

Creating select Interfaces for custom Annotation resources

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October 29, 2025

1 Introduction

The most common interface for retrieving data in *Bioconductor* is now the `select` method. The interface provides a simple way of extracting data.

There are really 4 methods that work together to allow a `select` interface. The 1st one is `columns`, which tells you about what kinds of values you can retrieve as columns in the final result.

```
library(RSQLite)
library(Homo.sapiens)

## Loading required package: OrganismDbi
## Loading required package: Seqinfo
## Loading required package: GenomicFeatures
## Loading required package: GenomicRanges
## Loading required package: GO.db
##
## Loading required package: org.Hs.eg.db
##
## Loading required package: TxDb.Hsapiens.UCSC.hg19.knownGene
columns(Homo.sapiens)

## [1] "ACCNUM"      "ALIAS"       "CDSCHROM"    "CDSSEND"
## [5] "CDSID"       "CDSNAME"     "CDSPHASE"    "CDSSTART"
## [9] "CDSSTRAND"   "DEFINITION"  "ENSEMBL"     "ENSEMBLPROT"
## [13] "ENSEMBLTRANS" "ENTREZID"    "ENZYME"      "EVIDENCE"
## [17] "EVIDENCEALL" "EXONCHROM"   "EXONEND"     "EXONID"
## [21] "EXONNAME"    "EXONRANK"    "EXONSTART"   "EXONSTRAND"
## [25] "GENEID"      "GENENAME"    "GENETYPE"    "GO"
## [29] "GOALL"       "GOID"        "IPI"         "MAP"
## [33] "OMIM"        "ONTOLOGY"    "ONTOLOGYALL" "PATH"
```

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```
## [37] "PFAM"          "PMID"          "PROSITE"       "REFSEQ"
## [41] "SYMBOL"        "TERM"          "TXCHROM"       "TXEND"
## [45] "TXID"          "TXNAME"        "TXSTART"       "TXSTRAND"
## [49] "TXTYPE"        "UCSCKG"        "UNIPROT"
```

The next method is `keytypes` which tells you the kinds of things that can be used as keys.

```
keytypes(Homo.sapiens)

## [1] "ACCNUM"        "ALIAS"         "CDSID"         "CDSNAME"
## [5] "DEFINITION"    "ENSEMBL"       "ENSEMBLPROT"  "ENSEMBLTRANS"
## [9] "ENTREZID"      "ENZYME"        "EVIDENCE"      "EVIDENCEALL"
## [13] "EXONID"        "EXONNAME"      "GENEID"        "GENENAME"
## [17] "GENETYPE"      "GO"            "GOALL"         "GOID"
## [21] "IPI"           "MAP"           "OMIM"          "ONTOLOGY"
## [25] "ONTOLOGYALL"   "PATH"          "PFAM"          "PMID"
## [29] "PROSITE"       "REFSEQ"        "SYMBOL"        "TERM"
## [33] "TXID"          "TXNAME"        "UCSCKG"        "UNIPROT"
```

The third method is `keys` which is used to retrieve all the viable keys of a particular type.

```
k <- head(keys(Homo.sapiens, keytype="ENTREZID"))
k
## [1] "1" "2" "9" "10" "11" "12"
```

And finally there is `select`, which extracts data by using values supplied by the other method.

```
result <- select(Homo.sapiens, keys=k,
                 columns=c("TXNAME", "TXSTART", "TXSTRAND"),
                 keytype="ENTREZID")

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

head(result)

##   ENTREZID          TXNAME TXSTRAND TXSTART
## 1         1 ENST00000596924.1_3      - 58856544
## 2         1 ENST00000263100.8_8      - 58856549
## 3         1 ENST00000850949.1_1      - 58856549
## 4         1 ENST00000850950.1_1      - 58856549
## 5         1 ENST00000600123.5_4      - 58858220
## 6         1 ENST00000595014.1_3      - 58858224
```

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But why would we want to implement these specific methods? It's a fair question. Why would we want to write a select interface for our annotation data? Why not just save a .rda file to the data directory and be done with it? There are basically two reasons for this. The 1st reason is convenience for end users. When your end users can access your data using the same four methods that they use everywhere else, they will have a more effortless time retrieving their data. And things that benefit your users benefit you.

The second reason is that by enabling a consistent interface across all annotation resources, we allow for things to be used in a programmatic manner. By implementing a select interface, we are creating a universal API for the whole project.

Lets look again at the example I described above and think about what is happening. The *Homo.sapiens* package is able to integrate data from many different resources largely because the separate resources all implemented a select method. This allows the *OrganismDbi* package to pull together resources from *org.Hs.eg.db*, *GO.db* and *TxDb.Hsapiens.UCSC.hg19.knownGene*.



Figure 1: Packages and relationships represented by the *Homo.sapiens* package

If these packages all exposed different interfaces for retrieving the data, then it would be a lot more challenging to retrieve it, and writing general code that retrieved the appropriate data would be a lost cause. So implementing a set of select methods is a way to convert your package from a data file into an actual resource.

2 Creating other kinds of Annotation packages

A few more automated options already exist for generating specific kinds of annotation packages. For users who seek to make custom chip packages, users should see the *SQLForge: An easy way to create a new annotation package with a standard database schema*. in the *AnnotationForge* package. And, for users who seek to make a probe package, there is another vignette called *Creating probe packages* that is also in the *AnnotationForge* package. And finally, for custom organism packages users should look at the manual page for `makeOrgPackageFromNCBI`. This function will attempt to make you an simplified organism package from NCBI resources. However, this function is not meant as a way to refresh annotation packages between releases. It is only meant for people who are working on less popular model organisms (so that annotations can be made available in this format).

But what if you had another kind of web resource or database and you wanted to expose it to the world using something like this new `select` method interface? How could you go about this?

3 Retrieving data from a web resource

If you choose to expose a web resource, then you will need to learn some skills for retrieving that data from the web. The *R* programming language has tools that can help you interact with web resources, pulling down files that are tab delimited or formatted as XML etc. There are also packages that can help you parse what you retrieve. In this section we will describe some of these resources in the context of the uniprot web service, and give examples to demonstrate how you can expose resources like this for your own purposes.

These days many web services are exposed using a representational state transfer or RESTful interface. An example of this are the services offered at Uniprot. Starting with the uniprot base URI you can add details to simply indicate what it is that you wish to retrieve.

So in the case of Uniprot the base URI for the service we want today is this:

