

# illuminaRatv1.db

February 3, 2010

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illuminaRatv1ACCNUM

*Map Manufacturer identifiers to Accession Numbers*

---

## Description

illuminaRatv1ACCNUM is an R object that contains mappings between a manufacturer's identifiers and manufacturers accessions.

## Details

For chip packages such as this, the ACCNUM mapping comes directly from the manufacturer. This is different from other mappings which are mapped onto the probes via an Entrez Gene identifier.

Each manufacturer identifier maps to a vector containing a GenBank accession number.

Mappings were based on data provided by: Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

## Examples

```
x <- illuminaRatv1ACCNUM
# Get the probe identifiers that are mapped to an ACCNUM
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0) {
  # Get the ACCNUM for the first five probes
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```

---

```
illuminaRatv1ALIAS2PROBE
```

*Map between Common Gene Symbol Identifiers and Manufacturer Identifiers*

---

### Description

illuminaRatv1ALIAS is an R object that provides mappings between common gene symbol identifiers and manufacturer identifiers.

### Details

Each gene symbol is mapped to a named vector of manufacturer identifiers. The name represents the gene symbol and the vector contains all manufacturer identifiers that are found for that symbol. An NA is reported for any gene symbol that cannot be mapped to any manufacturer identifiers.

This mapping includes ALL gene symbols including those which are already listed in the SYMBOL map. The SYMBOL map is meant to only list official gene symbols, while the ALIAS maps are meant to store all used symbols.

Mappings were based on data provided by: Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

### Examples

```
# Convert the object to a list
xx <- as.list(illuminaRatv1ALIAS2PROBE)
if(length(xx) > 0){
  # Get the probe identifiers for the first two aliases
  xx[1:2]
  # Get the first one
  xx[[1]]
}
```

---

```
illuminaRatv1.db Bioconductor annotation data package
```

---

### Description

Welcome to the illuminaRatv1.db annotation Package. The purpose of this package is to provide detailed information about the illuminaRatv1 platform. This package is updated biannually.

You can learn what objects this package supports with the following command:

```
ls("package:illuminaRatv1.db")
```

Each of these objects has their own manual page detailing where relevant data was obtained along with some examples of how to use it.

### Examples

```
ls("package:illuminaRatv1.db")
```

---

```
illuminaRatv1CHRENGTHS
```

*A named vector for the length of each of the chromosomes*

---

### Description

illuminaRatv1CHRENGTHS provides the length measured in base pairs for each of the chromosomes.

### Details

This is a named vector with chromosome numbers as the names and the corresponding lengths for chromosomes as the values.

Total lengths of chromosomes were derived by calculating the number of base pairs on the sequence string for each chromosome.

### Examples

```
tt <- illuminaRatv1CHRENGTHS
# Length of chromosome 1
tt["1"]
```

---

```
illuminaRatv1CHRLOC
```

*Map Manufacturer IDs to Chromosomal Location*

---

### Description

illuminaRatv1CHRLOC is an R object that maps manufacturer identifiers to the starting position of the gene. The position of a gene is measured as the number of base pairs.

The CHRLOCEND mapping is the same as the CHRLOC mapping except that it specifies the ending base of a gene instead of the start.

### Details

Each manufacturer identifier maps to a named vector of chromosomal locations, where the name indicates the chromosome. Due to inconsistencies that may exist at the time the object was built, these vectors may contain more than one chromosome and/or location. If the chromosomal location is unknown, the vector will contain an NA.

Chromosomal locations on both the sense and antisense strands are measured as the number of base pairs from the p (5' end of the sense strand) to q (3' end of the sense strand) arms. Chromosomal locations on the antisense strand have a leading "-" sign (e. g. -1234567).

Since some genes have multiple start sites, this field can map to multiple locations.

Mappings were based on data provided by: UCSC Genome Bioinformatics (Rattus norvegicus) [ftp://hgdownload.cse.ucsc.edu/goldenPath/currentGenomes/Rattus\\\_norvegicus](ftp://hgdownload.cse.ucsc.edu/goldenPath/currentGenomes/Rattus\_norvegicus) With a date stamp from the source of: 2009-Jul5

## Examples

```
x <- illuminaRatv1CHRLOC
# Get the probe identifiers that are mapped to chromosome locations
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0) {
  # Get the CHRLOC for the first five probes
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```

---

illuminaRatv1CHR    *Map Manufacturer IDs to Chromosomes*

---

## Description

illuminaRatv1CHR is an R object that provides mappings between a manufacturer identifier and the chromosome that contains the gene of interest.

