

AnnotationDbi: Introduction To Bioconductor Annotation Packages

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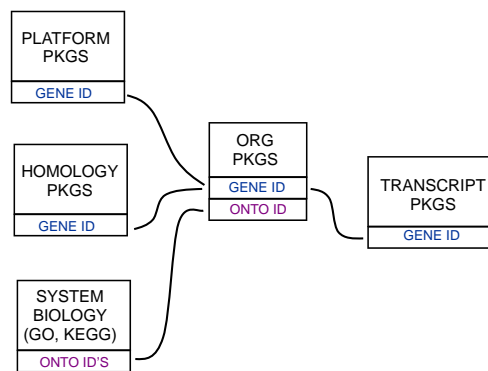


Figure 1: Annotation Packages: the big picture

Bioconductor provides extensive annotation resources. These can be *gene centric*, or *genome centric*. Annotations can be provided in packages curated by *Bioconductor*, or obtained from web-based resources. This vignette is primarily concerned with describing the annotation resources that are available as packages. More advanced users who wish to learn about how to make new annotation packages should see the vignette titled "Creating select Interfaces for custom Annotation resources" from the *AnnotationForge* package.

Gene centric *AnnotationDbi* packages include:

- Organism level: e.g. *org.Mm.eg.db*.
- Platform level: e.g. *hgu133plus2.db*, *hgu133plus2.probes*, *hgu133plus2.cdf*.
- Homology level: e.g. *hom.Dm.inp.db*.
- System-biology level: *GO.db*

Genome centric *GenomicFeatures* packages include

- Transcriptome level: e.g. *TxDb.Hsapiens.UCSC.hg19.knownGene*, *EnsDb.Hsapiens.v75*.
- Generic genome features: Can generate via *GenomicFeatures*

One web-based resource accesses [biomart](#), via the *biomaRt* package:

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- Query web-based 'biomart' resource for genes, sequence, SNPs, and etc.

The most popular annotation packages have been modified so that they can make use of a new set of methods to more easily access their contents. These four methods are named: `columns`, `keytypes`, `keys` and `select`. And they are described in this vignette. They can currently be used with all `chip`, `organism`, and `TxDb` packages along with the popular `GO.db` package.

For the older less popular packages, there are still convenient ways to retrieve the data. The *How to use bimap from the ".db" annotation packages* vignette in the `AnnotationDbi` package is a key reference for learnign about how to use bimap objects.

Finally, all of the '.db' (and most other *Bioconductor* annotation packages) are updated every 6 months corresponding to each release of *Bioconductor*. Exceptions are made for packages where the actual resources that the packages are based on have not themselves been updated.

1 AnnotationDb objects and the select method

As previously mentioned, a new set of methods have been added that allow a simpler way of extracting identifier based annotations. All the annotation packages that support these new methods expose an object named exactly the same as the package itself. These objects are collectively called *AnntoationDb* objects for the class that they all inherit from. The more specific classes (the ones that you will actually see in the wild) have names like `OrgDb`, `ChipDb` or `TxDb` objects. These names correspond to the kind of package (and underlying schema) being represented. The methods that can be applied to all of these objects are `columns`, `keys`, `keytypes` and `select`.

In addition, another accessor has recently been added which allows extraction of one column at a time. the `mapIds` method allows users to extract data into either a named character vector, a list or even a `SimpleCharacterList`. This method should work with all the different kinds of *AnntoationDb* objects described below.

2 ChipDb objects and the select method

An extremely common kind of Annotation package is the so called platform based or chip based package type. This package is intended to make the manufacturer labels for a series of probes or probesets to a wide range of gene-based features. A package of this kind will load an `ChipDb` object. Below is a set of examples to show how you might use the standard 4 methods to interact with an object of this type.