```
http://www.uniprot.org/uniprot/
```

This URI can be extended to retrieve individual uniprot records by specifying a query argument like this:

```
http://www.uniprot.org/uniprot/?query=P13368
```

We can then request multiple records like this:

```
http://www.uniprot.org/uniprot/?query=P13368+or+Q6GZX4
```

And we can ask that the records be returned to us in tabular form by adding another argument like this.

```
http://www.uniprot.org/uniprot/?query=P13368+or+Q6GZX4&format=tab
```

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As you might guess, each RESTful interface is a little different, but you can easily see how once you read the documentation for a given RESTful interface, you can start to retrieve the data in *R*. Here is an example.

```
uri <- 'http://rest.uniprot.org/uniprotkb/search?query='
ids <- c('P13368', 'Q6GZX4')
idStr <- paste(ids, collapse=utils::URLencode(" OR "))
format <- '&format=tsv'
fullUri <- paste0(uri,idStr,format)
uquery <- .get_file(fullUri)
read.delim(uquery)

##      Entry  Entry.Name Reviewed      Protein.names
## 1 P13368  7LESS_DROME reviewed  Protein sevenless (EC 2.7.10.1)
## 2 Q6GZX4  001R_FRG3G reviewed Putative transcription factor 001R
##           Gene.Names                Organism Length
## 1 sev HD-265 CG18085  Drosophila melanogaster (Fruit fly)  2554
## 2           FV3-001R Frog virus 3 (isolate Goorha) (FV-3)  256
```

Exercise 1

If you use the *columns* argument you can also specify which columns you want returned. So for example, you can choose to only have the sequence and id columns returned like this:

<http://www.uniprot.org/uniprot/?query=P13368+or+Q6GZX4&format=tab&columns=id,sequence>

Use this detail about the uniprot web service along with what was learned above to write a function that takes a character vector of uniprot IDs and another character vector of columns arguments and then returns the appropriate values. Be careful to filter out any extra records that the service returns.

Solution:

```
getUniprotGoodies <- function(query, columns)
{
  ## query and columns start as a character vectors
  qstring <- paste(query, collapse="+or+")
  cstring <- paste(columns, collapse=",")
  uri <- 'http://www.uniprot.org/uniprot/?query='
  fullUri <- paste0(uri,qstring,'&format=tab&columns=',cstring)
  dat <- read.delim(fullUri)
  ## now remove things that were not in the specific original query...
  dat <- dat[dat[,1] %in% query,]
  dat
}
```

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3.1 Parsing XML

Data for the previous example were downloaded from Uniprot in tab-delimited format. This is a convenient output to work with but unfortunately not always available. XML is still very common and it is useful to have some familiarity with parsing it. In this section we give a brief overview to using the *XML* package for navigating XML data.

The *XML* package provides functions to parse XML in both the tree-based DOM (document object model) or the event-driven SAX (Simple API for XML). We will use the DOM approach. The XML is first parsed into a tree-structure where the different elements of the data are nodes. The elements are processed by traversing the tree and generating a user-level representation of the nodes. XPath syntax is used to traverse the nodes. A detailed description of XPath can be found at <http://www.w3.org/xml>.

Retrieve the data: Data will be retrieved for the same id's as in the previous example. Unlike tab-delimited, the XML queries cannot be subset by column so the full record will be returned for each id. Details for what is possible with each type of data retrieval are found at <http://www.uniprot.org/faq/28>.

Parse the XML into a tree structure with `xmlTreeParse`. When `useInternalNodes=TRUE` and no `handlers` are specified the return value is a reference to C-level nodes. This storage mode allows us to traverse the tree of data in C instead of R objects.

```
library(XML)
uri <- "http://rest.uniprot.org/uniprotkb/search?query=P13368%20R%20Q6GZX4&format=xml"
fl <- tempfile()
download.file(uri, fl)
xml <- xmlTreeParse(fl, useInternalNodes=TRUE)
```

XML namespace: XML pages can have namespaces which facilitate the use of different XML vocabularies by resolving conflicts arising from identical tags. Namespaces are represented by a uri pointing to an XML schema page. When a namespace is defined on a node in an XML document it must be included in the XPath expression.

Use the `xmlNamespaceDefinitions` function to check if the XML has a namespace.

