

SA: Markov Sick-Sicker model in R

The DARTH workgroup

Developed by the Decision Analysis in R for Technologies in Health (DARTH) workgroup:

Fernando Alarid-Escudero, PhD (1)

Eva A. Enns, MS, PhD (2)

M.G. Myriam Hunink, MD, PhD (3,4)

Hawre J. Jalal, MD, PhD (5)

Eline M. Krijkamp, MSc (3)

Petros Pechlivanoglou, PhD (6,7)

Alan Yang, MSc (7)

In collaboration of:

1. Division of Public Administration, Center for Research and Teaching in Economics (CIDE), Aguascalientes, Mexico
2. University of Minnesota School of Public Health, Minneapolis, MN, USA
3. Erasmus MC, Rotterdam, The Netherlands
4. Harvard T.H. Chan School of Public Health, Boston, USA
5. University of Pittsburgh Graduate School of Public Health, Pittsburgh, PA, USA
6. University of Toronto, Toronto ON, Canada
7. The Hospital for Sick Children, Toronto ON, Canada

Please cite our publications when using this code:

- 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>
- Krijkamp EM, Alarid-Escudero F, Enns EA, Jalal HJ, Hunink MGM, Pechlivanoglou P. Microsimulation modeling for health decision sciences using R: A tutorial. *Med Decis Making*. 2018;38(3):400-22. <https://journals.sagepub.com/doi/abs/10.1177/0272989X18754513>
- Krijkamp EM, Alarid-Escudero F, Enns E, Pechlivanoglou P, Hunink MM, Jalal H. A Multidimensional Array Representation of State-Transition Model Dynamics. *Med Decis Making*. Feb;40(2):242-248. <https://doi.org/10.1177/0272989X19893973>
- Alarid-Escudero, F., Krijkamp, E. M., Enns, E. A., Hunink, M. G. M., Pechlivanoglou, P., & Jalal, H. (2020). Cohort state-transition models in R: From conceptualization to implementation. *ArXiv:2001.07824v1*, 1-31. <http://arxiv.org/abs/2001.07824>

Copyright 2017, THE HOSPITAL FOR SICK CHILDREN AND THE COLLABORATING INSTITUTIONS. All rights reserved in Canada, the United States and worldwide. Copyright, trademarks, trade names and any and all associated intellectual property are exclusively owned by THE HOSPITAL FOR SICK CHILDREN and the collaborating institutions. These materials may be used, reproduced, modified, distributed and adapted with proper attribution.

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

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

01 Load packages

```
if (!require('pacman')){
  install.packages('pacman')
  library(pacman) # use this package to conveniently install other packages
}
# load (install if required) packages from CRAN
pacman::p_load("dplyr", "devtools", "scales", "ellipse", "ggplot2", "lazyeval",
               "igraph", "truncnorm", "ggraph", "reshape2", "knitr", "stringr",
               "diagram", "dampack")
# load (install if required) packages from GitHub
# install_github("DARTH-git/darthtools", force = TRUE) Uncomment if there is a newer version
pacman::p_load_gh("DARTH-git/darthtools")
```

