Genome project tables in the genomes package

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The genomes package collects genome project metadata and provides tools to track, sort, group, summarize and plot the data. The genome project tables from the National Center for Biotechnology Information (NCBI) and the Genomes On Line Database (GOLD) are the primary sources of data and include a rapidly growing collection of organisms from all domains of life (viruses, archaea, bacteria, protists, fungi, plants, and animals) plus metagenomic sequences.

Genome tables are a defined class (*genomes*) in the package and each table is a data frame where rows are genome projects and columns are the fields describing the associated metadata. At a minimum, the table should have a column listing the project name, status, and release date. A number of methods are available that operate on genome tables including print, summary, plot and update.

There are a number of ways to install this package. If you are running R version 2.11, you can use the biocLite command.

```
R> source("http://bioconductor.org/biocLite.R")
R> biocLite("genomes")
```

You can also install the package on earlier versions of R using install.packages, but this has not been tested completely.

```
R> install.packages("genomes",
    repos="http://www.bioconductor.org/packages/release/bioC")
```

Finally, since the format of online genome tables may change (and then update commands may fail), I would recommend downloading the development version for fixes in between the six month release cycle.

```
R> install.packages("genomes",
    repos="http://www.bioconductor.org/packages/devel/bioC")
```

NCBI tables

Genome tables at NCBI are downloaded from the Genome Project database. The primary tables include a list of prokaryotic projects (lproks), eukaryotic projects (leuks), and metagenomic projects (lenvs). The print methods displays the first few rows and columns of the table (either select less than seven rows or convert the object to a data.frame to print all columns). The summary function displays the download date, a count of projects by status, and a list of recent submissions. The plot method displays a cumulative plot of genomes by release date in Figure 1 (use lines to add additional tables). The update method is not illustrated below, but can be used to download the latest version of the table from NCBI.

```
R> data(lproks)
R> lproks
```

A genomes data.frame with 4174 rows and 32 columns

```
refseq_pid
                                                                       status
                                                            name
1
          30807
                                          'Nostoc azollae' 0708
                                                                    Assembly
                              Abiotrophia defectiva ATCC 49176
2
          33011
                                                                    Assembly
3
          12997
                                Acaryochloris marina MBIC11017
                                                                    Complete
4
             NA
                                  Acaryochloris sp. CCMEE 5410 In Progress
5
             NA
                                Acetivibrio cellulolyticus CD2 In Progress
4174
             NA Zymomonas mobilis subsp. pomaceae ATCC 29192 In Progress
       released ...
     2009-03-06 ...
1
2
     2009-03-17 ...
3
     2007-10-16 ...
4
           <NA> ...
5
           <NA> ...
             . . . . . .
. . .
4174
           <NA> ...
R> summary(lproks)
$`Total genomes`
[1] 4174 genome projects on May 18, 2010
$`By status`
            Total
In Progress
             1768
Assembly
              1265
Complete
              1141
```

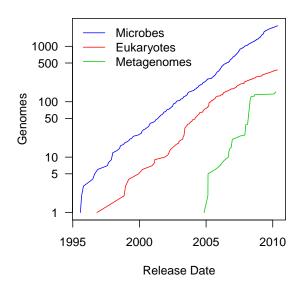


Figure 1: Cumulative plot of genome projects by release date at NCBI.

```
$`Recent submissions`
 RELEASED
                                                    STATUS
             NAME
1 2010-05-17 Arcobacter nitrofigilis DSM 7299
                                                    Complete
2 2010-05-17 Thermobispora bispora DSM 43833
                                                    Complete
3 2010-05-17 Thermosphaera aggregans DSM 11486
                                                    Complete
4 2010-05-13 Thermincola sp. JR
                                                    Complete
5 2010-05-11 Corynebacterium ammoniagenes DSM 20306 Assembly
R> plot(lproks, log = "y", las = 1)
R> data(leuks)
R> data(lenvs)
R> lines(leuks, col = "red")
R> lines(lenvs, col = "green3")
R> legend("topleft", c("Microbes", "Eukaryotes", "Metagenomes"),
     lty = 1, bty = "n", col = c("blue", "red", "green3"))
```

For microbial genome projects, the number of complete genomes doubles every 22 months and a new microbial genome is released about every other day. At least in 2008, fewer complete genomes were released than the previous year (Figure 2).

```
R> complete <- subset(lproks, status == "Complete")
R> doublingTime(complete)
days
674
```

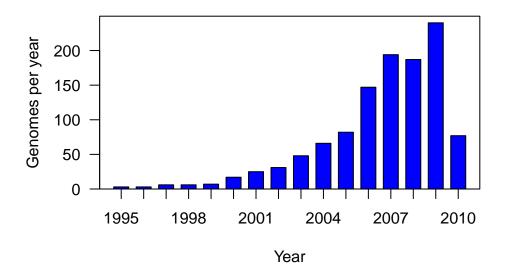


Figure 2: Number of complete microbial genomes released each year at NCBI

A number of functions are available to assist in sorting and grouping genomes. For example, the species and genus function can be used to extract the genus or species name. The table2 function formats and sorts a contingency table by counts.

