

# Efficient *R* Programming

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17-18 February, 2011

# Motivation

## Challenges

- ▶ Long calculations: bootstrap, MCMC, ....
- ▶ Big data: genome-wide association studies, re-sequencing, ....
- ▶ Long  $\times$  big: ...

## Solutions

- ▶ Avoid *R* programming pitfalls – *very* significant benefits
- ▶ Large data management
- ▶ Parallel evaluation, especially ‘embarrassingly parallel’ (not discussed in this course)

# Programming pitfalls: easy solutions

- ▶ Input only required data

```
> colClasses <-  
+   c("NULL", "integer", "numeric", "NULL")  
> df <- read.table("myfile", colClasses=colClasses)
```

- ▶ Preallocate-and-fill, not copy-and-append

```
> result <- numeric(nrow(df))  
> for (i in seq_len(nrow(df)))  
+   result[[i]] <- some_calc(df[i,])
```

- ▶ Vectorized calculations, not iteration

```
> x <- runif(100000); x2 <- x^2  
> m <- matrix(x2, nrow=1000); y <- rowSums(m)
```

- ▶ Avoid unnecessary character creation operations, e.g.,  
USE.NAMES=FALSE in sapply, use.names=FALSE in unlist.

## Programming pitfalls: moderate solutions

- ▶ Use appropriate functions, often from specialized packages.  

```
> library(limma) # microarray linear models  
> fit <- lmFit(eSet, design)
```
- ▶ Identify appropriate algorithms, e.g., %in% is  $O(N)$ , whereas naive might be  $O(N^2)$   

```
> x <- 1:100; s <- sample(x, 10)  
> inS <- x %in% s
```
- ▶ Use C or Fortran code. Requires knowledge of other programming languages, and how to integrate these into R

# Measuring performance: timing

- ▶ Use `system.time` to measure total evaluation time
  - ▶ `gcFirst=TRUE` for 'garbage collection'
- ▶ Use `replicate` to average over invocations

```
> m <- matrix(runif(200000), 20000)
> replicate(5, system.time(apply(m, 1, sum))[[1]])
```

```
[1] 0.23 0.28 0.29 0.25 0.25
```

```
> replicate(5, system.time(rowSums(m))[[1]])
```

```
[1] 0 0 0 0 0
```

- ▶ Cautionary tale: <http://tinyurl.com/29bd6xv>

## Measuring performance: comparison

- ▶ `identical` and `all.equal` ensure that 'optimizations' produce correct results!

```
> res1 <- apply(m, 1, sum)
```

```
> res2 <- rowSums(m)
```

```
> identical(res1, res2)
```

```
[1] TRUE
```

```
> identical(c(1, -1), c(x=1, y=-1))
```

```
[1] FALSE
```

```
> all.equal(c(1, -1), c(x=1, y=-1),  
+          check.attributes=FALSE)
```

```
[1] TRUE
```

## Measuring execution time: Rprof

```
> tmpf = tempfile()
> Rprof(tmpf)
> res1 <- apply(m, 1, sum)
> Rprof(NULL); summaryRprof(tmpf)
```

\$by.self

	self.time	self.pct	total.time	total.pct
"apply"	0.16	80	0.20	100
"FUN"	0.02	10	0.02	10
"lapply"	0.02	10	0.02	10
"unlist"	0.00	0	0.02	10

\$by.total

	total.time	total.pct	self.time	self.pct
"apply"	0.20	100	0.16	80
"FUN"	0.02	10	0.02	10
"lapply"	0.02	10	0.02	10
"unlist"	0.02	10	0.00	0

## Measuring memory use: `tracemem`

- ▶ Enable memory profiling

```
> ~/src/R-devel/configure --help  
> ~/src/R-devel/configure --enable-memory-profiling  
> make -j
```

- ▶ Copy-on-change semantics

```
> x <- 1:10; tracemem(x)  
[1] "<0x1b1a8f8>"  
> y <- x          # no change, so no copy  
> x[1] <- 2L      # x, y now differ, so copy  
tracemem[0x1b1a8f8 -> 0x1b1a8a0]:
```



## Measuring memory use: `tracemem`

### ► Copying in *R* functions

```
> l <- list(a=1:10, b=1:10); tracemem(l$a)
[1] "<0x1131ce0>"
> df0 <- as.data.frame(l)
tracemem[0x1131ce0 -> 0x1131bd8]: eval as.data.frame.list a
tracemem[0x1131bd8 -> 0x1131a20]: data.frame eval eval as.o
tracemem[0x1131a20 -> 0x11318c0]: as.data.frame.integer as.
> df1 <- data.frame(a=l$a, b=l$b)
tracemem[0x1131ce0 -> 0x11332c0]: data.frame
tracemem[0x11332c0 -> 0x1133160]: as.data.frame.integer as.
> identical(df0, df1)
[1] TRUE
```

## Case study: GWAS

- ▶ Subset of genome-wide association study data
- ▶ Manage sample and SNP annotations in SQL
- ▶ Manage SNP genotypes in netCDF
- ▶ Coordinate access using S4 classes
- ▶ Perform calculations (sliding window composite linkage disequilibrium) in C.
- ▶ Organize as *R* package *StudentGWAS*