The R package {bigstatsr}: memory- and computation-efficient tools for big matrices stored on disk

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Slides: https://privefl.github.io/R-presentation/bigstatsr.html

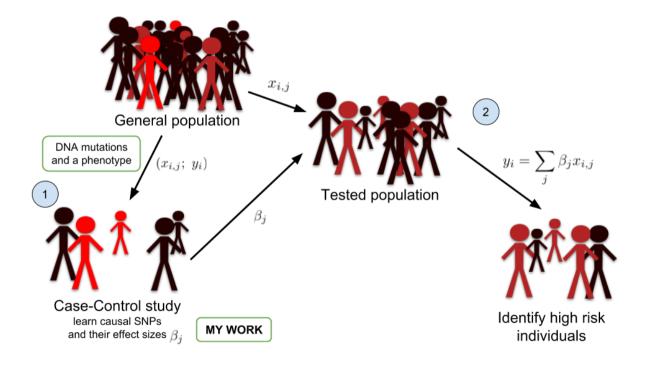
Installation: devtools::install_github("privefl/bigstatsr")

Motivation

My thesis work

I'm a PhD Student (2016-2019) in **Predictive Human Genetics**.

 $\overline{ ext{Disease} \sim ext{DNA mutations} + \cdots}$



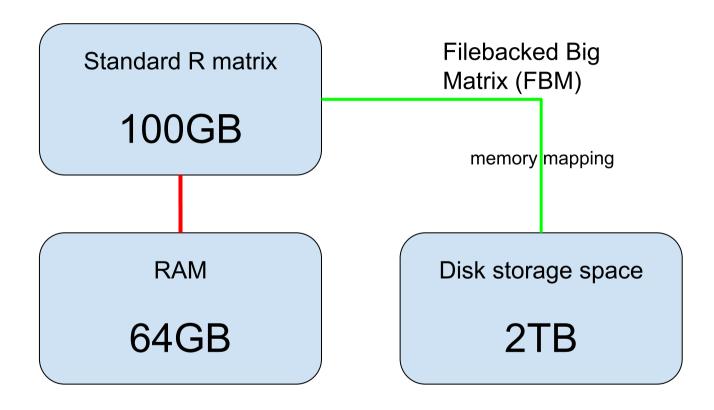
Very large genotype matrices

- previously: 15K x 280K, celiac disease (~30GB)
- currently: 500K x 500K, UK Biobank (~2TB)



But I still want to use **Q**...

The solution I found



Format FBM is very similar to format filebacked.big.matrix from package {bigmemory} (details in this vignette).

Simple accessors

Similar accessor as R matrices

```
X <- FBM(2, 5, init = 1:10, backingfile = "test")</pre>
X$backingfile
## [1] "/home/privef/Bureau/R-presentation/test.bk"
X[, 1] ## ok
## [1] 1 2
X[1, ] ## bad
## [1] 1 3 5 7 9
X[] ## super bad
## [,1] [,2] [,3] [,4] [,5]
## [1,] 1 3 5 7 9
## [2,] 2 4 6 8 10
```

Similar accessor as R matrices

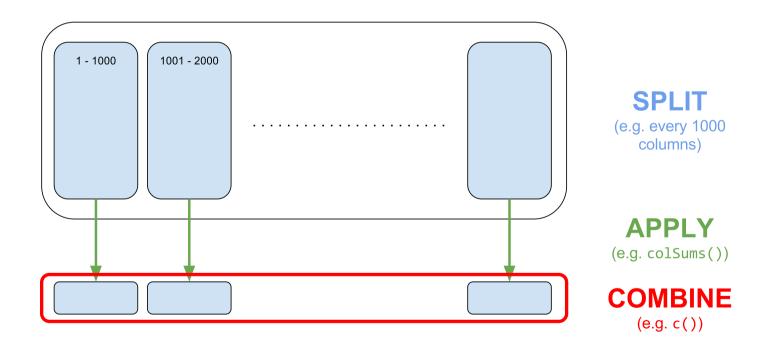
```
colSums(X[]) ## super bad
```

[1] 3 7 11 15 19



Split-(par)Apply-Combine Strategy

Apply standard R functions to big matrices (in parallel)



Implemented in big_apply().