```
defs <- xmlNamespaceDefinitions(xml, recursive=TRUE)
defs
## [[1]]
## $id
## [1] ""
##
## $uri
## [1] "http://uniprot.org/uniprot"
##
## $local
## [1] TRUE
```

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```
##  
## attr("class")  
## [1] "XMLNamespaceDefinition"  
##  
## $xsi  
## $id  
## [1] "xsi"  
##  
## $uri  
## [1] "http://www.w3.org/2001/XMLSchema-instance"  
##  
## $local  
## [1] TRUE  
##  
## attr("class")  
## [1] "XMLNamespaceDefinition"  
##  
## attr("class")  
## [1] "XMLNamespaceDefinitions"
```

The presence of uri's confirm there is a namespace. Alternatively we could have looked at the XML nodes for declarations of the form `xmlns:myNamespace="http://www.namespace.org"`. We organize the namespaces and will use them directly in parsing.

```
ns <- structure(sapply(defs, function(x) x$uri), names=names(defs))
```

Parsing with XPath: There are two high level 'entry' nodes which represent the two id's requested in the original query.

```
entry <- getNodeSet(xml, "//ns:entry", "ns")  
xmlSize(entry)  
  
## [1] 3
```

To get an idea of the data structure we first list the attributes of the top nodes and extract the names.

```
nms <- xpathSApply(xml, "//ns:entry/ns:name", xmlValue, namespaces="ns")  
attrs <- xpathSApply(xml, "//ns:entry", xmlAttrs, namespaces="ns")  
names(attrs) <- nms  
attrs  
  
## $`7LESS_DROME`  
##      dataset      created      modified      version  
## "Swiss-Prot" "1990-01-01" "2025-06-18"      "209"  
##
```

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```
## `$`001R_FRG3G`  
##      dataset      created      modified      version  
## "Swiss-Prot" "2011-06-28" "2025-04-09"      "45"  
##  
## $A0ABM1YJC1_AEDAL  
##      dataset      created      modified      version  
##      "TrEMBL" "2025-10-08" "2025-10-08"      "1"
```

Next, inspect the direct children of each node.

```
fun1 <- function(elt) unique(names(xmlChildren(elt)))  
xpathApply(xml, "//ns:entry", fun1, namespaces="ns")  
  
## [[1]]  
## [1] "accession"      "name"           "protein"  
## [4] "gene"           "organism"       "reference"  
## [7] "comment"        "dbReference"    "proteinExistence"  
## [10] "keyword"        "feature"        "evidence"  
## [13] "sequence"  
##  
## [[2]]  
## [1] "accession"      "name"           "protein"  
## [4] "gene"           "organism"       "organismHost"  
## [7] "reference"      "comment"        "dbReference"  
## [10] "proteinExistence" "keyword"        "feature"  
## [13] "evidence"      "sequence"  
##  
## [[3]]  
## [1] "accession"      "name"           "protein"  
## [4] "organism"      "reference"      "dbReference"  
## [7] "proteinExistence" "keyword"        "evidence"  
## [10] "sequence"
```

Query Q6GZX4 has 2 'feature' nodes and query P13368 has 48.

```
Q6GZX4 <- "//ns:entry[ns:accession='Q6GZX4']/ns:feature"  
xmlSize(getNodeSet(xml, Q6GZX4, namespaces="ns"))  
## [1] 1  
  
P13368 <- "//ns:entry[ns:accession='P13368']/ns:feature"  
xmlSize(getNodeSet(xml, P13368, namespaces="ns"))  
## [1] 53
```

List all possible values for the 'type' attribute of the 'feature' nodes.

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```
path <- "//ns:feature"
unique(xpathSApply(xml, path, xmlGetAttr, "type", namespaces="ns"))

## [1] "chain"                "topological domain"
## [3] "transmembrane region"  "domain"
## [5] "repeat"                "region of interest"
## [7] "compositionally biased region" "active site"
## [9] "binding site"         "modified residue"
## [11] "glycosylation site"   "mutagenesis site"
## [13] "sequence conflict"
```

XPath allows the construction of complex queries to pull out specific subsets of data. Here we extract the features with 'type=sequence conflict' for query P13368.

```
path <- "//ns:entry[ns:accession='P13368']/ns:feature[@type='sequence conflict']"
data.frame(t(xpathSApply(xml, path, xmlAttrs, namespaces="ns")))

##           type           description evidence ref
## 1 sequence conflict      In Ref. 1; AAA28882.      9  1
## 2 sequence conflict      In Ref. 3; AAF47992.      9  3
## 3 sequence conflict      In Ref. 3; AAF47992.      9  3
## 4 sequence conflict      In Ref. 3; AAF47992.      9  3
## 5 sequence conflict      In Ref. 3; AAF47992.      9  3
## 6 sequence conflict In Ref. 2; CAA31960/CAB55310.      9  2
## 7 sequence conflict      In Ref. 1; AAA28882.      9  1
```

Put the sequences in an AAStringSet and add the names.

```
library(Biostrings)

## Loading required package: XVector

##
## Attaching package: 'Biostrings'

## The following object is masked from 'package:base':
##
##   strsplit

path <- "//ns:entry/ns:sequence"
seqs <- xpathSApply(xml, path, xmlValue, namespaces="ns")
aa <- AAStringSet(unlist(lapply(seqs, function(elt) gsub("\n", "", elt)),
  use.names=FALSE))
names(aa) <- nms
aa

## AAStringSet object of length 3:
##      width seq                                     names
## [1] 2554 MTMFWQQNVDHQSDEQDKQAKG...TLREVPLKDKQLYANEGVSRL 7LESS_DROME
```

