An Introduction to GSEABase

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May 19, 2021

Abstract

The *GSEABase* package implements data structures and methods to represent, manipulate, and analyze gene sets in the context of gene set enrichment analysis. This includes construction of gene sets from reference resources, ID mapping, coloring according to phenotype association, and storing in gene set collections.

Package

GSEABase 1.54.0

Report issues on https://github.com/Bioconductor/GSEABase/issues

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1 GeneSet

A *GeneSet* stores a set of related gene identifiers. Important components of the gene set are a vector of identifiers, general descriptive information about the set, and information about how the gene set was constructed. To construct a gene set, use <u>GeneSet</u>. For example, to create a gene set from the identifiers in a subset of the sample *ExpressionSet* in the *Biobase* package use

```
data(sample.ExpressionSet) # from Biobase
egs <- GeneSet(sample.ExpressionSet[201:250,], setName="Sample")
egs

## setName: Sample
## geneIds: 31440_at, 31441_at, ..., 31489_at (total: 50)
## geneIdType: Annotation (hgu95av2)
## collectionType: ExpressionSet
## details: use 'details(object)'</pre>
```

Each gene set may have a name. The gene set contains 50 identifiers ('genes') from the *ExpressionSet*. These are accessible using geneIds, e.g.,

```
head(geneIds(egs))
## [1] "31440_at" "31441_at" "31442_at" "31443_at" "31444_s_at"
## [6] "31445_at"
```

The gene set records that the identifiers are probeset names from the annotation package hgu95av2.db, and that the source of the gene set was an ExpressionSet. Additional details are available:

```
details(egs)

## setName: Sample
## geneIds: 31440_at, 31441_at, ..., 31489_at (total: 50)

## geneIdType: Annotation (hgu95av2)

## collectionType: ExpressionSet

## setIdentifier: machv2.bioconductor.org:60128:Wed May 19 20:07:23 2021:1

## description: Smoking-Cancer Experiment

## (longDescription available)

## organism: Homo sapiens

## pubMedIds:

## urls: www.lab.not.exist

## contributor: Pierre Fermat

## setVersion: 0.0.1

## creationDate:
```

The set identifier, set version, and creation date provide mechanisms for carefully curating gene sets. Additional information is automatically copied from the expression set used to create egs.

```
## FIXME: GeneSet(AnnotationIdentifier("hgu95av2")) --> non-empty
## FIXME: GeneSet(AnnotationIdentifier("hgu95av2"),
## collectionType=GOCollection()) filters on GOCollection (or KEGG)
```

View additional methods for creating gene sets with:

```
showMethods("GeneSet", inherited=FALSE)

## Function: GeneSet (package GSEABase)

## type="BroadCollection"

## type="ExpressionSet"

## type="GOCollection"

## type="GeneIdentifierType"

## type="character"

## type="missing"
```

These are described on the GeneSet help page.

The identifier type of gene sets created from expression sets is AnnotationIdentifier. Additional predefined identifiers are available:

```
names(slot(getClass("GeneIdentifierType"), "subclasses"))

## [1] "NullIdentifier" "EnzymeIdentifier" "GenenameIdentifier"

## [4] "RefseqIdentifier" "SymbolIdentifier" "UniprotIdentifier"

## [7] "ENSEMBLIdentifier" "AnnotationIdentifier" "EntrezIdentifier"
```

```
## [10] "GOAllFrameIdentifier" "KEGGFrameIdentifier"
```

It is possible to map between identifier types (provided the corresponding map in the annotation package exists):

```
mapIdentifiers(egs, EntrezIdentifier())

## setName: Sample

## geneIds: 6932, 643332, ..., 4287 (total: 31)

## geneIdType: EntrezId (hgu95av2)

## collectionType: ExpressionSet

## details: use 'details(object)'
```

mapIdentifiers consults the gene set to determine that annotation (probeset) identifiers are to be converted to Entrez IDs based on the mapping in the <code>hgu95av2.db</code> package. The function <code>mapIdentifiers</code> can automatically determine many of the common maps; it is also possible to provide as a third argument an environment containing an arbitrary map. Use the <code>verbose</code> argument of <code>mapIdentifiers</code> to be informed when the map between identifier types is not 1:1.