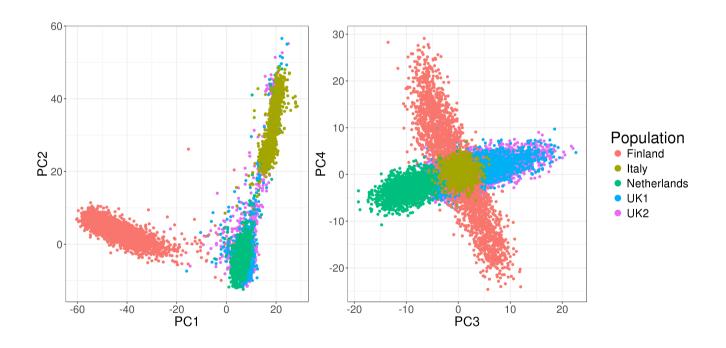
Similar accessor as Rcpp matrices

```
// [[Rcpp::depends(BH, bigstatsr)]]
#include <bigstatsr/BMAcc.h>
// [[Rcpp::export]]
NumericVector big_colsums(Environment BM) {
  XPtr<FBM> xpBM = BM["address"];
  BMAcc<double> macc(xpBM);
  size_t n = macc.nrow();
  size_t m = macc.ncol();
  NumericVector res(m);
  for (size_t j = 0; j < m; j++)</pre>
    for (size_t i = 0; i < n; i++)</pre>
      res[j] += macc(i, j);
  return res;
```

Some examples from my work

Partial Singular Value Decomposition

 $15K \times 100K - 10$ first PCs - 6 cores - 1 min (vs 2h in base R)

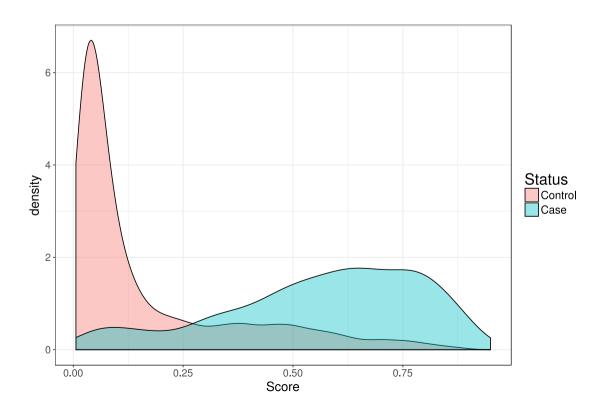


Implemented in big_randomSVD(), powered by R packages {RSpectra} and {Rcpp}.

Sparse linear models

Predicting complex diseases with a penalized logistic regression

 $15K \times 280K - 6 cores - 2 min$



Let us try some functions

Create an FBM object

```
X <- FBM(10e3, 1000, backingfile = "test2")
object.size(X)

## 648 bytes

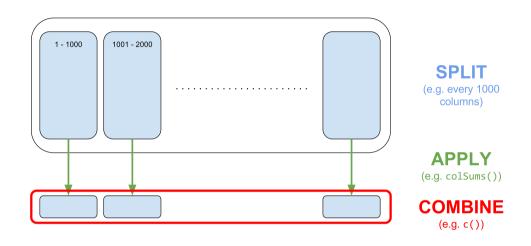
file.size(X$backingfile) ## 8 x 1e4 x 1e3

## [1] 8e+07

typeof(X)

## [1] "double"</pre>
```

Fill it with random values



```
big_apply(X, a.FUN = function(X, ind) {
   X[, ind] <- rnorm(nrow(X) * length(ind))
   NULL ## Here, you don't want to return anything
}, a.combine = 'c')</pre>
```

NULL

```
X[1:5, 1]
```

[1] 0.5372871 -0.7843506 1.7548309 -2.1620595 0.3548868

Correlation matrix

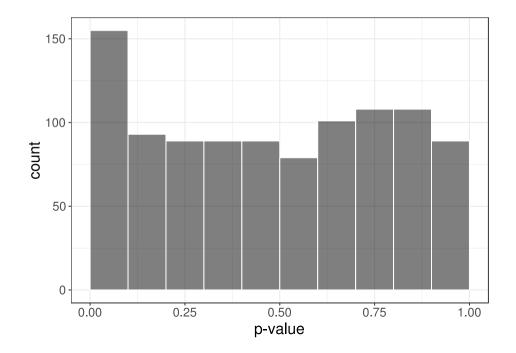
```
mat <- X[]
system.time(corr1 <- cor(mat))</pre>
##
    user system elapsed
##
   7.593 0.009 7.603
system.time(corr2 <- big_cor(X))</pre>
##
    user system elapsed
    0.434 0.032 0.466
##
all.equal(corr1, corr2[])
## [1] TRUE
```

Partial Singular Value Decomposition

```
system.time(svd1 <- svd(scale(mat), nu = 10, nv = 10))
##
    user system elapsed
##
    4.650 0.212 4.867
# Quadratic in the smallest dimension, linear in the other one
system.time(svd2 <- big SVD(X, fun.scaling = big scale(), k = 10))
## (1)
   user system elapsed
##
##
    0.761 0.027 0.792
# Linear in both dimensions
# Extremely useful if both dimensions are very large
system.time(svd3 <- big_randomSVD(X, fun.scaling = big_scale(), k =</pre>
##
    user system elapsed
##
    2.039 0.000 2.040
```

