

# Package ‘RichR’

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**Type** Package

**Title** Enrichment for Diseases in a Set of Genes

**Version** 1.0.0

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## Description

It provides a list of genes associated to diseases (g2d\$clean and g2d\$complete) based on the following 4 publications (GS2D, Fontaine (2016) <doi:10.18547/gcb.2016.vol2.iss1.e33>, DisGeNET, Pinero (2016) <doi:10.1093/nar/gkw943> Berto2016, Berto (2016) <doi:10.3389/fgene.2016.00031> and PsyGeNET, Sacristan (2015) <doi:10.1093/bioinformatics/btv301>). Those lists were combined and manually curated to have matching disease names. When provided a list of gene names, it calculates the disease enrichment of the gene set. The enrichment is calculated using proportion test and Fisher's exact test. Adjusted fdr p-values are returned alongside with p-values combined using the Fisher's method.

**License** GPL-2

**Imports** plyr, reshape2, magrittr, stats, metap

**Encoding** UTF-8

**LazyData** true

**Depends** R (>= 3.1)

**RoxygenNote** 6.1.1

**NeedsCompilation** no

**Repository** CRAN

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Enrichment

*Enrichment*

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### Description

Given a list of genes associated to diseases and a background list, it calculates the diseases enrichment. It calculates both pvalues from proportion test and Fisher's exact test. Adjusted BH pvalues are returned.

### Usage

```
Enrichment(Background, Genes2Dis = g2d$clean, Genes)
```

### Arguments

Background	The background list of genes. In general is the list of genes expressed.
Genes2Dis	A data.frame with the Gene names and the Diseases. The package has two inbuild lists. <code>g2d_clean</code> and <code>g2d_complete</code> . Both lists contains data from 4 publications (GS2D, DisGeNET, Berto2016 and PsyGeNET), however in the clean similar names were treated as the same disease.
Genes	A list of genes to test for enrichment

### Value

a list containing the enrichment of diseases

### Author(s)

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### Examples

```
data('g2d')

g2d_clean = g2d$clean

# The user can choose a particular disorder, or use the whole disease set to compare to

g2d_ASD = subset(g2d_clean, g2d_clean$Disease %in% c('AUTISM'))
Enrichment(Background = g2d_clean$Gene.symbol,
Genes2Dis = g2d_ASD,
Genes = g2d_ASD$Gene.symbol[1:100])
```

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*g2d**g2d*

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**Description**

*g2d* is a list with two data.frames based on the following 4 publications (GS2D, Fontaine (2016) <doi:10.18547/gcb.2016.vol2.iss1.e33>, DisGeNET, Pinero (2016) <doi:10.1093/nar/gkw943>, Berto2016, Berto (2016) <doi:10.3389/fgene.2016.00031> and PsyGeNET, Sacristan (2015) <doi:10.1093/bioinformatics/btv301>). Those lists were combined and manually curated to have matching disease names. The first list, clean, contains the curated data, the list complete contains complete data. In the former disease names might not match.

**Usage***g2d***Format**

An object of class `list` of length 2.

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