

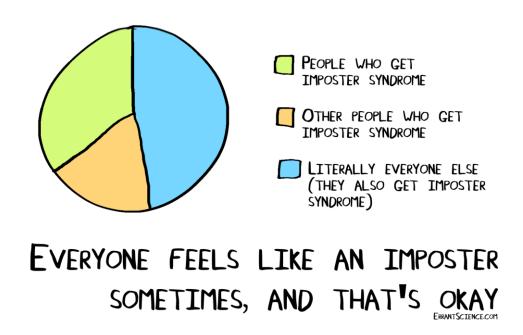
Improving the speed of a discrete event simulation model in breast cancer screening...

Stuart Wright, PhD stuart.j.wright@manchester.ac.uk

....the experience of an intermediate level R user!

Caveats

- Learnings from an ongoing process
- Huge amount of work went into original code
- Very happy to hear additional suggestions
- I apologise in advance for my novice markdown copy and pasting!

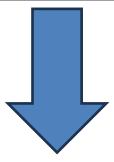


Evaluation of a Stratified National Breast Screening Program in the United Kingdom: An Early Model-Based Cost-Effectiveness Analysis



Ewan Gray, PhD^{1,2}, Anna Donten, MSc¹, Nico Karssemeijer, PhD^{1,3}, Carla van Gils, PhD^{1,4}, D. Gareth Evans, MD, FRCP^{1,5}, Sue Astley, PhD^{1,6}, Katherine Payne, PhD^{1,*}

¹Manchester Centre for Health Economics, University of Manchester, Manchester, UK; ²Usher Institute of Population Health Sciences and Informatics, University of Edinburgh, Edinburgh, UK; ³Radboud University Nijmegen Medical Centre, Nijmegen, Netherlands; ⁴University Medical Centre Utrecht, Utrecht, Netherlands; ⁵Genesis Breast Cancer Prevention Centre and Nightingale Breast Screening Centre, University Hospital of South Manchester, Manchester, UK; ⁶Department of Imaging Science and Biomedical Engineering, University of Manchester, Manchester, UK



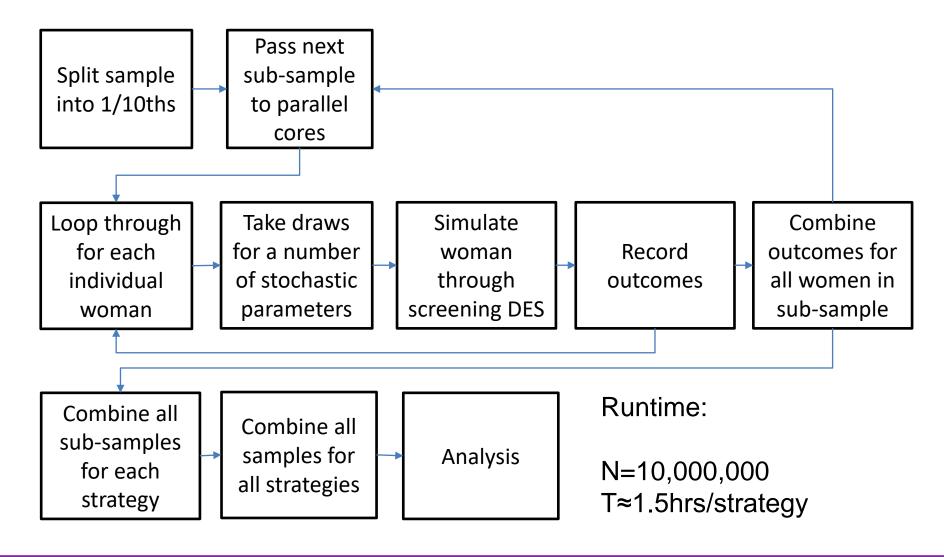
Applied Health Economics and Health Policy (2024) 22:527–542 https://doi.org/10.1007/s40258-024-00887-z

ORIGINAL RESEARCH ARTICLE

A structured process for the validation of a decision-analytic model: application to a cost-effectiveness model for risk-stratified national breast screening

Stuart J. Wright 10 · Ewan Gray 20 · Gabriel Rogers 10 · Anna Donten 10 · Katherine Payne 10

Rough Outline of Original Structure



Key Approaches to Speeding Up a Model

- Improve model code -> increase speed per run
- Reduce unnecessary variance -> reduce total N
- Improve computational power -> increase speed per run/improve parallelisation

Improve Model Code

- Everything in a loop is done i times
- Check for redundant code and where possible take these things out of a loop:
 - Defining new parameters
 - If statements (particularly if complex)
 - Objects that increase in size

Example: Function to see if cancer detected

- -Function is called at every screening event
- -That's around 4-5 times per individual



Don't need to define this vector 40mil times!