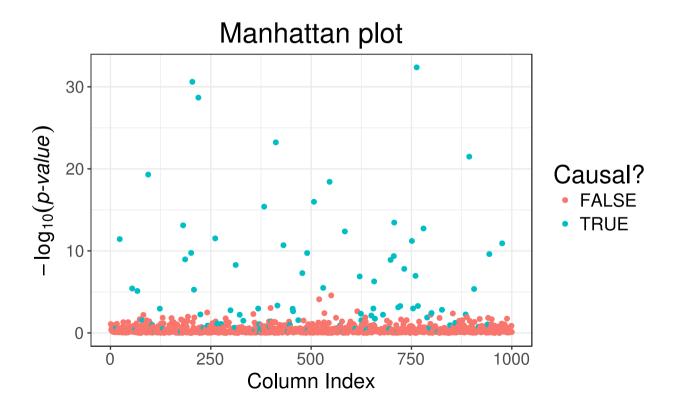
Multiple association

```
M <- 100 # number of causal variables
set <- sample(ncol(X), M)
y <- scale(X[, set]) %*% rnorm(M)
y <- y + rnorm(length(y), sd = 2 * sd(y))
mult_test <- big_univLinReg(X, y, covar.train = svd2$u)
plot(mult_test)</pre>
```



Multiple association

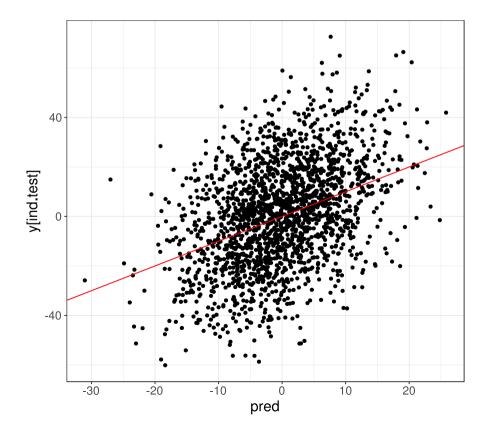
```
library(ggplot2)
plot(mult_test, type = "Manhattan") +
  aes(color = cols_along(X) %in% set) +
  labs(color = "Causal?")
```



Prediction

Prediction

```
# Plot true value vs prediction
qplot(pred, y[ind.test]) +
  geom_abline(intercept = 0, slope = 1, color = "red") +
  theme_bigstatsr()
```



Toy case:

Compute the sum for each column

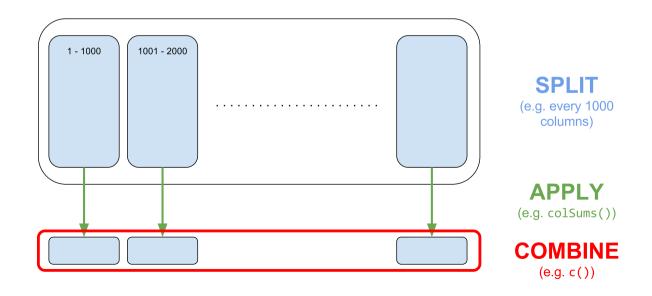
Brute force solution

sums1 <- colSums(X[]) ## /!\ access all the data in memory</pre>



Do it by blocks

[1] TRUE



Using Rcpp (1/3)

```
// [[Rcpp::depends(bigstatsr, BH)]]
#include <bigstatsr/BMAcc.h>
#include <Rcpp.h>
using namespace Rcpp;
// [[Rcpp::export]]
NumericVector bigcolsums(Environment BM) {
  XPtr<FBM> xpBM = BM["address"]; // get the external pointer
  BMAcc<double> macc(xpBM); // create an accessor to the data
  size_t i, j, n = macc.nrow(), m = macc.ncol();
  NumericVector res(m); // vector of m zeros
  for (j = 0; j < m; j++)
    for (i = 0; i < n; i++)
     res[j] += macc(i, j);
  return res;
```

Using Rcpp (1/3)

```
sums3 <- bigcolsums(X)
all.equal(sums3, sums1)</pre>
```

[1] TRUE

Using Rcpp (2/3): the bigstatsr way

```
// [[Rcpp::depends(bigstatsr, BH)]]
#include <bigstatsr/BMAcc.h>
#include <Rcpp.h>
using namespace Rcpp;
// [[Rcpp::export]]
NumericVector bigcolsums2(Environment BM,
                          const IntegerVector& rowInd,
                          const IntegerVector& colInd) {
  XPtr<FBM> xpBM = BM["address"];
  SubBMAcc<double> macc(xpBM, rowInd - 1, colInd - 1);
  size_t i, j, n = macc.nrow(), m = macc.ncol();
  NumericVector res(m); // vector of m zeros
  for (j = 0; j < m; j++)
   for (i = 0; i < n; i++)
     res[j] += macc(i, j);
  return res;
```