## Details

Each manufacturer identifier maps to a vector of chromosomes. Due to inconsistencies that may exist at the time the object was built, the vector may contain more than one chromosome (e.g., the identifier may map to more than one chromosome). If the chromosomal location is unknown, the vector will contain an NA.

Mappings were based on data provided by: Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

## Examples

```
x <- illuminaRatv1CHR
# Get the probe identifiers that are mapped to a chromosome
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0) {
  # Get the CHR for the first five probes
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```

---

`illuminaRatv1_dbconn`*Collect information about the package annotation DB*

---

## Description

Some convenience functions for getting a connection object to (or collecting information about) the package annotation DB.

## Usage

```
illuminaRatv1_dbconn()
illuminaRatv1_dbfile()
illuminaRatv1_dbschema(file="", show.indices=FALSE)
illuminaRatv1_dbInfo()
```

## Arguments

`file` A connection, or a character string naming the file to print to (see the `file` argument of the `cat` function for the details).

`show.indices` The CREATE INDEX statements are not shown by default. Use `show.indices=TRUE` to get them.

## Details

`illuminaRatv1_dbconn` returns a connection object to the package annotation DB. **IMPORTANT: Don't call `dbDisconnect` on the connection object returned by `illuminaRatv1_dbconn` or you will break all the `AnnDbObj` objects defined in this package!**

`illuminaRatv1_dbfile` returns the path (character string) to the package annotation DB (this is an SQLite file).

`illuminaRatv1_dbschema` prints the schema definition of the package annotation DB.

`illuminaRatv1_dbInfo` prints other information about the package annotation DB.

## Value

`illuminaRatv1_dbconn`: a `DBIConnection` object representing an open connection to the package annotation DB.

`illuminaRatv1_dbfile`: a character string with the path to the package annotation DB.

`illuminaRatv1_dbschema`: none (invisible NULL).

`illuminaRatv1_dbInfo`: none (invisible NULL).

## See Also

[dbGetQuery](#), [dbConnect](#), [dbconn](#), [dbfile](#), [dbschema](#), [dbInfo](#)

**Examples**

```
## Count the number of rows in the "probes" table:
dbGetQuery(illuminaRatv1_dbconn(), "SELECT COUNT(*) FROM probes")

## The connection object returned by illuminaRatv1_dbconn() was
## created with:
dbConnect(SQLite(), dbname=illuminaRatv1_dbfile(), cache_size=64000,
synchronous=0)

illuminaRatv1_dbschema()

illuminaRatv1_dbInfo()
```

---

```
illuminaRatv1ENSEMBL
```

*Map Ensembl gene accession numbers with Entrez Gene identifiers*

---

**Description**

illuminaRatv1ENSEMBL is an R object that contains mappings between Entrez Gene identifiers and Ensembl gene accession numbers.

**Details**

This object is a simple mapping of Entrez Gene identifiers <http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=gene> to Ensembl gene Accession Numbers.

Mappings were based on data provided by BOTH of these sources: <http://www.ensembl.org/biomart/martview/> <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA>

This mapping is a combination of NCBI to ensembl IDs from BOTH NCBI and ensembl. Users who wish to only use mappings from NCBI are encouraged to see the ncbi2ensembl table in the appropriate organism package. Users who wish to only use mappings from ensembl are encouraged to see the ensembl2ncbi table which is also found in the appropriate organism packages. These mappings are based upon the ensembl table which contains data from BOTH of these sources in an effort to maximize the chances that you will find a match.

Mappings were based on data provided by: Ensembl [ftp://ftp.ensembl.org/pub/current\\_fasta](ftp://ftp.ensembl.org/pub/current_fasta) With a date stamp from the source of: 2009-Mar6 and Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

**Examples**