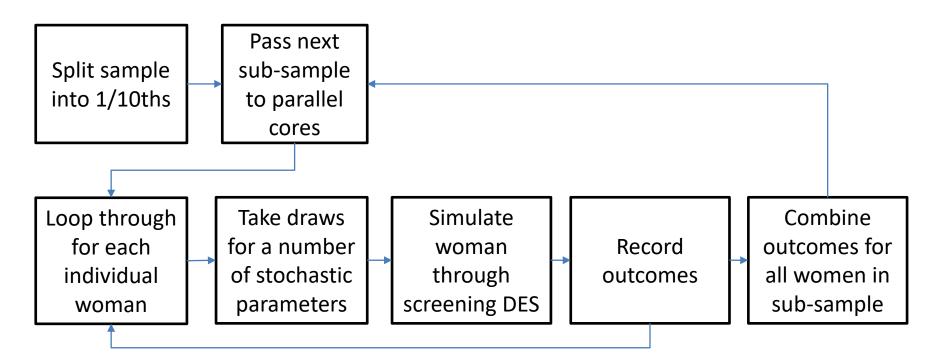
```
Sen_VDG<-c(0.85,0.776,0.695,0.61)
dense_OR <- (Sen_VDG[VDG]/(1-Sen_VDG[VDG]))/(Sen_VDG_av/(1-Sen_VDG_av))
Sensitivity <- ((Sensitivity/(1-Sensitivity))*dense_OR)/(1+((Sensitivity/(1-Sensitivity)))*dense_OR))</pre>
```

The Big One!

Pre-create a sample (data.frame) of individuals with pre-drawn stochastic parameters

Iterate over each row of the data.frame

The Problem



A: We have to do multiple <u>individual draws from parameter distributions</u> for each woman

B: Each sample of women for each strategy is <u>different</u> – they vary in ways that won't affect the results

A Useful Rule for Stochastic Models (inc PSA)

It is much quicker to take X random draws from a distribution than to do individually draw from the distribution X times

```
library(tictoc)
probdraw<-numeric(length(1000000))
tic()
for (i in 1:1000000) {
probdraw[i]<-rbeta(1,50,50) _
toc()
```

Vector of probabilities is different every time code is run

```
## 1.65 sec elapsed
```

```
probdraw<-numeric(length(1000000))
fairprobdraw<-numeric(length(1000000))
tic()
probdraw<-rbeta(1000000,50,50)
for (i in 1:1000000) {
  fairprobdraw[i] <-probdraw[i] 4
toc()
```

Vector of probabilities is the same every time code is run

BONUS!

By setting the seed, reviewers should be able to reproduce exact results

•	X1.VBD [‡]	X1.tenyrrisk ‡	X1.liferisk ‡	X1.risk_group ‡	X1.VDG [‡]	X1.MRI_screen ‡	X1.US_screen ‡	X1.risk_predicted [‡]	X1.feedback ‡
1017	14.39705	4.267940	16.76400	0	3	0	0	0	0
13499	13.06310	9.807890	31.36680	0	3	0	0	0	0
9941	9.16390	3.561810	17.07960	0	3	0	0	0	0
12204	5.87783	3.760580	15.05710	0	2	0	0	0	0
3476	5.31918	2.953430	10.84650	0	2	0	0	0	0
4633	7.46740	3.675890	16.67480	0	2	0	0	0	0
9888	11.69403	1.748370	7.24550	0	3	0	0	0	0
14835	3.45438	2.428690	12.04680	0	1	0	0	0	0
465	9.08413	1.458280	7.69616	0	3	0	0	0	0
13453	5.56860	2.646900	13.00200	0	2	0	0	0	0
14950	3.66738	1.181120	4.89676	0	1	0	0	0	0
6763	20.73423	3.004370	14.00670	0	4	0	0	0	0
6273	6.73515	1.403050	5.81391	0	2	0	0	0	0
501	5.35525	1.110270	4.53265	0	2	0	0	0	0
3747	3.93368	2.828720	11.45760	0	1	0	0	0	0
9509	11.15860	1.712360	8.74569	0	3	0	0	0	0
5939	3.11133	1.445100	5.97387	0	1	0	0	0	0
8943	8.11435	3.430440	13.11100	0	3	0	0	0	0
6611	12.95128	2.625970	12.90230	0	3	0	0	0	0
1101	3.66703	2.085600	8.47259	0	1	0	0	0	0

The parameters above this box are all generated with if statements – now outside the loop

BUT...