First we need to load the package:

```
suppressPackageStartupMessages({  
  library(hgu95av2.db)
```

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```
} )
```

If we list the contents of this package, we can see that one of the many things loaded is an object named after the package "hgu95av2.db":

```
ls("package:hgu95av2.db")  
  
## [1] "hgu95av2"                "hgu95av2.db"                "hgu95av2ACCCNUM"  
## [4] "hgu95av2ALIAS2PROBE"     "hgu95av2CHR"                "hgu95av2CHRENGTHS"  
## [7] "hgu95av2CHRL0C"         "hgu95av2CHRL0CEND"          "hgu95av2ENSEMBL "  
## [10] "hgu95av2ENSEMBL2PROBE"  "hgu95av2ENTREZID"           "hgu95av2ENZYME"  
## [13] "hgu95av2ENZYME2PROBE"   "hgu95av2GENENAME"           "hgu95av2G0"  
## [16] "hgu95av2G02ALLPROBES"   "hgu95av2G02PROBE"           "hgu95av2MAP"  
## [19] "hgu95av2MAPCOUNTS"     "hgu95av20MIM"                "hgu95av20ORGANISM"  
## [22] "hgu95av20RGPKG"         "hgu95av2PATH"                "hgu95av2PATH2PROBE"  
## [25] "hgu95av2PFAM"           "hgu95av2PMID"                "hgu95av2PMID2PROBE"  
## [28] "hgu95av2PROSITE"        "hgu95av2REFSEQ"              "hgu95av2SYMBOL "  
## [31] "hgu95av2UNIGENE"        "hgu95av2UNIPROT"             "hgu95av2_dbInfo"  
## [34] "hgu95av2_dbconn"        "hgu95av2_dbfile"             "hgu95av2_dbschema"
```

We can look at this object to learn more about it:

```
hgu95av2.db  
  
## ChipDb object:  
## | DBSCHEMAVERSION: 2.1  
## | Db type: ChipDb  
## | Supporting package: AnnotationDbi  
## | DBSCHEMA: HUMANCHIP_DB  
## | ORGANISM: Homo sapiens  
## | SPECIES: Human  
## | MANUFACTURER: Affymetrix  
## | CHIPNAME: Human Genome U95 Set  
## | MANUFACTURERURL: http://www.affymetrix.com/support/technical/byproduct.affx?product=hgu95  
## | EGSOURCEDATE: 2015-Sep27  
## | EGSOURCENAME: Entrez Gene  
## | EGSOURCEURL: ftp://ftp.ncbi.nlm.nih.gov/gene/DATA  
## | CENTRALID: ENTREZID  
## | TAXID: 9606  
## | GOSOURCENAME: Gene Ontology  
## | GOSOURCEURL: ftp://ftp.geneontology.org/pub/go/godatabase/archive/latest-lite/  
## | GOSOURCEDATE: 20150919  
## | GOEGSOURCEDATE: 2015-Sep27  
## | GOEGSOURCENAME: Entrez Gene  
## | GOEGSOURCEURL: ftp://ftp.ncbi.nlm.nih.gov/gene/DATA  
## | KEGGSOURCENAME: KEGG GENOME
```

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```
## | KEGGSOURCEURL: ftp://ftp.genome.jp/pub/kegg/genomes
## | KEGGSOURCEDATE: 2011-Mar15
## | GPSOURCENAME: UCSC Genome Bioinformatics (Homo sapiens)
## | GPSOURCEURL: ftp://hgdownload.cse.ucsc.edu/goldenPath/hg19
## | GPSOURCEDATE: 2010-Mar22
## | ENSOURCEDATE: 2015-Jul16
## | ENSOURCENAME: Ensembl
## | ENSOURCEURL: ftp://ftp.ensembl.org/pub/current_fasta
## | UPSOURCENAME: Uniprot
## | UPSOURCEURL: http://www.uniprot.org/
## | UPSOURCEDATE: Thu Oct 1 23:31:58 2015

##
## Please see: help('select') for usage information
```

If we want to know what kinds of data are retrievable via `select`, then we should use the `columns` method like this:

```
columns(hgu95av2.db)

## [1] "ACCNUM"      "ALIAS"      "ENSEMBL"    "ENSEMBLPROT"
## [5] "ENSEMBLTRANS" "ENTREZID"   "ENZYME"     "EVIDENCE"
## [9] "EVIDENCEALL" "GENENAME"   "GO"         "GOALL"
## [13] "IPI"        "MAP"        "OMIM"       "ONTOLOGY"
## [17] "ONTOLOGYALL" "PATH"      "PFAM"       "PMID"
## [21] "PROBEID"    "PROSITE"   "REFSEQ"     "SYMBOL"
## [25] "UCSCKG"    "UNIGENE"   "UNIPROT"
```