02 Load functions

```
# No function needed
```

03 Input model parameters

```
# Strategy names
v_names_str <- c("No Treatment", "Treatment")

# Markov model parameters
age      <- 25                # age at baseline
max_age  <- 55                # maximum age of follow up
n_t      <- max_age - age     # time horizon, number of cycles
v_names_states <- c("H", "S1", "S2", "D") # the 4 states of the model: Healthy (H), Sick (S1),
                                           # Sicker (S2), Dead (D)

v_init <- c("H" = 1,
            "S1" = 0,
            "S2" = 0,
            "D" = 0)          # initial cohort distribution (everyone allocated to the "healthy"

# Transition probabilities (per cycle)
p_HD      <- 0.005            # probability to die when healthy
p_HS1     <- 0.15             # probability to become sick when healthy, conditional on surviving
p_S1H     <- 0.5              # probability to become healthy when sick, conditional on surviving
p_S1S2    <- 0.105           # probability to become sicker when sick, conditional on surviving
hr_S1     <- 3                # hazard ratio of death in sick vs healthy
hr_S2     <- 10               # hazard ratio of death in sicker vs healthy
r_HD      <- - log(1 - p_HD)  # rate of death in healthy
r_S1D     <- hr_S1 * r_HD     # rate of death in sick
r_S2D     <- hr_S2 * r_HD     # rate of death in sicker
p_S1D     <- 1 - exp(-r_S1D)  # probability to die in sick
p_S2D     <- 1 - exp(-r_S2D)  # probability to die in sicker
```

```

# Cost and utility inputs
c_H   <- 2000 # cost of remaining one cycle in the healthy state
c_S1  <- 4000 # cost of remaining one cycle in the sick state
c_S2  <- 15000 # cost of remaining one cycle in the sicker state
c_trt <- 12000 # cost of treatment (per cycle)
c_D   <- 0    # cost of being in the death state
u_H   <- 1    # utility when healthy
u_S1  <- 0.75 # utility when sick
u_S2  <- 0.5  # utility when sicker
u_D   <- 0    # utility when dead
u_trt <- 0.95 # utility when being treated
d_e   <- 0.03 # discount rate per cycle for costs by 3%
d_c   <- 0.03 # discount rate per cycle for QALYs by 3%

n_str   <- length(v_names_str)    # Number of strategies
n_states <- length(v_names_states) # number of states

# Discount weights for costs and effects
v_dwc <- 1 / (1 + d_c) ^ (0:n_t)
v_dwe <- 1 / (1 + d_e) ^ (0:n_t)

```

Create a state-transition diagram of the cohort model

```

m_P_diag <- matrix(0,
                    nrow = n_states, ncol = n_states,
                    dimnames = list(v_names_states, v_names_states))
m_P_diag["H" , "S1"] = ""
m_P_diag["H" , "D" ] = ""
m_P_diag["H" , "H" ] = ""
m_P_diag["S1", "H" ] = ""
m_P_diag["S1", "S2"] = ""
m_P_diag["S1", "D" ] = ""
m_P_diag["S1", "S1"] = ""
m_P_diag["S2", "D" ] = ""
m_P_diag["S2", "S2"] = ""
m_P_diag["D", "D" ] = ""
layout.fig <- c(3, 1)

plotmat(t(m_P_diag), t(layout.fig), self.cex = 0.5, curve = 0, arr.pos = 0.7,
        latex = T, arr.type = "curved", relsize = 0.9, box.prop = 0.8,
        cex = 0.8, box.cex = 0.9, lwd = 1)

```

04 Define and initialize matrices and vectors

04.1 Cohort trace

```

# create the markov trace matrix M capturing the proportion of the cohort in
# each state at each cycle
m_M_notrt <- m_M_trt <- matrix(NA,
                                nrow = n_t + 1, ncol = n_states,
                                dimnames = list(paste("cycle", 0:n_t, sep = " "),
                                                  v_names_states))