A gene set can be created with different types of identifier, e.g., to create a gene set with Entrez IDs, use

```
library(annotate) # getEG
eids <- unique(getEG(geneIds(egs), "hgu95av2"))
eids <- eids[!is.na(eids)]
GeneSet(EntrezIdentifier(), geneIds=as.character(eids))

## setName: NA
## geneIds: 6932, 643332, ..., 4287 (total: 31)
## geneIdType: EntrezId
## collectionType: Null
## details: use 'details(object)'</pre>
```

The **collectionType** of a gene set provides additional information about a gene set. Available collection types are

```
names(slot(getClass("CollectionType"), "subclasses"))
## [1] "NullCollection"
                                  "ExpressionSetCollection"
   [3] "ComputedCollection"
                                  "CollectionIdType"
## [5] "BroadCollection"
                                  "KEGGCollection"
## [7] "OMIMCollection"
                                  "PMIDCollection"
## [9] "ChrCollection"
                                  "ChrlocCollection"
## [11] "MapCollection"
                                  "PfamCollection"
## [13] "PrositeCollection"
                                  "GOCollection"
## [15] "OBOCollection"
```

Collection types are most important when creating gene sets. For instance, the *GOCollection* class allows creation of gene sets based on gene ontology (GO) terms. The following command creates a gene set from two terms, including all genes with a particular evidence code.

```
geneIdType=EntrezIdentifier("org.Hs.eg.db"),
    setName="Sample GO Collection")

## setName: Sample GO Collection
## geneIds: (total: 0)

## geneIdType: EntrezId (org.Hs.eg.db)

## collectionType: GO

## ids: GO:0005488, GO:0019825 (2 total)

## evidenceCode: IDA

## ontology: CC MF BP

## details: use 'details(object)'
```

This creates a gene set by consulting an object in the *GO.db* package. A gene set created from an expression set, and with collection type *GOCollection* consults the appropriate environment in the annotation package associated with the expression set.

Gene sets encoded in XML following the schema and conventions of the Broad Institute can be read into R as follows:

```
fl <- system.file("extdata", "Broad1.xml", package="GSEABase")
bgs <- GeneSet(BroadCollection(), urls=fl)
bgs

## setName: chr16q24
## geneIds: GALNS, C160RF44, ..., TRAPPC2L (total: 129)
## geneIdType: Symbol
## collectionType: Broad
## bcCategory: c1 (Positional)
## bcSubCategory: NA
## details: use 'details(object)'</pre>
```

The example above uses a local file, but the source for the gene set might also be a web address accessible via the http:// protocol. The file name is added to the url of the gene set. The set name and category of the Broad collection indicate that the gene set contains gene symbols from band 24 of the q arm of chromosome 16. The probe sets in chip hgu95av2 corresponding to these symbols can be determined by mapping identifiers

```
bgs1 <- mapIdentifiers(bgs, AnnotationIdentifier("hgu95av2"))
bgs1

## setName: chr16q24

## geneIds: 32100_r_at, 32101_at, ..., 35807_at (total: 36)

## geneIdType: Annotation (hgu95av2)

## collectionType: Broad

## bcCategory: c1 (Positional)

## bcSubCategory: NA

## details: use 'details(object)'</pre>
```

Subsetting creates sets with just the symbols identified. Subsetting can use indices or symbol names.

```
bgs[1:5]
## setName: chr16q24
```

```
## geneIds: GALNS, C160RF44, ..., L0C646365 (total: 5)
## geneIdType: Symbol
## collectionType: Broad
## bcCategory: c1 (Positional)
## bcSubCategory: NA
## details: use 'details(object)'

bgs[c("GALNS", "L0C646365")]

## setName: chr16q24
## geneIds: GALNS, L0C646365 (total: 2)
## geneIdType: Symbol
## collectionType: Broad
## bcCategory: c1 (Positional)
## bcSubCategory: NA
## details: use 'details(object)'
```