```
x <- illuminaRatv1ENSEMBL
# Get the entrez gene IDs that are mapped to an Ensembl ID
mapped_genes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_genes])
if(length(xx) > 0) {
  # Get the Ensembl gene IDs for the first five genes
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```

```
#For the reverse map ENSEMBL2PROBE:
# Convert to a list
xx <- as.list(illuminaRatv1ENSEMBL2PROBE)
if(length(xx) > 0){
  # Gets the entrez gene IDs for the first five Ensembl IDs
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```

---

illuminaRatv1ENTREZID

*Map between Manufacturer Identifiers and Entrez Gene*

---

## Description

illuminaRatv1ENTREZID is an R object that provides mappings between manufacturer identifiers and Entrez Gene identifiers.

## Details

Each manufacturer identifier is mapped to a vector of Entrez Gene identifiers. An NA is assigned to those manufacturer identifiers that can not be mapped to an Entrez Gene identifier at this time.

If a given manufacturer identifier can be mapped to different Entrez Gene identifiers from various sources, we attempt to select the common identifiers. If a concensus cannot be determined, we select the smallest identifier.

Mappings were based on data provided by: Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

## References

<http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=gene>

## Examples

```
x <- illuminaRatv1ENTREZID
# Get the probe identifiers that are mapped to an ENTREZ Gene ID
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0) {
  # Get the ENTREZID for the first five probes
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```

---

illuminaRatv1ENZYME2PROBE

*Map between Enzyme Commission Numbers and Manufacturer Identifiers*

---

## Description

illuminaRatv1ENZYME2PROBE is an R object that maps Enzyme Commission (EC) numbers to manufacturer identifiers.

## Details

Each EC number maps to a named vector containing all of the manufacturer identifiers that correspond to the gene that produces that enzyme. The name of the vector corresponds to the EC number.

Enzyme Commission numbers are assigned by the Nomenclature Committee of the International Union of Biochemistry and Molecular Biology <http://www.chem.qmw.ac.uk/iubmb/enzyme/> to allow enzymes to be identified.

An Enzyme Commission number is of the format EC x.y.z.w, where x, y, z, and w are numeric numbers. In illuminaRatv1ENZYME2PROBE, EC is dropped from the Enzyme Commission numbers.

Enzyme Commission numbers have corresponding names that describe the functions of enzymes in such a way that EC x is a more general description than EC x.y that in turn is a more general description than EC x.y.z. The top level EC numbers and names are listed below:

EC 1 oxidoreductases

EC 2 transferases

EC 3 hydrolases

EC 4 lyases

EC 5 isomerases

EC 6 ligases

The EC name for a given EC number can be viewed at <http://www.chem.qmul.ac.uk/iupac/jcfn/index.html#6>

Mappings were based on data provided by: KEGG GENOME <ftp://ftp.genome.jp/pub/kegg/genomes>  
With a date stamp from the source of: 2009-Sep16

## References

<ftp://ftp.genome.ad.jp/pub/kegg/pathways>

## Examples

```
# Convert to a list
xx <- as.list(illuminaRatv1ENZYME2PROBE)
if(length(xx) > 0){
  # Get the probe identifiers for the first five enzyme
  #commission numbers
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```



---

`illuminaRatv1ENZYME`*Map between Manufacturer IDs and Enzyme Commission (EC) Numbers*

---

## Description

`illuminaRatv1ENZYME` is an R object that provides mappings between manufacturer identifiers and EC numbers.

## Details

Each manufacturer identifier maps to a named vector containing the EC number that corresponds to the enzyme produced by that gene. The name corresponds to the manufacturer identifier. If this information is unknown, the vector will contain an NA.

Enzyme Commission numbers are assigned by the Nomenclature Committee of the International Union of Biochemistry and Molecular Biology <http://www.chem.qmw.ac.uk/iubmb/enzyme/> to allow enzymes to be identified.

An Enzyme Commission number is of the format EC x.y.z.w, where x, y, z, and w are numeric numbers. In `illuminaRatv1ENZYME2PROBE`, EC is dropped from the Enzyme Commission numbers.

Enzyme Commission numbers have corresponding names that describe the functions of enzymes in such a way that EC x is a more general description than EC x.y that in turn is a more general description than EC x.y.z. The top level EC numbers and names are listed below:

EC 1 oxidoreductases

EC 2 transferases

EC 3 hydrolases

EC 4 lyases

EC 5 isomerases

EC 6 ligases

The EC name for a given EC number can be viewed at <http://www.chem.qmul.ac.uk/iupac/jcbrn/index.html#6>

Mappings between probe identifiers and enzyme identifiers were obtained using files provided by: KEGG GENOME <ftp://ftp.genome.jp/pub/kegg/genomes> With a date stamp from the source of: 2009-Sep16

## References

<ftp://ftp.genome.ad.jp/pub/kegg/pathways>

## Examples