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```
## [2] 256 MAFSAEDVLKEYDRRRRMEALL...VLYDDFRKIYTDLGWKFTPL 001R_FRG3G
## [3] 407 MADTYDEYEETHVTRKSVTTF...SQLRQFRFFQNIHRKQATMIQ A0ABM1YJC1_AEDAL
```

4 Setting up a package to expose a web service

In order to expose a web service using select, you will need to create an object that will be loaded at the time when the package is loaded. Unlike with a database, the purpose of this object is pretty much purely for dispatch. We just need `select` and it's friends to know which select method to call

The first step is to create an object. Creating an object is simple enough:

```
setClass("uniprot", representation(name="character"),
         prototype(name="uniprot"))
```

Once you have a class defined, all you need is to make an instance of this class. Making an instance is easy enough:

```
uniprot <- new("uniprot")
```

But of course it's a little more complicated because one of these objects will need to be spawned up whenever our package loads. This is accomplished by calling the `.onLoad` function in the `zzz.R` file. The following code will create an object, and then assign it to the package namespace as the package loads.

```
.onLoad <- function(libname, pkgname)
{
  ns <- asNamespace(pkgname)
  uniprot <- new("uniprot")
  assign("uniprot", uniprot, envir=ns)
  namespaceExport(ns, "uniprot")
}
```

5 Creating package accessors for a web service

At this point you have all that you need to know in order to implement `keytype`, `columns`, `keys` and `select` for your package. In this section we will explore how you could implement some of these if you were making a package that exposed uniprot.

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5.1 Example: creating `keytypes` and `columns` methods

The `keytype` and `columns` methods are always the 1st ones you should implement. They are the easiest, and their existence is required to be able to use `keys` or `select`. In this simple case we only have one value that can be used as a keytype, and that is a UNIPROT ID.

```
setMethod("keytypes", "uniprot", function(x){return("UNIPROT")})
uniprot <- new("uniprot")
keytypes(uniprot)

## [1] "UNIPROT"
```

So what about columns? Well it's not a whole lot more complicated in this case since we are limited to things that we can return from the web service. Since this is just an example, lets limit it to the following fields: "ID", "SEQUENCE", "ORGANISM".

```
setMethod("columns", "uniprot",
          function(x){return(c("ID", "SEQUENCE", "ORGANISM"))})
columns(uniprot)

## [1] "ID"      "ORGANISM" "SEQUENCE"
```

Also, notice how for both `keytypes` and `columns` I am using all capital letters. This is a style adopted throughout the project.

5.2 Example 2: creating a `select` method

At this point we have enough to be able to make a select method.

Exercise 2

Using what you have learned above, and the helper function from earlier, define a `select` method. This `select` method will have a default `keytype` of "UNIPROT".

Solution:

```
.select <- function(x, keys, columns){
  colsTranslate <- c(id='ID', sequence='SEQUENCE', organism='ORGANISM')
  columns <- names(colsTranslate)[colsTranslate %in% columns]
  getUniprotGoodies(query=keys, columns=columns)
}
setMethod("select", "uniprot",
          function(x, keys, columns, keytype)
{
  .select(keys=keys, columns=columns)
})
```

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```
select(uniprot, keys=c("P13368", "P20806"), columns=c("ID", "ORGANISM"))
```

6 Retrieving data from a database resource

If your package is retrieving data from a database, then there are some additional skills you will need to be able to interface with this database from *R*. This section will introduce you to those skills.

6.1 Getting a connection

If all you know is the name of the SQLite database, then to get a DB connection you need to do something like this:

```
drv <- SQLite()
library("org.Hs.eg.db")
con_hs <- dbConnect(drv, dbname=org.Hs.eg_dbfile())
con_hs
```

In cases where the connection is created on package load, we can do something like below:

```
org.Hs.eg.db$conn
## or better we can use a helper function to wrap this:
AnnotationDbi::dbconn(org.Hs.eg.db)
## or we can just call the provided convenience function
## from when this package loads:
org.Hs.eg_dbconn()
```

6.2 Getting data out

Now we just need to get our data out of the DB. There are several useful functions for doing this. Most of these come from the *RSQLite* or *DBI* packages. For the sake of simplicity, I will only discuss those that are immediately useful for exploring and extracting data from a database in this vignette. One pair of useful methods are the `dbListTables` and `dbListFields` which are useful for exploring the schema of a database.

```
head(dbListTables(con_hs))
## [1] "accessions"          "alias"                "chrlengths"
## [4] "chromosome_locations" "chromosomes"          "cytogenetic_locations"
dbListFields(con_hs, "alias")
## [1] "_id"                 "alias_symbol"
```