Logical operations provide a convenient way to identify genes with particular properties. For instance, the intersection

```
egs & bgs1

## setName: (Sample & chr16q24)

## geneIds: (total: 0)

## geneIdType: Annotation (hgu95av2)

## collectionType: Computed

## details: use 'details(object)'
```

is empty (note that the identifiers in the two sets were of the same type), indicating that none of the identifiers in egs are on 16q24. Additional operations on sets include union (performed with |) and difference (setdiff).

Methods exist to directly subset expression sets using gene sets

```
sample.ExpressionSet[bgs,]

## ExpressionSet (storageMode: lockedEnvironment)

## assayData: 1 features, 26 samples

## element names: exprs, se.exprs

## protocolData: none

## phenoData

## sampleNames: A B ... Z (26 total)

## varLabels: sex type score

## varMetadata: labelDescription

## experimentData: use 'experimentData(object)'

## Annotation: hgu95av2
```

Remember that sample. ExpressionSet contains just 500 probe sets, so the small size of the subset is not surprising. Note also that subsetting required mapping of symbol identifiers in bgs to *AnnotationIdentifiers*; this map used the annotation information in the expression set, and is not necessarily 1:1.

2 GeneColorSet

A *GeneColorSet* is a gene set with "coloring" to indicate how features of genes and phenotypes are associated. The following sample data describes how changes in expression levels of several genes (with Entrez and Symbol names) influence cisplatin resistance phenotype.

```
tbl
##
     Entrez.ID Gene.Symbol Expression.level Phenotype.response
## 1
          1244
                      ABCC2
                                    Increase
                                                       Resistant
           538
## 2
                      ATP7A
                                    Increase
                                                       Resistant
## 3
           540
                      ATP7B
                                    Increase
                                                       Resistant
## 4
          9961
                        MVP
                                    Increase
                                                       Resistant
## 5
          7507
                        XPA
                                    Increase
                                                       Resistant
## 6
          2067
                      ERCC1
                                    Increase
                                                       Resistant
## 7
           672
                      BRCA1
                                    Increase
                                                       Resistant
## 8
          3725
                        JUN
                                    Increase
                                                       Resistant
## 9
          2730
                       GCLM
                                    Increase
                                                       Resistant
```

Note that three different aspects of data influence coloring: the phenotype under consideration (cisplatin resistance), whether expression responses refer to increasing or decreasing levels of gene expression, and whether the phenotypic response represents greater resistance or sensitivity to cisplatin. Here is the resulting gene color set:

```
gcs <- GeneColorSet(EntrezIdentifier(),</pre>
                    setName="A color set",
                    geneIds=as.character(tbl$Entrez.ID),
                    phenotype="Cisplatin resistance",
                    geneColor=tbl$Expression.level,
                    phenotypeColor=tbl$Phenotype.response)
qcs
## setName: A color set
## geneIds: 1244, 538, ..., 2730 (total: 9)
## geneIdType: EntrezId
## collectionType: Null
## phenotype: Cisplatin resistance
## geneColor: Increase, Increase, ..., Increase
     levels: Increase
## phenotypeColor: Resistant, Resistant, ..., Resistant
     levels: Resistant
## details: use 'details(object)'
```

Gene color sets can be used in the same way as gene sets, e.g., for subsetting expression sets (provided the map between identifiers is 1:1, so that coloring corresponding to identifiers can be determined). The coloring method allows access to the coloring information with a data frame interface; phenotype, geneColor and phenotypeColor provide additional accessors.