```
x <- illuminaRatv1ENZYME
# Get the probe identifiers that are mapped to an EC number
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0) {
  # Get the ENZYME for the first five probes
```

```

xx[1:5]
# Get the first one
xx[[1]]
}

```

---

```
illuminaRatv1GENENAME
```

*Map between Manufacturer IDs and Genes*

---

### Description

illuminaRatv1GENENAME is an R object that maps manufacturer identifiers to the corresponding gene name.

### Details

Each manufacturer identifier maps to a named vector containing the gene name. The vector name corresponds to the manufacturer identifier. If the gene name is unknown, the vector will contain an NA.

Gene names currently include both the official (validated by a nomenclature committee) and preferred names (interim selected for display) for genes. Efforts are being made to differentiate the two by adding a name to the vector.

Mappings were based on data provided by: Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

### Examples

```

x <- illuminaRatv1GENENAME
# Get the probe identifiers that are mapped to a gene name
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0) {
  # Get the GENENAME for the first five probes
  xx[1:5]
  # Get the first one
  xx[[1]]
}

```

---

```
illuminaRatv1GO2ALLPROBES
```

*Map between Gene Ontology (GO) Identifiers and all Manufacturer Identifiers in the subtree*

---

### Description

illuminaRatv1GO2ALLPROBES is an R object that provides mappings between a given GO identifier and all manufacturer identifiers annotated at that GO term or one of its children in the GO ontology.

## Details

GO consists of three ontologies—molecular function (MF), biological process (BP), and cellular component (CC). All ontologies are structured as directed acyclic graphs (DAGs). Each node in each DAG (tree) is a GO term (id) associated with a named vector of manufacturer identifiers. The name associated with each manufacturer id corresponds to the evidence code for that GO identifier. This object `illuminaRatv1GO2ALLPROBES` maps between a given GO identifier and all manufacturer identifiers annotated at that GO term or one of its children in the GO ontology.

The evidence code indicates what kind of evidence supports the association between the GO and Entrez Gene identifiers. Evidence codes currently in use include:

IMP - inferred from mutant phenotype  
 IGI - inferred from genetic interaction  
 IPI - inferred from physical interaction  
 ISS - inferred from sequence similarity  
 IDA - inferred from direct assay  
 IEP - inferred from expression pattern  
 IEA - inferred from electronic annotation  
 TAS - traceable author statement  
 NAS - non-traceable author statement  
 ND - no biological data available  
 IC - inferred by curator

A GO identifier may be mapped to the same manufacturer identifier more than once but the evidence code can be different. Mappings between Gene Ontology identifiers and Gene Ontology terms and other information are available in a separate data package named `GO`.

Mappings were based on data provided by:

Gene Ontology <ftp://ftp.geneontology.org/pub/go/godatabase/archive/latest-lite/> With a date stamp from the source of: 20090830 and Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

## References

<ftp://ftp.ncbi.nlm.nih.gov/gene/DATA/>

## Examples

