnpredict`'

## Usage

```
matplotlibProbesPDNN(x, type="l", ...)
```

## Arguments

<code>x</code>	a matrix (and attributes) as returned by <code>pmcorrect.pdnn</code> or <code>pmcorrect.pdnnpredict</code> .
<code>type</code>	type of plot (same as in <code>matplotlib</code> )
<code>...</code>	optional arguments to be passed to <code>matplotlib</code>

## Details

The crosses are the probe intensities which are considered ‘ok’ by the outlier detection part of the algorithm, while the circles are the ones considered ‘outliers’

## Value

Only used for its side-effect.

## See Also

[pmcorrect.pdnn](#) and [pmcorrect.pdnnpredict](#)

## Examples

```
# see 'pmcorrect.pdnn'
```

`params.dilution`      *Parameters for the Dilution dataset*

## Description

PDNN parameters for the Dilution dataset

## Usage

```
data(params.dilution)
```

## Format

The format is: List of 6 \$ lambda :List of 12625 (probesets) ... \$ Bs : num [1:4] 111.9 57.3 120.5 50.1 \$ Ns : num [1:4] 2967 2998 2992 2999 \$ Fs : num [1:4] 0.607 0.662 0.600 0.656 \$ names.abatch: chr [1:12625] "1000\_at" "1001\_at" "1002\_f\_at" ... \$ names.i : int [1:12625] 1 2 3 4 5 6 7 8 9 10 ...

## Details

These data are provided to reduce the time needed to run the vignette.

## Examples

```
data(params.dilution)
```

`pdnn.params.chiptype` *A function to fit PDNN parameters*

## Description

A function to fit PDNN parameters that are chip-type specific

## Usage

```
pdnn.params.chiptype(energy.param.file, probes.file = NULL, probes.pack= NULL,
                      probes.data.frame = NULL,
                      seq.name, x.name, y.name, affyid.name, verbose = TRUE)
```

## Arguments

<code>energy.param.file</code>	Path to the energy data file (see details)
<code>probes.file</code>	Path to the probe files (see details)
<code>probes.pack</code>	Name of the probe pack (see details)
<code>probes.data.frame</code>	A <code>data.frame</code>
<code>seq.name, x.name, y.name, affyid.name</code>	The names of the columns in the <code>data.frame</code> from <code>probes.pack</code> or <code>probes.file</code> for the probe sequences, the X positions, the Y positions and the probe set ID respectively
<code>verbose</code>	verbosity (TRUE or FALSE)

## Details

The parameters `probes.file`, `probes.pack` and `probes.data.frame` are mutually exclusive. The function fits PDNN parameters that are specific to chip-types (hence specific to the probe sequences). It requires data files like the one found on Li Zhang's web page: (<http://odin.mdacc.tmc.edu/~zhangli/PerfectMatch/>). This should be computed once for all for a given chip type. Computed values for the chips are included in the package. This allows 'automagic' use of them when these chip types are used (as done in the function [expressopdnn](#)).

## Value

A list of:

Eg	environment. One entry per dinucleotide.
Wg	numerical vector
En	environment. One entry per dinucleotide.
Wn	numerical vector
params.gene	environment. One entry per gene, each entry is a list of elements Sg, Sn, xy and gene.i

## warning

The X and Y positions in the `data.frame` are expected to be original ones in the Affymetrix files (starting at zero). They are offset by one within this function.

## See Also

[find.params.pdnn](#)

## Examples

```
if (interactive()) {
  energy.file <- system.file("exampleData", "pdnn-energy-parameter_hg-u95av2.txt", package="affypdnn")

  params.chiptype <- pdnn.params.chiptype(energy.file, probes.pack="hgu95av2probe")
}
```

*transform.ProbeSet*      A function to transform a *ProbeSet*

## Description

A function to transform the PMs and MMs in a *ProbeSet*.

## Usage

```
## S3 method for class 'ProbeSet'
transform(`_data`, fun = I, ...)
```

**Arguments**

- \_data object of [ProbeSet-class](#).
- fun a function. The identity function by default.
- ... optional arguments for fun.

**Details**

The function `fun` is applied to the slots `pm` and `mm`. The function `vsnh` in the package `vsn` is a recommended argument for `fun`.

**Value**

An object of class `ProbeSet`.

**Note**

This function should make to the package `affy` for the version 1.4.x.

**Author(s)**

Laurent

**Examples**

```
library(affydata)

data(Dilution)

ppset.name <- sample(featureNames(Dilution), 1)
ppset <- probeset(Dilution, ppset.name)[[1]]
ppset.log <- transform(ppset, fun=log)

par(mfrow=c(1,2))
plot(ppset)
plot(ppset.log)
```

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