If we are further curious to know more about those values for columns, we can consult the help pages. Asking about any of these values will pull up a manual page describing the different fields and what they mean.

```
help("SYMBOL")
```

If we are curious about what kinds of fields we could potentially use as keys to query the database, we can use the `keytypes` method. In a perfect world, this method will return values very similar to what was returned by `columns`, but in reality, some kinds of values make poor keys and so this list is often shorter.

```
keytypes(hgu95av2.db)

## [1] "ACCNUM"      "ALIAS"      "ENSEMBL"    "ENSEMBLPROT"
## [5] "ENSEMBLTRANS" "ENTREZID"   "ENZYME"     "EVIDENCE"
## [9] "EVIDENCEALL" "GENENAME"   "GO"         "GOALL"
## [13] "IPI"        "MAP"        "OMIM"       "ONTOLOGY"
## [17] "ONTOLOGYALL" "PATH"      "PFAM"       "PMID"
## [21] "PROBEID"    "PROSITE"   "REFSEQ"     "SYMBOL"
```

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```
## [25] "UCSCKG" "UNIGENE" "UNIPROT"
```

If we want to extract some sample keys of a particular type, we can use the `keys` method.

```
head(keys(hgu95av2.db, keytype="SYMBOL"))
## [1] "A1BG" "A2M" "A2MP1" "NAT1" "NAT2" "NATP"
```

And finally, if we have some keys, we can use `select` to extract them. By simply using appropriate argument values with `select` we can specify what keys we want to look up values for (keys), what we want returned back (columns) and the type of keys that we are passing in (keytype)

```
#1st get some example keys
k <- head(keys(hgu95av2.db, keytype="PROBEID"))
# then call select
select(hgu95av2.db, keys=k, columns=c("SYMBOL", "GENENAME"), keytype="PROBEID")

## 'select()' returned 1:1 mapping between keys and columns

##   PROBEID SYMBOL
## 1  1000_at  MAPK3
## 2  1001_at  TIE1
## 3 1002_f_at CYP2C19
## 4 1003_s_at CXCR5
## 5  1004_at  CXCR5
## 6  1005_at  DUSP1
##
##                                GENENAME
## 1                mitogen-activated protein kinase 3
## 2 tyrosine kinase with immunoglobulin like and EGF like domains 1
## 3                cytochrome P450 family 2 subfamily C member 19
## 4                                C-X-C motif chemokine receptor 5
## 5                                C-X-C motif chemokine receptor 5
## 6                dual specificity phosphatase 1
```

And as you can see, when you call the code above, `select` will try to return a data.frame with all the things you asked for matched up to each other.

Finally if you wanted to extract only one column of data you could instead use the `mapIds` method like this:

```
#1st get some example keys
k <- head(keys(hgu95av2.db, keytype="PROBEID"))
# then call mapIds
mapIds(hgu95av2.db, keys=k, column=c("GENENAME"), keytype="PROBEID")

## 'select()' returned 1:1 mapping between keys and columns
```

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```
##                                     1000_at
##                                     "mitogen-activated protein kinase 3"
##                                     1001_at
## "tyrosine kinase with immunoglobulin like and EGF like domains 1"
##                                     1002_f_at
## "cytochrome P450 family 2 subfamily C member 19"
##                                     1003_s_at
## "C-X-C motif chemokine receptor 5"
##                                     1004_at
## "C-X-C motif chemokine receptor 5"
##                                     1005_at
## "dual specificity phosphatase 1"
```

3 OrgDb objects and the select method

An organism level package (an 'org' package) uses a central gene identifier (e.g. Entrez Gene id) and contains mappings between this identifier and other kinds of identifiers (e.g. GenBank or Uniprot accession number, RefSeq id, etc.). The name of an org package is always of the form *org.<Ab>.<id>.db* (e.g. *org.Sc.sgd.db*) where *<Ab>* is a 2-letter abbreviation of the organism (e.g. *Sc* for *Saccharomyces cerevisiae*) and *<id>* is an abbreviation (in lower-case) describing the type of central identifier (e.g. *sgd* for gene identifiers assigned by the Saccharomyces Genome Database, or *eg* for Entrez Gene ids).

Just as the chip packages load a *ChipDb* object, the org packages will load a *OrgDb* object. The following exercise should acquaint you with the use of these methods in the context of an organism package.