```
# Convert to a list
xx <- as.list(illuminaRatv1GO2ALLPROBES)
if(length(xx) > 0){
  # Get the probe identifiers for the top 2nd and 3rd GO identifiers
  goids <- xx[2:3]
  # Get all the probe identifiers for the first element of goids
  goids[[1]]
  # Evidence code for the mappings
  names(goids[[1]])
}
```

---

`illuminaRatv1GO2PROBE`*Map between Gene Ontology (GO) and Manufacturer Identifiers*

---

**Description**

`illuminaRatv1GO2PROBE` is an R object that provides mappings between GO identifiers and manufacturer identifiers.

**Details**

Each GO term maps to a named vector of manufacturer identifiers. The name associated with each manufacturer identifier corresponds to the evidence code for that GO identifier. The evidence code indicates what kind of evidence supports the association between the GO and Entrez Gene identifiers. Evidence codes currently in use include:

IMP - inferred from mutant phenotype

IGI - inferred from genetic interaction

IPI - inferred from physical interaction

ISS - inferred from sequence similarity

IDA - inferred from direct assay

IEP - inferred from expression pattern

IEA - inferred from electronic annotation

TAS - traceable author statement

NAS - non-traceable author statement

ND - no biological data available

IC - inferred by curator

A GO identifier may be mapped to the same probe identifier more than once but the evidence code can be different. Mappings between Gene Ontology identifiers and Gene Ontology terms and other information are available in a separate data package named `GO`.

Mappings were based on data provided by:

Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

**References**

<ftp://ftp.ncbi.nlm.nih.gov/gene/DATA/>

**Examples**

```
# Convert to a list
xx <- as.list(illuminaRatv1GO2PROBE)
if(length(xx) > 0){
  # Get the probe identifiers for the top 2nd and 3rd GO identifiers
  goids <- xx[2:3]
  # Get the probe identifiers for the first element of goids
  goids[[1]]
  # Evidence code for the mappings
  names(goids[[1]])
}
```

---

illuminaRatv1GO      *Map between Manufacturer IDs and Gene Ontology (GO)*

---

## Description

illuminaRatv1GO is an R object that provides mappings between manufacturer identifiers and the GO identifiers that they are directly associated with.

## Details

Each Entrez Gene identifier is mapped to a list of lists. The names on the outer list are GO identifiers. Each inner list consists of three named elements: GOID, Ontology, and Evidence.

The GOID element matches the GO identifier named in the outer list and is included for convenience when processing the data using 'lapply'.

The Ontology element indicates which of the three Gene Ontology categories this identifier belongs to. The categories are biological process (BP), cellular component (CC), and molecular function (MF).

The Evidence element contains a code indicating what kind of evidence supports the association of the GO identifier to the Entrez Gene id. The evidence codes in use include:

IMP: inferred from mutant phenotype

IGI: inferred from genetic interaction

IPI: inferred from physical interaction

ISS: inferred from sequence similarity

IDA: inferred from direct assay

IEP: inferred from expression pattern

IEA: inferred from electronic annotation

TAS: traceable author statement

NAS: non-traceable author statement

ND: no biological data available

IC: inferred by curator

Mappings between probe identifiers and GO information were obtained through their mappings to Entrez Gene identifiers. NAs are assigned to probe identifiers that can not be mapped to any Gene Ontology information. Mappings between Gene Ontology identifiers and Gene Ontology terms and other information are available in a separate data package named GO.

Mappings were based on data provided by: Gene Ontology <ftp://ftp.geneontology.org/pub/go/godatabase/archive/latest-lite/> With a date stamp from the source of: 20090830

## Examples

```
x <- illuminaRatv1GO
# Get the probe identifiers that are mapped to a GO ID
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0) {
  # Try the first one
```

```

got <- xx[[1]]
got[[1]][["GOID"]]
got[[1]][["Ontology"]]
got[[1]][["Evidence"]]

}

```

---

```
illuminaRatv1MAPCOUNTS
```

*Number of mapped keys for the maps in package illuminaRatv1.db*

---

### Description

illuminaRatv1MAPCOUNTS provides the "map count" (i.e. the count of mapped keys) for each map in package illuminaRatv1.db.

### Details

This "map count" information is precalculated and stored in the package annotation DB. This allows some quality control and is used by the [checkMAPCOUNTS](#) function defined in AnnotationDbi to compare and validate different methods (like `count.mappedkeys(x)` or `sum(!is.na(as.list(x)))`) for getting the "map count" of a given map.

### See Also

[mappedkeys](#), [count.mappedkeys](#), [checkMAPCOUNTS](#)

### Examples

```

illuminaRatv1MAPCOUNTS
mapnames <- names(illuminaRatv1MAPCOUNTS)
illuminaRatv1MAPCOUNTS[mapnames[1]]
x <- get(mapnames[1])
sum(!is.na(as.list(x)))
count.mappedkeys(x) # much faster!