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For actually executing SQL to retrieve data, you probably want to use something like `dbGetQuery`. The only caveat is that this will actually require you to know a little SQL.

```
dbGetQuery(con_hs, "SELECT * FROM metadata")

##           name                                     value
## 1  DBSCHEMAVERSION                               2.1
## 2           Db type                               OrgDb
## 3 Supporting package                             AnnotationDbi
## 4           DBSCHEMA                             HUMAN_DB
## 5           ORGANISM                             Homo sapiens
## 6           SPECIES                               Human
## 7  EGSOURCEDATE                                  2025-Sep24
## 8  EGSOURCENAME                                 Entrez Gene
## 9  EGSOURCEURL                                  ftp://ftp.ncbi.nlm.nih.gov/gene/DATA
## 10 CENTRALID                                     EG
## 11          TAXID                                 9606
## 12  GOSOURCENAME                                 Gene Ontology
## 13  GOSOURCEURL  http://current.geneontology.org/ontology/go-basic.obo
## 14  GOSOURCEDATE                                  2025-07-22
## 15  GOEGSOURCEDATE                                2025-Sep24
## 16  GOEGSOURCENAME                               Entrez Gene
## 17  GOEGSOURCEURL                                  ftp://ftp.ncbi.nlm.nih.gov/gene/DATA
## 18  KEGGSOURCENAME                               KEGG GENOME
## 19  KEGGSOURCEURL                                  ftp://ftp.genome.jp/pub/kegg/genomes
## 20  KEGGSOURCEDATE                                2011-Mar15
## 21  GPSOURCENAME                                UCSC Genome Bioinformatics (Homo sapiens)
## 22  GPSOURCEURL  ftp://hgdownload.cse.ucsc.edu/goldenPath/hg38/database
## 23  GPSOURCEDATE                                  UTC-Sep26
## 24  ENSOURCEDATE                                  2025-Sep03
## 25  ENSOURCENAME                                 Ensembl
## 26  ENSOURCEURL                                  ftp://ftp.ensembl.org/pub/current_fasta
## 27  UPSOURCENAME                                 Uniprot
## 28  UPSOURCEURL                                  http://www.UniProt.org/
## 29  UPSOURCEDATE                                  Sat Oct 4 09:48:11 2025
```

6.3 Some basic SQL

The good news is that SQL is pretty easy to learn. Especially if you are primarily interested in just retrieving data from an existing database. Here is a quick run-down to get you started on writing simple SELECT statements. Consider a table that looks like this:

This statement:

```
SELECT bar FROM sna;
```

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Table sna	
foo	bar
1	baz
2	boo

Tells SQL to get the "bar" field from the "foo" table. If we wanted the other field called "sna" in addition to "bar", we could have written it like this:

```
SELECT foo, bar FROM sna;
```

Or even this (* is a wildcard character here)

```
SELECT * FROM sna;
```

Now lets suppose that we wanted to filter the results. We could also have said something like this:

```
SELECT * FROM sna WHERE bar='boo';
```

That query will only retrieve records from foo that match the criteria for bar. But there are two other things to notice. First notice that a single = was used for testing equality. Second notice that I used single quotes to demarcate the string. I could have also used double quotes, but when working in *R* this will prove to be less convenient as the whole SQL statement itself will frequently have to be wrapped as a string.

What if we wanted to be more general? Then you can use LIKE. Like this:

```
SELECT * FROM sna WHERE bar LIKE 'boo\%';
```

That query will only return records where bar starts with "boo", (the % character is acting as another kind of wildcard in this context).

You will often find that you need to get things from two or more different tables at once. Or, you may even find that you need to combine the results from two different queries. Sometimes these two queries may even come from the same table. In any of these cases, you want to do a join. The simplest and most common kind of join is an inner join. Lets suppose that we have two tables:

Table sna		Table fu	
foo	bar	foo	bo
1	baz	1	hi
2	boo	2	ca

And we want to join them where the records match in their corresponding "foo" columns. We can do this query to join them:

```
SELECT * FROM sna, fu WHERE sna.foo=fu.foo;
```

Something else we can do is tidy this up by using aliases like so:

```
SELECT * FROM sna AS s, fu AS f WHERE s.foo=f.foo;
```

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This last trick is not very useful in this particular example since the query ended up being longer than we started with, but is still great for other cases where queries can become really long.

6.4 Exploring the SQLite database from R

Now that we know both some SQL and also about some of the methods in *DBI* and *RSQLite* we can begin to explore the underlying database from R. How should we go about this? Well the 1st thing we always want to know are what tables are present. We already know how to learn this:

```
head(dbListTables(con_hs))  
## [1] "accessions"          "alias"                "chrlengths"  
## [4] "chromosome_locations" "chromosomes"          "cytogenetic_locations"
```

And we also know that once we have a table we are curious about, we can then look up it's fields using `dbListFields`

```
dbListFields(con_hs, "chromosomes")  
## [1] "_id"          "chromosome"
```

And once we know something about which fields are present in a table, we can compose a SQL query. perhaps the most straightforward query is just to get all the results from a given table. We know that the SQL for that should look like:

```
SELECT * FROM chromosomes;
```

So we can now call a query like that from R by using `dbGetQuery`:

```
head(dbGetQuery(con_hs, "SELECT * FROM chromosomes"))  
##   _id chromosome  
## 1   1         19  
## 2   2         12  
## 3   3          8  
## 4   4          8  
## 5   5          8  
## 6   6         14
```

Exercise 3

Now use what you have learned to explore the [org.Hs.eg.db](#) database. Now find the table for chromosome locations in the [org.Hs.eg.db](#) database and extract it into R. How many chromosomes are present in this table? Write a SQL query that will retrieve chromosome locations from this table that are in chromosome 1.

