

# Package: malani (via r-universe)

September 17, 2024

**Type** Package

**Title** Machine Learning Assisted Network Inference

**Version** 1.0

**Author** Mehrab Ghanat Bari

**Maintainer** Mehrab Ghanat Bari <m.ghanatbari@gmail.com>

**Description** Find dark genes. These genes are often disregarded due to no detected mutation or differential expression, but are important in coordinating the functionality in cancer networks.

**License** GPL-3

**LazyData** TRUE

**Depends** e1071, stats

**RoxygenNote** 5.0.1

**NeedsCompilation** no

**Date/Publication** 2016-09-26 23:44:53

**Repository** <https://mehrabgbari.r-universe.dev>

**RemoteUrl** <https://github.com/cran/malani>

**RemoteRef** HEAD

**RemoteSha** a122c14004b5e6d3fe044adb42a72ece819dbe92

## Contents

dat . . . . .	2
grp . . . . .	2
Gsvmod . . . . .	2
intGenes . . . . .	3
malanidata . . . . .	4
pairmod . . . . .	4
<b>Index</b>	<b>5</b>

`dat` *A matrix of expression values.*

---

**Description**

A numeric matrix 100\*20.

**Usage**

`dat`

**Format**

matrix.

---

`grp` *A vector of class labels for [dat](#).*

---

**Description**

Vector length of 20.

**Usage**

`grp`

**Format**

vector

---

`Gsvmod` *G SVM models.*

---

**Description**

Returns accuracy performance of all genes. G support vector machine (SVM) classifiers trained using G different data matrixes, are used to predict labels in test data. Models are ranked based on prediction performances.

**Usage**

`Gsvmod(dat.train, lab.train, dat.test, lab.test)`

**Arguments**

dat.train	Train data with G features and $(k-1)*S/k$ samples. Parameter k comes from cross-validation scheme and is specified by user (default is 2).
lab.train	Class labels for train data.
dat.test	Test data with G features and S/k samples.
lab.test	Class labels for test data.

**Value**

Accuracy scores for models. Each model represents one gene.

---

intGenes	<i>Select initial gene list from original data matrix.</i>
----------	--

---

**Description**

Train G-1 SVM models in k-fold cross validation scheme to select initial genes list.

**Usage**

```
intGenes(dat, grp, nfolds.out = 2, top.per = 0.05)
```

**Arguments**

dat	Original gene expression data matrix with G rows (number of genes) and S column (number of samples).
grp	Class labels.
nfolds.out	Outer cross validation number (default is 2).
top.per	All genes are ranked based on their models performance and top.per% of them are selected as initial genes.

**Value**

Selected initial genes.

**Examples**

```
data(malanidata)
int <- intGenes(dat,grp)
print(int$top.genes)
```

---

malanidata	<i>Dataset for malani package</i>
------------	-----------------------------------

---

**Description**

A numeric matrix  $G \times S$  contains gene expressions data.  $G$  are the genes (rows) and  $S$  are the samples (columns).

**Usage**

```
malanidata
```

**Format**

A matrix of numeric values, 100 genes, 20 samples and class labels.

**Examples**

```
data(malanidata)
```

---

pairmod	<i>Find best performing pairs</i>
---------	-----------------------------------

---

**Description**

Combine each gene in initial set with all genes in the original set. Top  $n_{pair}$  pairs are selected to construct the  $Q$  matrix.

**Usage**

```
pairmod(X, LX, theta, npair = 10)
```

**Arguments**

$X$	Original gene expression data matrix. With $G$ rows (number of genes) and $S$ column (number of samples).
$LX$	Class labels.
$\theta$	Initial gene set.
$n_{pair}$	Given a gene in initial set, top $n_{pair}$ best performing pairs correspond to that gene are selected (Default is 10).

**Value**

Best  $(n_{pair} \times G / 20)$  performing pairs.

# Index

## \* datasets

dat, 2

grp, 2

malanidata, 4

dat, 2, 2

grp, 2

Gsvmod, 2

intGenes, 3

malanidata, 4

pairmod, 4