Exercise 1

Display the *OrgDb* object for the *org.Hs.eg.db* package.

Use the `columns` method to discover which sorts of annotations can be extracted from it. Is this the same as the result from the `keytypes` method? Use the `keytypes` method to find out.

Finally, use the `keys` method to extract UNIPROT identifiers and then pass those keys in to the `select` method in such a way that you extract the gene symbol and KEGG pathway information for each. Use the help system as needed to learn which values to pass in to `columns` in order to achieve this.

Solution:

```
library(org.Hs.eg.db)
columns(org.Hs.eg.db)
```

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```
## [1] "ACCNUM"      "ALIAS"        "ENSEMBL"     "ENSEMBLPROT"  
## [5] "ENSEMBLTRANS" "ENTREZID"     "ENZYME"      "EVIDENCE"  
## [9] "EVIDENCEALL"  "GENENAME"    "GO"          "GOALL"  
## [13] "IPI"          "MAP"         "OMIM"        "ONTOLOGY"  
## [17] "ONTOLOGYALL" "PATH"        "PFAM"        "PMID"  
## [21] "PROSITE"     "REFSEQ"      "SYMBOL"      "UCSCKG"  
## [25] "UNIGENE"     "UNIPROT"
```

```
help("SYMBOL") ## for explanation of these columns and keytypes values
```

```
keytypes(org.Hs.eg.db)
```

```
## [1] "ACCNUM"      "ALIAS"        "ENSEMBL"     "ENSEMBLPROT"  
## [5] "ENSEMBLTRANS" "ENTREZID"     "ENZYME"      "EVIDENCE"  
## [9] "EVIDENCEALL"  "GENENAME"    "GO"          "GOALL"  
## [13] "IPI"          "MAP"         "OMIM"        "ONTOLOGY"  
## [17] "ONTOLOGYALL" "PATH"        "PFAM"        "PMID"  
## [21] "PROSITE"     "REFSEQ"      "SYMBOL"      "UCSCKG"  
## [25] "UNIGENE"     "UNIPROT"
```

```
uniKeys <- head(keys(org.Hs.eg.db, keytype="UNIPROT"))
```

```
cols <- c("SYMBOL", "PATH")
```

```
select(org.Hs.eg.db, keys=uniKeys, columns=cols, keytype="UNIPROT")
```

```
## 'select()' returned 1:many mapping between keys and columns
```

```
##   UNIPROT SYMBOL  PATH  
## 1  P04217  A1BG  <NA>  
## 2  V9HWD8  A1BG  <NA>  
## 3  P01023  A2M  04610  
## 4  P18440  NAT1  00232  
## 5  P18440  NAT1  00983  
## 6  P18440  NAT1  01100  
## 7  Q400J6  NAT1  00232  
## 8  Q400J6  NAT1  00983  
## 9  Q400J6  NAT1  01100  
## 10 F5H5R8  NAT1  00232  
## 11 F5H5R8  NAT1  00983  
## 12 F5H5R8  NAT1  01100
```

So how could you use `select` to annotate your results? This next exercise should help you to understand how that should generally work.

Exercise 2

Please run the following code snippet (which will load a fake data result that I have provided for the purposes of illustration):

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```
load(system.file("extdata", "resultTable.Rda", package="AnnotationDbi"))
head(resultTable)

##           logConc      logFC LR.statistic      PValue      FDR
## 100418920 -9.639471 -4.679498    378.0732 3.269307e-84 2.613484e-80
## 100419779 -10.638865 -4.264830    291.1028 2.859424e-65 1.142912e-61
## 100271867 -11.448981 -4.009603    222.3653 2.757135e-50 7.346846e-47
## 100287169 -11.026699 -3.486593    206.7771 6.934967e-47 1.385953e-43
## 100287735 -11.036862  3.064980    204.1235 2.630432e-46 4.205535e-43
## 100421986 -12.276297 -4.695736    190.5368 2.427556e-43 3.234314e-40
```

The rownames of this table happen to provide entrez gene identifiers for each row (for human). Find the gene symbol and gene name for each of the rows in resultTable and then use the merge method to attach those annotations to it.