Solution:

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```
head(dbGetQuery(con_hs, "SELECT * FROM chromosome_locations"))
## Then only retrieve human records
## Query: SELECT * FROM Anopheles_gambiae WHERE species='HOMSA'
head(dbGetQuery(con_hs, "SELECT * FROM chromosome_locations WHERE seqname='1'"))
dbDisconnect(con_hs)
```

7 Setting up a package to expose a SQLite database object

For the sake of simplicity, let's look at an existing example of this in the [org.Hs.eg.db](#) package. This package contains a .sqlite database inside of the extdata directory. There are a couple of important details though about databases like these. The 1st is that we recommend that the database have the same name as the package, but end with the extension .sqlite. The second detail is that we recommend that the metadata table contain some important fields. This is the metadata from the current [org.Hs.eg.db](#) package.

##	name	value
## 1	DBSCHEMAVERSION	2.1
## 2	Db type	OrgDb
## 3	Supporting package	AnnotationDbi
## 4	DBSCHEMA	HUMAN_DB
## 5	ORGANISM	Homo sapiens
## 6	SPECIES	Human
## 7	EGSOURCEDATE	2025-Sep24
## 8	EGSOURCENAME	Entrez Gene
## 9	EGSOURCEURL	ftp://ftp.ncbi.nlm.nih.gov/gene/DATA
## 10	CENTRALID	EG
## 11	TAXID	9606
## 12	GOSOURCENAME	Gene Ontology
## 13	GOSOURCEURL	http://current.geneontology.org/ontology/go-basic.obo
## 14	GOSOURCEDATE	2025-07-22
## 15	GOEGSOURCEDATE	2025-Sep24
## 16	GOEGSOURCENAME	Entrez Gene
## 17	GOEGSOURCEURL	ftp://ftp.ncbi.nlm.nih.gov/gene/DATA
## 18	KEGGSOURCENAME	KEGG GENOME
## 19	KEGGSOURCEURL	ftp://ftp.genome.jp/pub/kegg/genomes
## 20	KEGGSOURCEDATE	2011-Mar15
## 21	GPSOURCENAME	UCSC Genome Bioinformatics (Homo sapiens)
## 22	GPSOURCEURL	ftp://hgdownload.cse.ucsc.edu/goldenPath/hg38/database
## 23	GPSOURCEDATE	UTC-Sep26
## 24	ENSOURCEDATE	2025-Sep03
## 25	ENSOURCENAME	Ensembl

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```
## 26      ENSOURCEURL      ftp://ftp.ensembl.org/pub/current_fasta
## 27      UPSOURCENAME      Uniprot
## 28      UPSOURCEURL      http://www.UniProt.org/
## 29      UPSOURCEDATE      Sat Oct  4 09:48:11 2025
```

As you can see there are a number of very useful fields stored in the metadata table and if you list the equivalent table for other packages you will find even more useful information than you find here. But the most important fields here are actually the ones called "package" and "Db type". Those fields specify both the name of the package with the expected class definition, and also the name of the object that this database is expected to be represented by in the R session respectively. If you fail to include this information in your metadata table, then `loadDb` will not know what to do with the database when it is called. In this case, the class definition has been stored in the `AnnotationDbi` package, but it could live anywhere you need it too. By specifying the metadata field, you enable `loadDb` to find it.

Once you have set up the metadata you will need to create a class for your package that extends the `AnnotationDb` class. In the case of the `org.Hs.eg.db` package, the class is defined to be a `OrgDb` class.

```
showClass("OrgDb")
```

Finally the `.onLoad` call for your package will have to contain code that will call the `loadDb` method. This is what it currently looks like in the `org.Hs.eg.db` package.

```
sPkgname <- sub(".db$", "", pkgname)
db <- loadDb(system.file("extdata", paste(sPkgname,
    ".sqlite", sep=""), package=pkgname, lib.loc=libname),
    packageName=pkgname)
dbNewname <- AnnotationDbi::dbObjectName(pkgname, "OrgDb")
ns <- asNamespace(pkgname)
assign(dbNewname, db, envir=ns)
namespaceExport(ns, dbNewname)
```

When the code above is run (at load time) the name of the package (AKA "pkgname", which is a parameter that will be passed into `.onLoad`) is then used to derive the name for the object. Then that name, is used by `onload` to create an `InparanoidDb` object. This object is then assigned to the namespace for this package so that it will be loaded for the user.

8 Creating package accessors for databases

At this point, all that remains is to create the means for accessing the data in the database. This should prove a lot less difficult than it may initially sound. For the new interface, only the four methods that were described earlier are really required: `columns`, `keytypes`, `keys` and `select`.

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In order to do this you need to know a small amount of SQL and a few tricks for accessing the database from R. The point of providing these 4 accessors is to give users of these packages a more unified experience when retrieving data from the database. But other kinds of accessors (such as those provided for the *TxDB* objects) may also be warranted.