## Check the "map count" of all the maps in package illuminaRatv1.db
checkMAPCOUNTS("illuminaRatv1.db")

```

---

```
illuminaRatv1MAP
```

*Map between Manufacturer Identifiers and cytogenetic maps/bands*

---

### Description

illuminaRatv1MAP is an R object that provides mappings between manufacturer identifiers and cytoband locations.

**Details**

Each manufacturer identifier is mapped to a vector of cytoband locations. The vector length may be one or longer, if there are multiple reported chromosomal locations for a given gene. An NA is reported for any manufacturer identifiers that cannot be mapped to a cytoband at this time.

The physical location of each band on a chromosome can be obtained from another object named "organism"CYTOLOC in a separate data package for human(humanCHRLOC), mouse(mouseCHRLOC), and rat(ratCHRLOC).

Mappings were based on data provided by: Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

**References**

<http://www.ncbi.nlm.nih.gov>

**Examples**

```
x <- illuminaRatv1MAP
# Get the probe identifiers that are mapped to any cytoband
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0) {
  # Get the MAP for the first five probes
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```

---

illuminaRatv1ORGANISM

*The Organism information for illuminaRatv1*

---

**Description**

illuminaRatv1ORGANISM is an R object that contains a single item: a character string that names the organism for which illuminaRatv1 was built. illuminaRatv1ORGPKG is an R object that contains a character vector with the name of the organism package that a chip package depends on for its gene-centric annotation.

**Details**

Although the package name is suggestive of the organism for which it was built, illuminaRatv1ORGANISM provides a simple way to programmatically extract the organism name. illuminaRatv1ORGPKG provides a simple way to programmatically extract the name of the parent organism package. The parent organism package is a strict dependency for chip packages as this is where the gene centric information is ultimately extracted from. The full package name will always be this string plus the extension ".db". But most programatic acces will not require this extension, so its more convenient to leave it out.

**Examples**

```
illuminaRatv1ORGANISM
illuminaRatv1ORGPKG
```

---

```
illuminaRatv1PATH2PROBE
```

*Map between Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway identifiers and Manufacturer Identifiers*

---

### Description

illuminaRatv1PATH2PROBE is an R object that provides mappings between KEGG identifiers and manufacturer identifiers.

### Details

Each KEGG identifier is mapped to a named vector of manufacturer identifiers. The name represents the KEGG identifier and the vector contains all manufacturer identifiers that are found in that particular pathway. An NA is reported for any KEGG identifier that cannot be mapped to any manufacturer identifiers.

Pathway name for a given pathway identifier can be obtained using the KEGG data package that can either be built using AnnBuilder or downloaded from Bioconductor <http://www.bioconductor.org>.

Graphic presentations of pathways are searchable at <http://www.genome.ad.jp/kegg/pathway.html> using pathway identifiers as keys.

Mappings were based on data provided by: KEGG GENOME <ftp://ftp.genome.jp/pub/kegg/genomes> With a date stamp from the source of: 2009-Sep16

### References

<http://www.genome.ad.jp/kegg/>

### Examples

```
# Convert the object to a list
xx <- as.list(illuminaRatv1PATH2PROBE)
if (length(xx) > 0) {
  # Get the probe identifiers for the first two pathway identifiers
  xx[1:2]
  # Get the first one
  xx[[1]]
}
```

---

```
illuminaRatv1PATH Mappings between probe identifiers and KEGG pathway identifiers
```

---

### Description

KEGG (Kyoto Encyclopedia of Genes and Genomes) maintains pathway data for various organisms. illuminaRatv1PATH maps probe identifiers to the identifiers used by KEGG for pathways in which the genes represented by the probe identifiers are involved



## Details

Each KEGG pathway has a name and identifier. Pathway name for a given pathway identifier can be obtained using the KEGG data package that can either be built using AnnBuilder or downloaded from Bioconductor <http://www.bioconductor.org>.

Graphic presentations of pathways are searchable at url <http://www.genome.ad.jp/kegg/pathway.html> by using pathway identifiers as keys.