Solution:

```
annots <- select(org.Hs.eg.db, keys=rownames(resultTable),
                 columns=c("SYMBOL", "GENENAME"), keytype="ENTREZID")

## 'select()' returned 1:1 mapping between keys and columns

resultTable <- merge(resultTable, annots, by.x=0, by.y="ENTREZID")
head(resultTable)

##   Row.names  logConc      logFC LR.statistic      PValue      FDR
## 1 100127888 -10.57050  2.758937    182.8937 1.131473e-41 1.130624e-38
## 2 100131223 -12.37808 -4.654318    179.2331 7.126423e-41 6.329847e-38
## 3 100271381 -12.06340  3.511937    188.4824 6.817155e-43 7.785191e-40
## 4 100271867 -11.44898 -4.009603    222.3653 2.757135e-50 7.346846e-47
## 5 100287169 -11.02670 -3.486593    206.7771 6.934967e-47 1.385953e-43
## 6 100287735 -11.03686  3.064980    204.1235 2.630432e-46 4.205535e-43
##           SYMBOL                                     GENENAME
## 1  SLC04A1-AS1                                       SLC04A1 antisense RNA 1
## 2  LOC100131223 ADP ribosylation factor like GTPase 8B pseudogene
## 3   RPS28P8                                           ribosomal protein S28 pseudogene 8
## 4   MPVQTL1                                           Mean platelet volume QTL1
## 5      <NA>                                           <NA>
## 6   TTTY13B                                           testis-specific transcript, Y-linked 13B
```


4 Using select with GO.db

When you load the GO.db package, a *GODb* object is also loaded. This allows you to use the `columns`, `keys`, `keytypes` and `select` methods on the contents of the GO ontology. So if for example, you had a few GO IDs and wanted to know more about it, you could do it like this:

```
library(GO.db)
GOIDs <- c("GO:0042254", "GO:0044183")
select(GO.db, keys=GOIDs, columns="DEFINITION", keytype="GOID")

## 'select()' returned 1:1 mapping between keys and columns
##          GOID
## 1 GO:0042254
## 2 GO:0044183
##
## 1          A cellular process that results in the biosynthesis of constituent macromolecules, assembl
## 2 Interacting selectively and non-covalently with any protein or protein complex (a complex of t
```

5 Using select with TxDb packages

A *TxDb* package (a 'TxDb' package) connects a set of genomic coordinates to various transcript oriented features. The package can also contain Identifiers to features such as genes and transcripts, and the internal schema describes the relationships between these different elements. All TxDb containing packages follow a specific naming scheme that tells where the data came from as well as which build of the genome it comes from.

Exercise 3

Display the TxDb object for the [TxDb.Hsapiens.UCSC.hg19.knownGene](#) package.

As before, use the `columns` and `keytypes` methods to discover which sorts of annotations can be extracted from it.

Use the `keys` method to extract just a few gene identifiers and then pass those keys in to the `select` method in such a way that you extract the transcript ids and transcript starts for each.

Solution:

```
library(TxDb.Hsapiens.UCSC.hg19.knownGene)

## Loading required package: GenomicFeatures
## Loading required package: GenomeInfoDb
```

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```
## Loading required package: GenomicRanges

txdb <- TxDb.Hsapiens.UCSC.hg19.knownGene
txdb

## TxDb object:
## # Db type: TxDb
## # Supporting package: GenomicFeatures
## # Data source: UCSC
## # Genome: hg19
## # Organism: Homo sapiens
## # Taxonomy ID: 9606
## # UCSC Table: knownGene
## # Resource URL: http://genome.ucsc.edu/
## # Type of Gene ID: Entrez Gene ID
## # Full dataset: yes
## # miRBase build ID: GRCh37
## # transcript_nrow: 82960
## # exon_nrow: 289969
## # cds_nrow: 237533
## # Db created by: GenomicFeatures package from Bioconductor
## # Creation time: 2015-10-07 18:11:28 +0000 (Wed, 07 Oct 2015)
## # GenomicFeatures version at creation time: 1.21.30
## # RSQLite version at creation time: 1.0.0
## # DBSCHEMAVERSION: 1.1

columns(txdb)