8.1 Examples: creating a `columns` and `keytypes` method

Now lets suppose that we want to define a `columns` method for our *org.Hs.eg.db* object. And lets also suppose that we want is for it to tell us about the actual organisms for which we can extract identifiers. How could we do that?

```
.cols <- function(x)
{
  con <- AnnotationDbi::dbconn(x)
  list <- dbListTables(con)
  ## drop unwanted tables
  unwanted <- c("map_counts", "map_metadata", "metadata")
  list <- list[!list %in% unwanted]
  # use on.exit to disconnect
  # on.exit(dbDisconnect(con))
  ## Then just to format things in the usual way
  toupper(list)
}

## Then make this into a method
setMethod("columns", "OrgDb", .cols(x))
## Then we can call it
columns(org.Hs.eg.db)
```

Notice again how I formatted the output to all uppercase characters? This is just done to make the interface look consistent with what has been done before for the other `select` interfaces. But doing this means that we will have to do a tiny bit of extra work when we implement our other methods.

Exercise 4

Now use what you have learned to define a method for `keytypes` on *org.Hs.eg.db*. The `keytypes` method should return the same results as `columns` (in this case). What if you needed to translate back to the lowercase table names? Also write a quick helper function to do that.

Solution:

```
setMethod("keytypes", "OrgDb", function(x) .cols(x))
## Then we can call it
keytypes(org.Hs.eg.db)
```

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```
## refactor of .cols
.getLCcolnames <- function(x)
{
  con <- AnnotationDbi::dbconn(x)
  list <- dbListTables(con)
  ## drop unwanted tables
  unwanted <- c("map_counts", "map_metadata", "metadata")
  # use on.exit to disconnect
  # on.exit(dbDisconnect(con))
  list[!list %in% unwanted]
}
.cols <- function(x)
{
  list <- .getLCcolnames(x)
  ## Then just to format things in the usual way
  toupper(list)
}
## Test:
columns(org.Hs.eg.db)

## new helper function:
.getTableNames <- function(x)
{
  LC <- .getLCcolnames(x)
  UC <- .cols(x)
  names(UC) <- LC
  UC
}
.getTableNames(org.Hs.eg.db)
```

8.2 Example: creating a `keys` method

Exercise 5

Now define a method for `keys` on the example `org.Hs.eg.db`. This function can be a helper function to make it easy to extract column values after filtering the rows. The `keys` method should return the keys from a given organism based on the appropriate keytype. For example, if a table has rows that correspond to both human and non-human IDs, it will be necessary to filter out the human rows from the result.

Solution:

```
.keys <- function(x, keytype)
{
```

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```
## translate keytype back to table name
tabNames <- .getTableNames(x)
lckeytype <- names(tabNames[tabNames %in% keytype])
## get a connection
con <- AnnotationDbi::dbconn(x)
sql <- paste("SELECT _id FROM", lckeytype, "WHERE species != 'HOMSA'")
res <- dbGetQuery(con, sql)
res <- as.vector(t(res))
dbDisconnect(con)
res
}

setMethod("keys", "ExampleDbClass", .keys)
## Then we would call it
keys(example.db, "TRICHOPLAX_ADHAERENS")
```

9 Creating a database resource from available data

Sometimes you may have a lot of data that you want to organize into a database. Or you may have another existing database that you wish to convert into a SQLite database. This section will deal with some simple things you can do to create and import a SQLite database of your own.

9.1 Making a new connection

Then lets make a new database. Notice that we specify the database name with "dbname" This allows it to be written to disc instead of just memory.

```
drv <- dbDriver("SQLite")
dbname <- file.path(tempdir(), "myNewDb.sqlite")
con <- dbConnect(drv, dbname=dbname)
```

9.2 Importing data

Imagine that we want to reate a database and then put a table in it called genePheno to store the genes mutated and a phenotypes associated with each. Plan for genePheno to hold the following gene IDs and phenotypes (as a toy example):

```
data <- data.frame(id=c(22,7,15), string=c("Blue", "Red", "Green"))
```

Making the table is very simple, and just involves a create table statement.

```
CREATE Table genePheno (id INTEGER, string TEXT);
```

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The SQL create table statement just indicates what the table is to be called, as well as the different fields that will be present and the type of data each field is expected to contain.

```
dbExecute(con, "CREATE Table genePheno (id INTEGER, string TEXT)")  
## [1] 0
```

Then we can populate the SQL table:

```
sql <- "INSERT INTO genePheno VALUES ($id, $string)"  
dbExecute(con, sql, params=data)  
## [1] 3
```

We can check the content of the table with:

```
dbReadTable(con, "genePheno")  
##   id string  
## 1 22   Blue  
## 2  7    Red  
## 3 15   Green
```

9.3 Attaching other database resources

In SQLite it is possible to attach another database to your session and then query across both resources as if they were the same DB.

The SQL what we want looks quite simple:

```
ATTACH "TxDb.Hsapiens.UCSC.hg19.knownGene.sqlite" AS db;
```

So in R we need to do something similar to this:

```
db <- system.file("extdata", "TxDb.Hsapiens.UCSC.hg19.knownGene.sqlite",  
                  package="TxDb.Hsapiens.UCSC.hg19.knownGene")  
dbExecute(con, sprintf("ATTACH '%s' AS db",db))  
## [1] 0
```

Here we have attached a DB from one of the packages that this vignette required you to have installed, but we could have attached any SQLite database that we provided a path to.