R> table2(species(lproks\$name))

	Total
Escherichia coli	283
Streptococcus pneumoniae	194
Salmonella enterica	127
Staphylococcus aureus	78
Mycobacterium tuberculosis	76
Enterococcus faecalis	57
Bacillus cereus	53
Vibrio cholerae	48
Brucella melitensis	41
Helicobacter pylori	41

Because subsets of tables are often needed, the binary operator like allows pattern matching using wildcards. The plotby function below expands on the default plot method

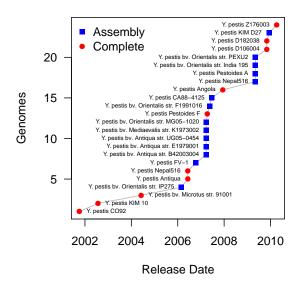


Figure 3: Cumulative plot of *Yersinia pestis* genomes by release date.

and adds the ability to plot by groups (default is status) using either labeled points or multiple lines like Figure 1. For example, the release dates of complete and draft sequences of *Yersinia pestis* are displayed in Figure 3.

```
R> yp <- subset(lproks, name %like% "Yersinia pestis*")
R> plotby(yp, labels = TRUE, cex = 0.5, lbty = "n")
```

GOLD and other tables

The Genomes Online Database (GOLD) is a comprehensive resource that collects detailed project metadata from over 7,000 genomes. There are currenlty over 100 columns in this large table with specific fields relating to the organism, host, environment, and sequencing methods. Just two of the hundreds of possible queries are illustated below. In first example, a list of endosymbiotic intracellular organisms is divided into pathogens and commensal bacteria. In the second example, the comma-separated list of phenotypes is split and a new table is created listing the GOLD identifier, name, and a single phenotype. Then genomes matching "Arsenic metabolizer" are displayed.

```
0
                                  47
                                        47
  Chlamydia
  Rickettsia
                         2
                                  18
                                        20
  Rhizobium
                        18
                                  0
                                        18
  Wolbachia
                        17
                                  0
                                        17
                                  12
                                        12
  Chlamydophila
                         0
  Buchnera
                                  0
                                        11
                        11
                                  7
  Coxiella
                         0
                                         7
                                  7
                                         7
  Ehrlichia
                         0
                         0
                                   6
                                         6
  Anaplasma
  Mesorhizobium
                         5
                                   0
                                         5
R> x <- subset(gold, phenotype != "")</pre>
R> x2 <- strsplit(x$phenotype, ", ")</pre>
R> gold2 <- as.data.frame(cbind(goldstamp = rep(x$goldstamp,
     sapply(x2, length)), name = rep(x$name, sapply(x2, length)),
     phenotype = unlist(x2))
R> table2(gold2$phenotype)
                        Total
Pathogen
                         1952
Non-Pathogen
                          249
Intracellular pathogen
                          114
                           71
Acidophile
Parasite
                           58
Probiotic
                           50
Meticillin resistant
                           44
Radiation resistant
                           37
Catalase positive
                           34
```

Commensal Pathogen Total

R> subset(gold2, phenotype %like% "Arsenic metabol*")

32

Symbiont

	goldstamp		name		phenotype
130	Gc00422	Alkalilimnicola ehrlichei	MLHE-1	Arsenic	metabolizer
133	Gc00666	Alkaliphilus oremlandii	OhILAs	Arsenic	metabolizer
303	Gi00970	Bacillus selenitireducens	MLMS-1	Arsenic	metabolizer
304	Gi00921	Bacillus selenitireducens	MLS-10	Arsenic	metabolizer
1787	Gc00526	Herminiimonas arsenicoxydans	ULPAs1	Arsenic	metabolizer
2994	Gi00788	Thiomor	nas sp.	Arsenic	metabolizer

Finally, genome data from the Human Microbiome Project is stored in the hmp dataset and includes additional information such as the primary body site occupied by a sequenced organism.