Mappings were based on data provided by: KEGG GENOME <ftp://ftp.genome.jp/pub/kegg/genomes> With a date stamp from the source of: 2009-Sep16

## References

<http://www.genome.ad.jp/kegg/>

## Examples

```
x <- illuminaRatv1PATH
# Get the probe identifiers that are mapped to a KEGG pathway ID
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0) {
  # Get the PATH for the first five probes
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```

---

illuminaRatv1PFAM *Map Manufacturer IDs to Pfam IDs*

---

## Description

illuminaRatv1PFAM is an R object that provides mappings between a manufacturer identifier and the associated Pfam identifiers.

## Details

Each manufacturer identifier maps to a named vector of Pfam identifiers. The name for each Pfam identifier is the IPI accession number where this Pfam identifier is found.

If the Pfam is a named NA, it means that the associated Entrez Gene id of this manufacturer identifier is found in an IPI entry of the IPI database, but there is no Pfam identifier in the entry.

If the Pfam is a non-named NA, it means that the associated Entrez Gene id of this manufacturer identifier is not found in any IPI entry of the IPI database.

Mappings were based on data provided by: The International Protein Index <ftp://ftp.ebi.ac.uk/pub/databases/IPI/current> With a date stamp from the source of: 2009-Jul29

## References

Kersey P. J., Duarte J., Williams A., Karavidopoulou Y., Birney E., Apweiler R. The International Protein Index: An integrated database for proteomics experiments. *Proteomics* 4(7): 1985-1988 (2004) <http://www.ebi.ac.uk/IPI/IPIhelp.html>

**Examples**

```
x <- illuminaRatv1PFAM
# Get the probe identifiers that are mapped to any Pfam ID
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
# randomly display 10 probes
sample(xx, 10)
```

---

illuminaRatv1PMID2PROBE

*Map between PubMed Identifiers and Manufacturer Identifiers*

---

**Description**

illuminaRatv1PMID2PROBE is an R object that provides mappings between PubMed identifiers and manufacturer identifiers.

**Details**

Each PubMed identifier is mapped to a named vector of manufacturer identifiers. The name represents the PubMed identifier and the vector contains all manufacturer identifiers that are represented by that PubMed identifier. The length of the vector may be one or longer, depending on how many manufacturer identifiers are mapped to a given PubMed identifier.

Titles, abstracts, and possibly full texts of articles can be obtained from PubMed by providing a valid PubMed identifier. The `pubmed` function of `annotate` can also be used for the same purpose

Mappings were based on data provided by: Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

**Examples**

```
# Convert the object to a list
xx <- as.list(illuminaRatv1PMID2PROBE)
if(length(xx) > 0){
  # Get the probe identifiers for the first two PubMed identifiers
  xx[1:2]
  # Get the first one
  xx[[1]]
  if(interactive() && require(annotate)){
    # Get article information as XML files for a PubMed id
    xmls <- pubmed(names(xx)[1], disp = "data")
    # View article information using a browser
    pubmed(names(xx)[1], disp = "browser")
  }
}
```

---

illuminaRatv1PMID *Map between Manufacturer Identifiers and PubMed Identifiers*

---

## Description

illuminaRatv1PMID is an R object that provides mappings between manufacturer identifiers and PubMed identifiers.

## Details

Each manufacturer identifier is mapped to a named vector of PubMed identifiers. The name associated with each vector corresponds to the manufacturer identifier. The length of the vector may be one or greater, depending on how many PubMed identifiers a given manufacturer identifier is mapped to. An NA is reported for any manufacturer identifier that cannot be mapped to a PubMed identifier.

Titles, abstracts, and possibly full texts of articles can be obtained from PubMed by providing a valid PubMed identifier. The `pubmed` function of `annotate` can also be used for the same purpose.

Mappings were based on data provided by: Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

## References

<http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=PubMed>

## Examples