Once we have attached the database, we can join to it's tables as if they were in our own database. All that is required is a prefix, and some knowledge about how to do joins in SQL. In the end the SQL to take advantage of the attached database looks like this:

```
SELECT * FROM db.gene AS dbg, genePheno AS gp
```

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```
WHERE dbg.gene_id=gp.id;
```

Then in R:

```
sql <- "SELECT * FROM db.gene AS dbg,  
       genePheno AS gp WHERE dbg.gene_id=gp.id"  
res <- dbGetQuery(con, sql)  
res  
  
##   gene_id _tx_id id string  
## 1     15 309633 15  Green  
## 2     15 309632 15  Green  
## 3     15 309630 15  Green  
## 4     15 309631 15  Green  
## 5     22 377545 22   Blue  
## 6     22 377546 22   Blue  
## 7     22 377547 22   Blue  
## 8     22 377548 22   Blue  
## 9     22 377549 22   Blue  
## 10    22 377550 22   Blue  
## 11    22 377551 22   Blue  
## 12    22 377552 22   Blue  
## 13    22 377553 22   Blue  
## 14    22 377554 22   Blue  
## 15    22 377555 22   Blue  
## 16    22 377556 22   Blue  
## 17    22 377557 22   Blue  
## 18    22 377558 22   Blue  
## 19    22 377559 22   Blue  
## 20    22 377560 22   Blue  
## 21    22 377561 22   Blue
```

9.4 Closing the DB connection

```
dbDisconnect(con)
```

10 Session information

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

```
sessionInfo()  
  
## R version 4.5.1 Patched (2025-08-23 r88802)  
## Platform: x86_64-pc-linux-gnu  
## Running under: Ubuntu 24.04.3 LTS  
##
```

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```
## Matrix products: default
## BLAS: /home/biocbuild/bbs-3.22-bioc/R/lib/libRblas.so
## LAPACK: /usr/lib/x86_64-linux-gnu/lapack/liblapack.so.3.12.0 LAPACK version 3.12.0
##
## locale:
## [1] LC_CTYPE=en_US.UTF-8 LC_NUMERIC=C
## [3] LC_TIME=en_GB 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
##
## time zone: America/New_York
## tzcode source: system (glibc)
##
## attached base packages:
## [1] stats4 stats graphics grDevices utils datasets methods
## [8] base
##
## other attached packages:
## [1] Biostrings_2.78.0
## [2] XVector_0.50.0
## [3] XML_3.99-0.19
## [4] Homo.sapiens_1.3.1
## [5] TxDb.Hsapiens.UCSC.hg19.knownGene_3.22.1
## [6] org.Hs.eg.db_3.22.0
## [7] GO.db_3.22.0
## [8] OrganismDbi_1.52.0
## [9] GenomicFeatures_1.62.0
## [10] GenomicRanges_1.62.0
## [11] Seqinfo_1.0.0
## [12] RSQLite_2.4.3
## [13] knitr_1.50
## [14] AnnotationForge_1.52.0
## [15] AnnotationDbi_1.72.0
## [16] IRanges_2.44.0
## [17] S4Vectors_0.48.0
## [18] Biobase_2.70.0
## [19] BiocGenerics_0.56.0
## [20] generics_0.1.4
## [21] BiocStyle_2.38.0
##
## loaded via a namespace (and not attached):
## [1] tidyselect_1.2.1 dplyr_1.1.4
## [3] blob_1.2.4 filelock_1.0.3
```

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```
## [5] bitops_1.0-9          fastmap_1.2.0
## [7] RCurl_1.98-1.17       BiocFileCache_3.0.0
## [9] GenomicAlignments_1.46.0 digest_0.6.37
## [11] lifecycle_1.0.4      KEGGREST_1.50.0
## [13] magrittr_2.0.4       compiler_4.5.1
## [15] rlang_1.1.6          sass_0.4.10
## [17] tools_4.5.1          yaml_2.3.10
## [19] rtracklayer_1.70.0   S4Arrays_1.10.0
## [21] bit_4.6.0            curl_7.0.0
## [23] DelayedArray_0.36.0  abind_1.4-8
## [25] BiocParallel_1.44.0  withr_3.0.2
## [27] purrr_1.1.0          grid_4.5.1
## [29] preprocessCore_1.72.0 SummarizedExperiment_1.40.0
## [31] cli_3.6.5            rmarkdown_2.30
## [33] crayon_1.5.3         httr_1.4.7
## [35] rjson_0.2.23         DBI_1.2.3
## [37] cachem_1.1.0         affy_1.88.0
## [39] parallel_4.5.1       BiocManager_1.30.26
## [41] restfulr_0.0.16     matrixStats_1.5.0
## [43] vctrs_0.6.5         Matrix_1.7-4
## [45] jsonlite_2.0.0       bookdown_0.45
## [47] bit64_4.6.0-1       RBGL_1.86.0
## [49] jquerylib_0.1.4     affyio_1.80.0
## [51] glue_1.8.0          codetools_0.2-20
## [53] BiocIO_1.20.0       tibble_3.3.0
## [55] pillar_1.11.1       rappdirs_0.3.3
## [57] htmltools_0.5.8.1   graph_1.88.0
## [59] httr2_1.2.1         R6_2.6.1
## [61] dbplyr_2.5.1        evaluate_1.0.5
## [63] lattice_0.22-7      highr_0.11
## [65] png_0.1-8           Rsamtools_2.26.0
## [67] cigarillo_1.0.0     memoise_2.0.1
## [69] bslib_0.9.0         SparseArray_1.10.0
## [71] xfun_0.53           MatrixGenerics_1.22.0
## [73] pkgconfig_2.0.3
```