3 GeneSetCollection

A GeneSetCollection is a collection of gene sets. Sets in the collection must have distinct setNames, but can be a mix of GeneSet and GeneColorSet. Two convenient ways to create a gene set collection are by specifying a source of identifiers (e.g., an ExpressionSet or AnnotationIdentifier) and how the identifiers are to be induced into sets (e.g., by consulting the GO or KEGG ontologies):

```
gsc <- GeneSetCollection(sample.ExpressionSet[201:250,], setType=GOCollection())</pre>
gsc
## GeneSetCollection
     names: G0:0000122, G0:0000186, ..., G0:1990837 (340 total)
    unique identifiers: 31471_at, 31468_f_at, ..., 31477_at (32 total)
##
    types in collection:
       geneIdType: AnnotationIdentifier (1 total)
       collectionType: GOCollection (1 total)
gsc[["G0:0005737"]]
## setName: G0:0005737
## geneIds: 31450_s_at, 31451_at, ..., 31489_at (total: 11)
## geneIdType: Annotation (hgu95av2)
## collectionType: G0
    ids: G0:0005737 (1 total)
    evidenceCode: EXP IDA IPI IMP IGI IEP HTP HDA HMP HGI HEP ISS ISO ISA ISM IGC IBA IBD IKR IRD RCA TAS N
    ontology: CC MF BP
## details: use 'details(object)'
```

In this example, the annotation identifiers of the sample expression set are organized into gene sets based on their presence in GO pathways. Providing arguments such as evidenceCode to GOCollection act to select just those pathways satisfying the GO collection constraint:

Sets in the collection are named after the GO terms, and can be accessed by numeric index or name.

A file or url containing several gene sets defined by Broad XML can be used to create a gene set collection, e.g.,

```
## FIXME: BroadCollection default to paste("c", 1:4, sep="")
## FIXME: GeneSetCollection(BroadCollection(), urls=fl); filters on bcCategory
fl <- system.file("extdata", "Broad.xml", package="GSEABase")
gss <- getBroadSets(fl)
gss</pre>
```

```
## GeneSetCollection
## names: chr5q23, chr16q24 (2 total)
## unique identifiers: ZNF474, CCDC100, ..., TRAPPC2L (215 total)
## types in collection:
## geneIdType: SymbolIdentifier (1 total)
## collectionType: BroadCollection (1 total)
names(gss)
## [1] "chr5q23" "chr16q24"
```

Identifiers within a gene set collection can be mapped to a common type (provided maps are available) with, for example,

```
gsc <- mapIdentifiers(gsc, EntrezIdentifier())</pre>
qsc
## GeneSetCollection
    names: G0:0000122, G0:0000186, ..., G0:1990837 (340 total)
## unique identifiers: 641339, 2911, ..., 7033 (27 total)
    types in collection:
       geneIdType: EntrezIdentifier (1 total)
##
       collectionType: GOCollection (1 total)
gsc[["G0:0005737"]]
## setName: G0:0005737
## geneIds: 6014, 392, ..., 4287 (total: 8)
## geneIdType: EntrezId (hgu95av2)
## collectionType: G0
   ids: G0:0005737 (1 total)
## evidenceCode: EXP IDA IPI IMP IGI IEP HTP HDA HMP HGI HEP ISS ISO ISA ISM IGC IBA IBD IKR IRD RCA TAS N.
## ontology: CC MF BP
## details: use 'details(object)'
```

A convenient way to visualize a GeneSetCollection is with the ReportingTools package.

```
## 'interesting' gene sets
idx <- sapply(gsc, function(x) length(geneIds(x))) > 2
library(ReportingTools)
## Loading required package: knitr
## Registered S3 method overwritten by 'GGally':
     method from
##
     +.gg
           ggplot2
##
gscReport <- HTMLReport(</pre>
    shortName="gsc_example",
    title="GSEABase Vignette GeneSetCollection",
    basePath=tempdir())
publish(gsc[idx], gscReport, annotation.db="org.Hs.eg")
url <- finish(gscReport)</pre>
```

GSEABase

The report can be viewed with

browseURL(url)

This concludes a brief tour of gene sets and gene set collections available in the *GSEABase* package.