```
x <- illuminaRatv1PMID
# Get the probe identifiers that are mapped to any PubMed ID
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0){
  # Get the PubMed identifiers for the first two probe identifiers
  xx[1:2]
  # Get the first one
  xx[[1]]
  if(interactive() && !is.null(xx[[1]]) && !is.na(xx[[1]])
    && require(annotate)){
    # Get article information as XML files
    xmls <- pubmed(xx[[1]], disp = "data")
    # View article information using a browser
    pubmed(xx[[1]], disp = "browser")
  }
}
```

---

`illuminaRatv1PROSITE`*Map Manufacturer IDs to PROSITE ID*

---

### Description

`illuminaRatv1PROSITE` is an R object that provides mappings between a manufacturer identifier and the associated PROSITE identifiers.

### Details

Each manufacturer identifier maps to a named vector of PROSITE identifiers. The name for each PROSITE identifier is the IPI accession number where this PROSITE identifier is found.

If the PROSITE is a named NA, it means that the associated Entrez Gene id of this manufacturer identifier is found in an IPI entry of the IPI database, but there is no PROSITE identifier in the entry.

If the PROSITE is a non-named NA, it means that the associated Entrez Gene id of this manufacturer identifier is not found in any IPI entry of the IPI database.

Mappings were based on data provided by: The International Protein Index <ftp://ftp.ebi.ac.uk/pub/databases/IPI/current>  
With a date stamp from the source of: 2009-Jul29

### References

Kersey P. J., Duarte J., Williams A., Karavidopoulou Y., Birney E., Apweiler R. The International Protein Index: An integrated database for proteomics experiments. *Proteomics* 4(7): 1985-1988 (2004) <http://www.ebi.ac.uk/IPI/IPIhelp.html>

### Examples

```
x <- illuminaRatv1PROSITE
# Get the probe identifiers that are mapped to any PROSITE ID
mapped_probes <- mappedkeys(x)
# Convert to a list
xxx <- as.list(x[mapped_probes])
# randomly display 10 probes
xxx[sample(1:length(xxx), 10)]
```

---

`illuminaRatv1REFSEQ`*Map between Manufacturer Identifiers and RefSeq Identifiers*

---

### Description

`illuminaRatv1REFSEQ` is an R object that provides mappings between manufacturer identifiers and RefSeq identifiers.



**Details**

Each manufacturer identifier is mapped to an abbreviation for the corresponding gene. An NA is reported if there is no known abbreviation for a given gene.

Symbols typically consist of 3 letters that define either a single gene (ABC) or multiple genes (ABC1, ABC2, ABC3). Gene symbols can be used as key words to query public databases such as Entrez Gene.

Mappings were based on data provided by: Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

**References**

<http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=gene>

**Examples**

```
x <- illuminaRatv1SYMBOL
# Get the probe identifiers that are mapped to a gene symbol
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0) {
  # Get the SYMBOL for the first five probes
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```

---

illuminaRatv1UNIGENE

*Map between Manufacturer Identifiers and UniGene cluster identifiers*

---

**Description**

illuminaRatv1UNIGENE is an R object that provides mappings between manufacturer identifiers and UniGene identifiers.

**Details**

Each manufacturer identifier is mapped to a UniGene identifier. An NA is reported if the manufacturer identifier cannot be mapped to UniGene at this time.

A UniGene identifier represents a cluster of sequences of a gene. Using UniGene identifiers one can query the UniGene database for information about the sequences or the Entrez Gene database for information about the genes.

Mappings were based on data provided by: Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

**References**

<http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=gene>

**Examples**

```
x <- illuminaRatv1UNIGENE
# Get the probe identifiers that are mapped to an UNIGENE ID
mapped_probes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_probes])
if(length(xx) > 0) {
  # Get the UNIGENE for the first five probes
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```

---

illuminaRatv1UNIPROT

*Map Uniprot accession numbers with Entrez Gene identifiers*

---

**Description**

illuminaRatv1UNIPROT is an R object that contains mappings between Entrez Gene identifiers and Uniprot accession numbers.

**Details**

This object is a simple mapping of Entrez Gene identifiers <http://www.ncbi.nlm.nih.gov/entrez/query.fcgi?db=gene> to Uniprot Accessions.

Mappings were based on data provided by: Entrez Gene <ftp://ftp.ncbi.nlm.nih.gov/gene/DATA> With a date stamp from the source of: 2009-Sep1

**Examples**

```
x <- illuminaRatv1UNIPROT
# Get the entrez gene IDs that are mapped to an Uniprot ID
mapped_genes <- mappedkeys(x)
# Convert to a list
xx <- as.list(x[mapped_genes])
if(length(xx) > 0) {
  # Get the Uniprot IDs for the first five genes
  xx[1:5]
  # Get the first one
  xx[[1]]
}
```

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