## [1] "CDSCHROM" "CSEND" "CDSID" "CDSNAME" "CDSSTART"
## [6] "CDSSTRAND" "EXONCHROM" "EXONEND" "EXONID" "EXONNAME"
## [11] "EXONRANK" "EXONSTART" "EXONSTRAND" "GENEID" "TXCHROM"
## [16] "TXEND" "TXID" "TXNAME" "TXSTART" "TXSTRAND"
## [21] "TXTYPE"

keytypes(txdb)

## [1] "CDSID" "CDSNAME" "EXONID" "EXONNAME" "GENEID" "TXID"
## [7] "TXNAME"

keys <- head(keys(txdb, keytype="GENEID"))
cols <- c("TXID", "TXSTART")
select(txdb, keys=keys, columns=cols, keytype="GENEID")

## 'select()' returned 1:many mapping between keys and columns

## GENEID TXID TXSTART
## 1 1 70455 58858172
## 2 1 70456 58859832
## 3 10 31944 18248755
```

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```
## 4      100 72132 43248163
## 5     1000 65378 25530930
## 6     1000 65379 25530930
## 7    10000 7895 243651535
## 8    10000 7896 243663021
## 9    10000 7897 243663021
## 10 100008586 75890 49217763
```

As is widely known, in addition to providing access via the `select` method, `TxDb` objects also provide access via the more familiar `transcripts`, `exons`, `cds`, `transcriptsBy`, `exonsBy` and `cdsBy` methods. For those who do not yet know about these other methods, more can be learned by seeing the vignette called: *Making and Utilizing TxDb Objects* in the `GenomicFeatures` package.

6 Using select with EnsDb packages

Similar to the `TxDb` objects/packages discussed in the previous section, `EnsDb` objects/packages provide genomic coordinates of gene models along with additional annotations (e.g. gene names, biotypes etc) but are tailored to annotations provided by Ensembl. The central methods `columns`, `keys`, `keytypes` and `select` are all implemented for `EnsDb` objects. In addition, these methods allow also the use of the `EnsDb` specific filtering framework to retrieve only selected information from the database (see vignette of the `ensemldb` package for more information).

In the example below we first evaluate which columns, keys and keytypes are available for `EnsDb` objects and fetch then the transcript ids, genomic start coordinate and transcript biotype for some genes.

```
library(EnsDb.Hsapiens.v75)

## Loading required package: ensemblDb

## Loading required package: AnnotationFilter

##
## Attaching package: 'ensemldb'

## The following object is masked from 'package:stats':
##
##   filter

edb <- EnsDb.Hsapiens.v75
edb

## EnsDb for Ensembl:
## |Backend: SQLite
```

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```
## |Db type: EnsDb
## |Type of Gene ID: Ensembl Gene ID
## |Supporting package: ensemblldb
## |Db created by: ensemblldb package from Bioconductor
## |script_version: 0.3.0
## |Creation time: Thu May 18 09:15:45 2017
## |ensembl_version: 75
## |ensembl_host: localhost
## |Organism: homo_sapiens
## |taxonomy_id: 9606
## |genome_build: GRCh37
## |DBSCHEMAVERSION: 2.0
## | No. of genes: 64102.
## | No. of transcripts: 215647.
## |Protein data available.

## List all columns
columns(edb)

## [1] "ENTREZID"           "EXONID"           "EXONIDX"
## [4] "EXONSEQEND"        "EXONSEQSTART"    "GENEBIOTYPE"
## [7] "GENEID"            "GENENAME"         "GENESEQEND"
## [10] "GENESEQSTART"      "INTERPROACCESSION" "ISCIRCULAR"
## [13] "PROTDOMEND"        "PROTDOMSTART"    "PROTEINDOMAINID"
## [16] "PROTEINDOMAINSOURCE" "PROTEINID"        "PROTEINSEQUENCE"
## [19] "SEQCOORDSYSTEM"    "SEQLENGTH"        "SEQNAME"
## [22] "SEQSTRAND"         "SYMBOL"           "TXBIOTYPE"
## [25] "TXCDSSEQEND"       "TXCDSSEQSTART"   "TXID"
## [28] "TXNAME"            "TXSEQEND"         "TXSEQSTART"
## [31] "UNIPROTDB"         "UNIPROTID"        "UNIPROTMAPPINGTYPE"

