How to manage data

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Lecturer in Epigenetic Epidemiology





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- e.g. Given data for 100,000 individuals, calculate mean income by county.

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Here is one way to speed it up.

Step 1. Peek at the first few rows to determine the data class of each column:

```
CHR POS SNP
1 721290 rs12565286:G:C
1 723891 rs2977670:G:C
1 752566 rs3094315:G:A
                        CHR
                       ###
```

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                                 ## 1
## 2
```

Step 2. Use fread() from the data.table library to load the file:

```
classes <- c(chr="character", pos="integer", snp="character")
dat <- fread("big-massive-file.txt",
    header=T, sep=" ",</pre>
                                                                                                                            colClasses=classes)
library(data.table)
```

Some spreadsheets are just too large to load, e.g. files larger than 1/2 Gb.

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One solution is to load parts of the table as you need them.

```
dat <- read.table.ffdf("big-massive-file.txt", header=T, sep=" ")</pre>
```

You can now access dat just like a data frame.

```
## retrieve the first three rows
## retreive the column named "CHR"
dat[1:3,]
dat$CHR
```

However, you might notice that retrievals are much slower than a normal data frame.

R can handle many other file formats.

The following functions all read the file and return a data frame:

file format R package function(s)

ta()	ta13()	ta())ss())ss()	read xls() and read xlsx()
read.dta()	read.d	read_dta()	read.spss()	read_spss()	read x
foreign	readstata13 read.dta13()	haven		haven	readxl
STATA (.dat) foreign			SPSS (.sav) foreign		Excel

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	readstata13	readstata13 read.dta13()
	haven	read_dta()
SPSS (.sav) foreign	foreign	read.spss()
	haven	read_spss()
Excel	readxl	read_xls() and read_xlsx()

Note

- This is not a complete list. See the foreign and haven packages for other formats.
- read.dta13() is specifically for STATA v13 files.
- read_dta() loads variable comments, read.dta13() does not

Save data to a file: csv

The most common way in R to save a data frame is to use write.csv() or write.table().

```
write.csv(dat, row.names=F, quote=F)
write.table(dat, row.names=F, quote=F, sep=",")
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```

The newer function fwrite() is 30-60 times faster but otherwise identical.

```
library(data.table)
fwrite(dat, row.names=F, quote=F, sep=",")
```

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If a dataset will only be used by R, then it is most efficient to use the RData/RDS

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RData

The save() function saves variables to to an RData file.

```
save(dat, dat.norm, ret, file="datasets
```

The Load() function loads these variables into R.

```
load("datasets.rda", verbose=T)
## Loading objects:
## dat
## data.norm
## ret
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```

RDS files each save only one object.

```
saveRDS(dat, file="dataset.rds")
```

That single object can be loaded into R with any given variable name.

```
mydat <- readRDS("dataset.rds")</pre>
```

Transform a variable

Suppose we have a numerical variable x.

We can convert it to 3 categories corresponding to three ranges of values.

```
x.3 <- rep('null', length(x))
x.3[x < -0.5] <- 'neg'
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```
x.3 <- factor(x.3, levels=c("neg", "null", "pos"), ordered=T)
```

We could combine the 'null' and 'pos' categories and replace call the resulting category 'nonneg'.

```
x.2 <- as.character(x.3)
x.2[x.3=="null" | x.3=="pos"] <- "nonneg"
x.2 <- factor(x.2, levels=c("neg", "nonneg"), ordered=T)</pre>
```

Align datasets

For a population sample, we have gene expression levels and phenotype information.

- gene.dat is a matrix rows=genes and columns=samples
- pheno.dat is a data frame with rows=samples and columns=variables

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To test associations between gene expression and phenotypes, we need to make sure that the samples are in the same order.

```
idx <- match(pheno.dat$id, colnames(gene.dat))
gene.dat <- gene.dat[,idx]</pre>
```

Now we can test associations.

```
fit <- lm(gene.dat["MYC",] ~ ., data=pheno.dat)</pre>
```

Summarize a variable by group

Suppose we have data from an experiment in which guinea pigs were each given one of two supplements at one of 3 different doses. We want to know which supplement/dose maximizes tooth growth.

```
> head(ToothGrowth)
```

We calculate the mean growth by supplement/dose using aggregate().



"The tidyverse is an opinionated collection of R packages designed for data science. All packages share an underlying design philosophy, grammar, and data structures."

Some of the more popular packages are:

- **dplyr** (https://dplyr.tidyverse.org/) is for manipulating data.
- **ggplot2** (https://ggplot2.tidyverse.org/) is

for creating plots.