Using Rcpp (2/3): the bigstatsr way

```
sums4 <- bigcolsums2(X, rows_along(mat), cols_along(mat))
all.equal(sums4, sums1)

## [1] TRUE

sums5 <- bigcolsums2(X, rows_along(mat), 1:10)
all.equal(sums5, sums1[1:10])

## [1] TRUE</pre>
```

Using Rcpp (3/3): already implemented

```
sums6 <- big_colstats(X)
str(sums6)

## 'data.frame': 1000 obs. of 2 variables:
## $ sum: num -78.4 51.8 29.14 9.45 70.98 ...
## $ var: num 1.009 1.01 0.999 0.985 1.017 ...

all.equal(sums6$sum, sums1)</pre>
## [1] TRUE
```

Parallelism

Most of the functions are parallelized

```
ind.rep <- rep(cols_along(X), each = 100) ## size: 100,000
 (NCORES <- nb_cores())</pre>
## [1] 2
system.time(
  mult_test2 <- big_univLinReg(X, y, covar.train = svd2$u,</pre>
                                ind.col = ind.rep)
## user system elapsed
## 9.677 0.007 9.701
system.time(
  mult_test3 <- big_univLinReg(X, y, covar.train = svd2$u,</pre>
                                ind.col = ind.rep, ncores = NCORES)
##
   user system elapsed
    0.036 0.031 6.001
##
```

Parallelize your own functions

```
system.time(
  mult_test4 <- big_parallelize(</pre>
    X, p.FUN = function(X, ind, y, covar) {
      bigstatsr::big_univLinReg(X, y, covar.train = covar,
                                 ind.col = ind)
    }, p.combine = "rbind", ind = ind.rep,
    ncores = NCORES, y = y, covar = svd2$u)
##
    user system elapsed
    0.125 0.072 5.679
##
all.equal(mult_test4, mult_test3)
## [1] TRUE
```

Conclusion

I'm able to run algorithms on 100GB of data in \bigcirc on my computer

Advantages of using FBM objects

- you can apply algorithms on data larger than your RAM,
- you can easily **parallelize** your algorithms because the data on disk is shared,
- you write more efficient algorithms (you do less copies and think more about what you're doing),
- you can use **different types of data**, for example, in my field, I'm storing my data with only 1 byte per element (rather than 8 bytes for a standard R matrix). See the documentation of the FBM class for details.

R Packages

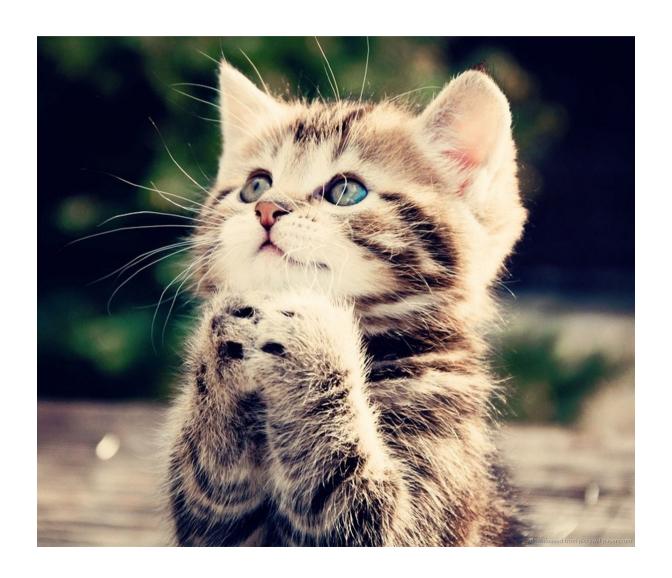
Efficient analysis of large-scale genome-wide data with two R packages: bigstatsr and bigsnpr 3

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Bioinformatics, bty185, https://doi.org/10.1093/bioinformatics/bty185

- {bigstatsr}: to be used by any field of research
- {bigsnpr}: algorithms specific to my field of research

Contributors are welcomed!



Make sure to grab an hex sticker



Thanks!

Presentation: https://privefl.github.io/R-presentation/bigstatsr.html

Package's website: https://privefl.github.io/bigstatsr/

DOI: 10.1093/bioinformatics/bty185



Slides created via the R package **xaringan**.