## List all keytypes
keytypes(edb)

## [1] "ENTREZID"           "EXONID"           "GENEBIOTYPE"
## [4] "GENEID"            "GENENAME"         "PROTDOMID"
## [7] "PROTEINDOMAINID"   "PROTEINDOMAINSOURCE" "PROTEINID"
## [10] "SEQNAME"           "SEQSTRAND"        "SYMBOL"
## [13] "TXBIOTYPE"         "TXID"             "TXNAME"
## [16] "UNIPROTID"

## Get the first
keys <- head(keys(edb, keytype="GENEID"))

## Get the data
select(edb, keys=keys, columns=c("TXID", "TXSEQSTART", "TXBIOTYPE"),
       keytype="GENEID")
```

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##	GENEID	TXID	TXSEQSTART	TXBIOTYPE
## 1	ENSG00000000003	ENST00000373020	99883667	protein_coding
## 2	ENSG00000000003	ENST00000496771	99887538	processed_transcript
## 3	ENSG00000000003	ENST00000494424	99888439	processed_transcript
## 4	ENSG00000000005	ENST00000373031	99839799	protein_coding
## 5	ENSG00000000005	ENST00000485971	99848621	processed_transcript
## 6	ENSG000000000419	ENST00000371588	49551404	protein_coding
## 7	ENSG000000000419	ENST00000466152	49551404	processed_transcript
## 8	ENSG000000000419	ENST00000371582	49551404	protein_coding
## 9	ENSG000000000419	ENST00000494752	49551433	processed_transcript
## 10	ENSG000000000419	ENST00000371584	49551482	protein_coding
## 11	ENSG000000000419	ENST00000413082	49552685	protein_coding
## 12	ENSG000000000419	ENST00000371583	49551490	protein_coding
## 13	ENSG000000000457	ENST00000367771	169818772	protein_coding
## 14	ENSG000000000457	ENST00000367770	169822215	protein_coding
## 15	ENSG000000000457	ENST00000423670	169823652	protein_coding
## 16	ENSG000000000457	ENST00000470238	169828260	processed_transcript
## 17	ENSG000000000457	ENST00000367772	169821804	protein_coding
## 18	ENSG000000000460	ENST00000498289	169631245	processed_transcript
## 19	ENSG000000000460	ENST00000472795	169763871	processed_transcript
## 20	ENSG000000000460	ENST00000496973	169764184	processed_transcript
## 21	ENSG000000000460	ENST00000481744	169764184	processed_transcript
## 22	ENSG000000000460	ENST00000359326	169764190	protein_coding
## 23	ENSG000000000460	ENST00000466580	169764220	processed_transcript
## 24	ENSG000000000460	ENST00000459772	169764220	processed_transcript
## 25	ENSG000000000460	ENST00000286031	169764550	protein_coding
## 26	ENSG000000000460	ENST00000456684	169764220	protein_coding
## 27	ENSG000000000460	ENST00000413811	169764181	protein_coding
## 28	ENSG000000000938	ENST00000374005	27938575	protein_coding
## 29	ENSG000000000938	ENST00000374004	27938811	protein_coding
## 30	ENSG000000000938	ENST00000374003	27939180	protein_coding
## 31	ENSG000000000938	ENST00000457296	27942025	protein_coding
## 32	ENSG000000000938	ENST00000475472	27949895	processed_transcript
## 33	ENSG000000000938	ENST00000468038	27949918	processed_transcript
## 34	ENSG000000000938	ENST00000545953	27938803	protein_coding
## 35	ENSG000000000938	ENST00000399173	27938803	protein_coding

We can modify the queries above to retrieve only genes encoded on chromosome Y. To this end we use filter objects available for *EnsDb* objects and its methods.

```
## Retrieve all gene IDs of all lincRNAs encoded on chromosome Y
linkY <- keys(edb,
              filter=list(GeneBiotypeFilter("lincRNA"), SeqNameFilter("Y")))
length(linkY)
```

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```
## [1] 48

## We get now all transcripts for these genes.
txs <- select(edb, keys=linkY, columns=c("TXID", "TXSEQSTART", "TXBIOTYPE"),
              keytype="GENEID")
nrow(txs)

## [1] 66

## Alternatively, we could specify/pass the filters with the keys argument.
txs <- select(edb, keys=list(GeneBiotypeFilter("lincRNA"), SeqNameFilter("Y")),
              columns=c("TXID", "TXSEQSTART", "TXBIOTYPE"))

## Note: ordering of the results might not match ordering of keys!
nrow(txs)

## [1] 66
```