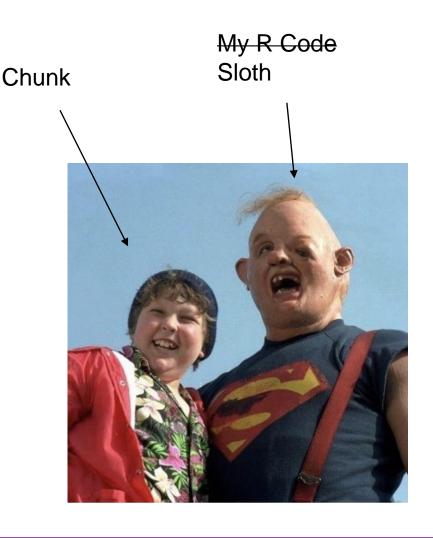
1. R holds data in RAM – making a huge data.frame means it will run slowly or not at all

2. Iterating over the rows of the frame rather than over a simple sequence of numbers requires a bit of a tweak



Solution 1: Chunking

- Split the big data.frame into chunks of smaller ones
- Save the chunks to drive
- Remove the data.frames from memory
- In the outer loop load one of the chunks
- Run through model
- Save results
- Load next chunk...

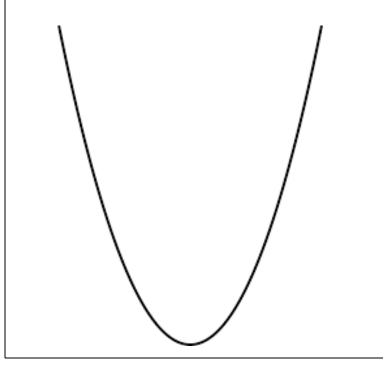


A note of caution on chunks

runtime

 More chunks means less data in memory

- BUT running each chunk requires two slow processes
 - Loading data in R
 - Initialising the parallelisation
- So need to find the optimal number of chunks which depends on N



N chunks

Solution 2: Parallelising a data.frame

```
results <- foreach(i=n/chunks,.combine = 'rbind',.packages = c('MASS','dqrng','tidyverse')) %dopar% {
```

- -This is called an iterator
- -Essentially R makes a sequential vector with the numbers 1:n/chunks
- -PROBLEM: each core gets the same rows (1:n/chunks)
- -Actually need each core to work on separate set of rows

Tip: supposedly a faster random number generator than in base R

Different iterators

 Using the iterators package you can iterate over different types of objects

```
Chunk of main data.frame

itx<-iter(splitsample,by="row")

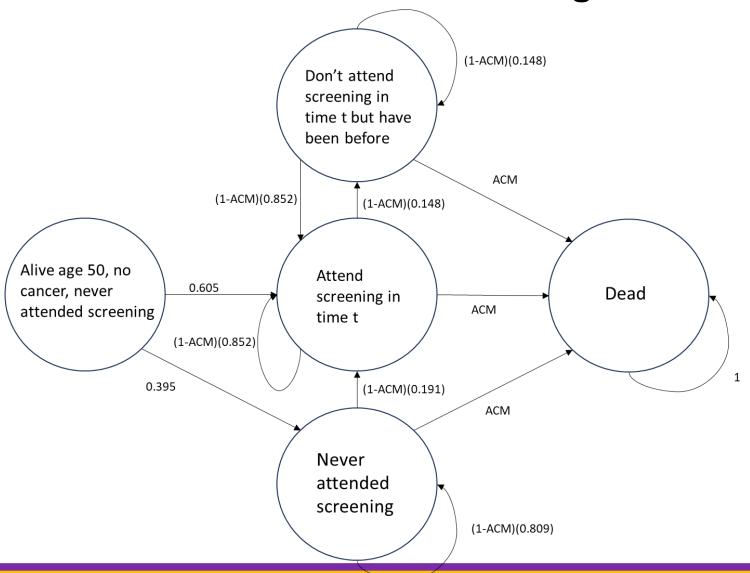
results <- foreach(i=itx,.combine = 'rbind',.packages = c('MASS','dqrng','tidyverse')) %dopar% {
```

This one line took me about a 6 months to figure out

Results

	Original Model	MANC-RISK- SCREEN
N per strategy	10,000,000	3,000,000
Run time per strategy (mins)	~90	10-15 depending on complexity

~87% of women in the model don't get cancer



Screening models: pushing the limits of R?

- Building a cervical cancer screening model
- Infectious disease model of HPV transmission
- Dynamic effect of vaccination over time
- Pre-cancer and cancer growth model
- Very complex screening and triage strategies
- Considering using Python for data generation, R for analysis

Thank You!