Calibrating a 3-state cancer model

Directed search using Nelder-mead

The DARTH workgroup

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Please cite our publications when using this code:

- Alarid-Escudero F, Maclehose RF, Peralta Y, Kuntz KM, Enns EA. Non-identifiability in model calibration and implications for medical decision making. Med Decis Making. 2018; 38(7):810-821. https://pubmed.ncbi.nlm.nih.gov/30248276/
- Jalal H, Pechlivanoglou P, Krijkamp E, Alarid-Escudero F, Enns E, Hunink MG. An Overview of R in Health Decision Sciences. Med Decis Making. 2017; 37(3): 735-746. https://journals.sagepub.com/doi/abs/10.1177/0272989X16686559

A walkthrough of the code could be found in the following link: - https://darth-git.github.io/calibSMDM2018-materials/

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Change eval to TRUE if you want to knit this document.

```
rm(list = ls())  # clear memory (removes all the variables from the workspace)
```

00 Calibration Specifications

Model: 3-State Cancer Relative Survival (CRS) Markov Model

Inputs to be calibrated: p_Mets, p_DieMets

Targets: Surv

Calibration method: Directed search using Nelder-mead

Goodness-of-fit measure: Sum of Log-Likelihood

01 Load packages

```
if (!require('pacman')) install.packages('pacman'); library(pacman) # use this package to conveniently
# load (install if required) packages from CRAN
p_load("lhs", "plotrix", "psych")
```

02 Load target data

```
load("CRS CalibTargets.RData")
lst_targets <- CRS_targets</pre>
# Plot the targets
# TARGET 1: Survival ("Surv")
plotrix::plotCI(x = lst_targets$Surv$time, y = lst_targets$Surv$value,
                ui = lst_targets$Surv$ub,
                li = lst_targets$Surv$lb,
                ylim = c(0, 1),
                xlab = "Time", ylab = "Pr Survive")
# TARGET 2: (if you had more...)
\# plotrix::plotCI(x = lst_targets$Target2$time, y = lst_targets$Target2$value,
                  ui = lst_targets$Target2$ub,
#
                  li = lst targets$Target2$lb,
#
                  ylim = c(0, 1),
                  xlab = "Time", ylab = "Target 2")
#
```

03 Load model as a function

```
# - inputs are parameters to be estimated through calibration
# - outputs correspond to the target data
source("CRS_MarkovModel_Function.R") # creates the function run_crs_markov()
# Check that it works
v_params_test <- c(p_Mets = 0.10, p_DieMets = 0.05)
run_crs_markov(v_params_test) # It works!</pre>
```

04 Specify calibration parameters

```
# Specify seed (for reproducible sequence of random numbers)
set.seed(072218)

# number of initial starting points
n_init <- 100

# names and number of input parameters to be calibrated
v_param_names <- c("p_Mets", "p_DieMets")
n_param <- length(v_param_names)

# range on input search space
1b <- c(p_Mets = 0.04, p_DieMets = 0.04) # lower bound
ub <- c(p_Mets = 0.16, p_DieMets = 0.16) # upper bound

# number of calibration targets
v_target_names <- c("Surv")
n_target <- length(v_target_names)</pre>
```

05 Calibration functions

06 Calibrate!

```
# record start time of calibration
t_init <- Sys.time()</pre>
### Sample multiple random starting values for Nelder-Mead ###
v_params_init <- matrix(nrow=n_init, ncol=n_param)</pre>
for (i in 1:n_param){
 v_params_init[,i] <- runif(n_init,min=lb[i],max=ub[i])</pre>
colnames(v_params_init) <- v_param_names</pre>
### Run Nelder-Mead for each starting point ###
m_calib_res <- matrix(nrow = n_init, ncol = n_param+1)</pre>
colnames(m_calib_res) <- c(v_param_names, "Overall_fit")</pre>
for (j in 1:n_init){
  ### use optim() as Nelder-Mead ###
 fit_nm <- optim(v_params_init[j,], f_gof,</pre>
                  control = list(fnscale = -1, # switches from minimization to maximization
                                 maxit = 1000), hessian = T)
 m_calib_res[j,] <- c(fit_nm$par,fit_nm$value)</pre>
  ### to use a simulated annealing instead ###
  # fit_sa <- optim(v_params_init[j,], f_gof,</pre>
                    method = c("SANN"), # switches to using simulated annealing
  #
                    control = list(temp = 10, tmax = 10, # algorithm tuning parameters
  #
                                    fnscale = -1, maxit = 1000),
                    hessian = T)
  # m_calib_res[j,] = c(fit_sa$par,fit_sa$value)
  ### to use a genetic algorithm instead ###
  # library(DEoptim)
  # f fitness <- function(params){</pre>
  \# names(params) = v_param_names
```

```
# return(-f_gof(params))}
# fit_ga = DEoptim(f_fitness, lower=lb, upper=ub)
# m_calib_res[j,] = c(fit_ga$optim$bestmem,-1*fit_ga$optim$bestval)
}
# Calculate computation time
comp_time <- Sys.time() - t_init</pre>
```

07 Exploring best-fitting input sets

```
# Arrange parameter sets in order of fit
m_calib_res <- m_calib_res[order(-m_calib_res[,"Overall_fit"]),]</pre>
# Examine the top 10 best-fitting sets
m_calib_res[1:10, ]
# Plot the top 10 (top 10%)
plot(m_calib_res[1:10,1],m_calib_res[1:10,2],
     xlim=c(lb[1],ub[1]),ylim=c(lb[2],ub[2]),
     xlab = colnames(m calib res)[1], ylab = colnames(m calib res)[2])
# Pairwise comparison of top 10 sets
pairs.panels(m_calib_res[1:10,v_param_names])
### Plot model-predicted output at mean vs targets ###
v_out_best <- run_crs_markov(m_calib_res[1,])</pre>
# TARGET 1: Survival ("Surv")
plotrix::plotCI(x = lst_targets$Surv$time, y = lst_targets$Surv$value,
                ui = lst_targets$Surv$ub,
                li = lst targets$Surv$lb,
                ylim = c(0, 1),
                xlab = "Time", ylab = "Pr Survive")
points(x = lst_targets$Surv$time,
       y = v_out_best$Surv,
       pch = 8, col = "red")
legend("topright",
       legend = c("Target", "Model-predicted output"),
       col = c("black", "red"), pch = c(1, 8))
# TARGET 2: (if you had more...)
\# plotrix::plotCI(x = lst_targets$Target2$time, y = lst_targets$Target2$value,
                  ui = lst\_targets\$Target2\$ub,
#
                  li = lst_targets$Target2$lb,
#
                  ylim = c(0, 1),
                  xlab = "Time", ylab = "Target 2")
# points(x = lst_targets$Target2$time,
y = v \text{ out best$Target2},
        pch = 8, col = "red")
```

```
# legend("topright",
# legend = c("Target", "Model-predicted output"),
# col = c("black", "red"), pch = c(1, 8))
```