The version number of R and packages loaded for generating the vignette were:

```
## R version 4.0.0 (2020-04-24)
## Platform: x86_64-pc-linux-gnu (64-bit)
## Running under: Ubuntu 18.04.4 LTS
##
## Matrix products: default
## BLAS: /home/biocbuild/bbs-3.11-bioc/R/lib/libRblas.so
## LAPACK: /home/biocbuild/bbs-3.11-bioc/R/lib/libRlapack.so
##
## locale:
## [1] LC_CTYPE=en_US.UTF-8 LC_NUMERIC=C
## [3] LC_TIME=en_US.UTF-8 LC_COLLATE=C
## [5] LC_MONETARY=en_US.UTF-8 LC_MESSAGES=en_US.UTF-8
## [7] LC_PAPER=en_US.UTF-8 LC_NAME=C
## [9] LC_ADDRESS=C LC_TELEPHONE=C
## [11] LC_MEASUREMENT=en_US.UTF-8 LC_IDENTIFICATION=C
##
## attached base packages:
## [1] parallel stats4 stats graphics grDevices utils datasets
## [8] methods base
##
## other attached packages:
## [1] EnsDb.Hsapiens.v75_2.99.0
## [2] ensemblDb_2.12.0
## [3] AnnotationFilter_1.12.0
## [4] TxDb.Hsapiens.UCSC.hg19.knownGene_3.2.2
## [5] GenomicFeatures_1.40.0
## [6] GenomicRanges_1.40.0
```

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```
## [7] GenomeInfoDb_1.24.0
## [8] GO.db_3.10.0
## [9] hgu95av2.db_3.2.3
## [10] AnnotationForge_1.30.0
## [11] org.Hs.eg.db_3.10.0
## [12] AnnotationDbi_1.50.0
## [13] IRanges_2.22.0
## [14] S4Vectors_0.26.0
## [15] Biobase_2.48.0
## [16] BiocGenerics_0.34.0
## [17] DBI_1.1.0
## [18] knitr_1.28
##
## loaded via a namespace (and not attached):
## [1] httr_1.4.1             bit64_0.9-7
## [3] assertthat_0.2.1      askpass_1.1
## [5] BiocManager_1.30.10   highr_0.8
## [7] BiocFileCache_1.12.0  blob_1.2.1
## [9] GenomeInfoDbData_1.2.3 Rsamtools_2.4.0
## [11] yaml_2.2.1            progress_1.2.2
## [13] pillar_1.4.3          RSQLite_2.2.0
## [15] lattice_0.20-41       glue_1.4.0
## [17] digest_0.6.25         XVector_0.28.0
## [19] htmltools_0.4.0       Matrix_1.2-18
## [21] XML_3.99-0.3          pkgconfig_2.0.3
## [23] biomaRt_2.44.0        zlibbioc_1.34.0
## [25] purrr_0.3.4           BiocParallel_1.22.0
## [27] tibble_3.0.1          openssl_1.4.1
## [29] ellipsis_0.3.0        SummarizedExperiment_1.18.0
## [31] lazyeval_0.2.2        magrittr_1.5
## [33] crayon_1.3.4          memoise_1.1.0
## [35] evaluate_0.14         tools_4.0.0
## [37] prettyunits_1.1.1     hms_0.5.3
## [39] BiocStyle_2.16.0      lifecycle_0.2.0
## [41] matrixStats_0.56.0    stringr_1.4.0
## [43] DelayedArray_0.14.0   Biostrings_2.56.0
## [45] compiler_4.0.0        rlang_0.4.5
## [47] grid_4.0.0            RCurl_1.98-1.2
## [49] rappdirs_0.3.1        bitops_1.0-6
## [51] rmarkdown_2.1         curl_4.3
## [53] R6_2.4.1              GenomicAlignments_1.24.0
## [55] dplyr_0.8.5           rtracklayer_1.48.0
## [57] bit_1.1-15.2          ProtGenerics_1.20.0
## [59] stringi_1.4.6         Rcpp_1.0.4.6
## [61] vctrs_0.2.4           dbplyr_1.4.3
```

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```
## [63] tidyselect_1.0.0      xfun_0.13
```