```

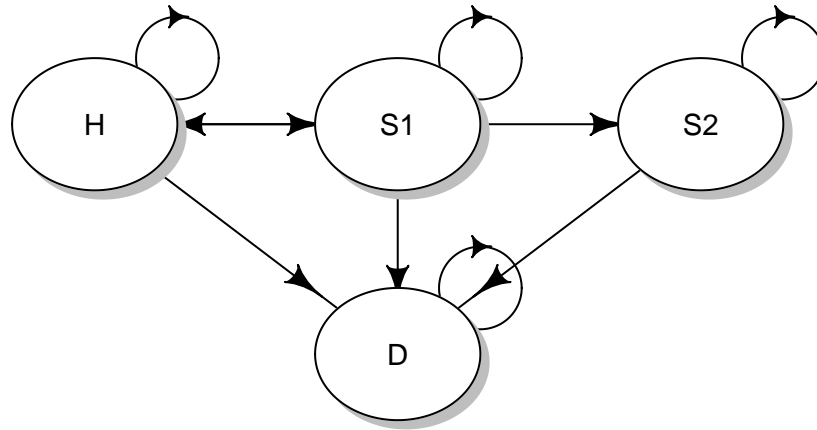


Figure 1: State-transition diagram of the time-independent Sick-Sicker cohort state-transition model.

```
head(m_M_notrt) # show first 6 rows of the matrix
```

```
##      H S1 S2 D
## cycle 0 NA NA NA NA
## cycle 1 NA NA NA NA
## cycle 2 NA NA NA NA
## cycle 3 NA NA NA NA
## cycle 4 NA NA NA NA
## cycle 5 NA NA NA NA
```

```
# The cohort starts as healthy
```

```
m_M_notrt[1, ] <- c(1, 0, 0, 0) # initiate first cycle of cohort trace
```

```
m_M_trt[1, ] <- c(1, 0, 0, 0) # initiate first cycle of cohort trace
```

04.2 Transition probability matrices

```
# create the transition probability matrix for both treatment arms
```

```
m_P_notrt <- m_P_trt <- matrix(0,
                                nrow = n_states,
                                ncol = n_states,
                                dimnames = list(v_names_states,
                                                  v_names_states)) # name the columns and rows of the matrix
```

```
m_P_notrt
```

```
##      H S1 S2 D
## H    0  0  0  0
## S1   0  0  0  0
## S2   0  0  0  0
## D    0  0  0  0
```

Fill in the transition probability matrix:

```
# from Healthy
```

```
m_P_notrt["H", "H"] <- (1 - p_HD) * (1 - p_HS1)
```

```
m_P_notrt["H", "S1"] <- (1 - p_HD) * p_HS1
```

```
m_P_notrt["H", "D"] <- p_HD
```

```
# from Sick
```

```
m_P_notrt["S1", "H"] <- (1 - p_S1D) * p_S1H
```

```

m_P_notrt["S1", "S1"] <- (1 - p_S1D) * (1 - (p_S1H + p_S1S2))
m_P_notrt["S1", "S2"] <- (1 - p_S1D) * p_S1S2
m_P_notrt["S1", "D" ] <- p_S1D
# from Sicker
m_P_notrt["S2", "S2"] <- 1 - p_S2D
m_P_notrt["S2", "D" ] <- p_S2D
# from Dead
m_P_notrt["D", "D" ] <- 1

# Check that transition probabilities are in [0, 1]
check_transition_probability(m_P_notrt, verbose = TRUE)
# Check that all rows sum to 1
check_sum_of_transition_array(m_P_notrt, n_states = n_states, verbose = TRUE)

# Overwrite the transition probability matrix for treatment with the no treatment values as they are the same
m_P_trt <- m_P_notrt

```

05 Run Markov model

```

for (t in 1:n_t){      # loop through the number of cycles
  m_M_notrt[t + 1, ] <- t(m_M_notrt[t, ]) %*% m_P_notrt # estimate the Markov trace
                                                            # for the next cycle (t + 1)
  m_M_trt[t + 1, ]    <- t(m_M_trt[t, ]) %*% m_P_trt    # estimate the Markov trace
                                                            # for the next cycle (t + 1)
} # close the loop

head(m_M_notrt) # show the first 6 lines of the matrix

```

```

##           H           S1           S2           D
## cycle 0 1.0000000 0.0000000 0.00000000 0.00000000
## cycle 1 0.8457500 0.1492500 0.00000000 0.00500000
## cycle 2 0.7888043 0.1843020 0.01543735 0.01145632
## cycle 3 0.7579069 0.1894418 0.03374551 0.01890581
## cycle 4 0.7343069 0.1868303 0.05169021 0.02717260
## cycle 5 0.7130610 0.1822918 0.06848747 0.03615973

```

06 Compute and Plot Epidemiological Outcomes

06.1 Cohort trace

```

# create a plot of the data
matplot(m_M_notrt, type = 'l',
        ylab = "Probability of state occupancy",
        xlab = "Cycle",
        main = "Cohort Trace")
# add a legend to the graph
legend("topright", v_names_states,
       col = 1:n_states, lty = 1:n_states, bty = "n")

```

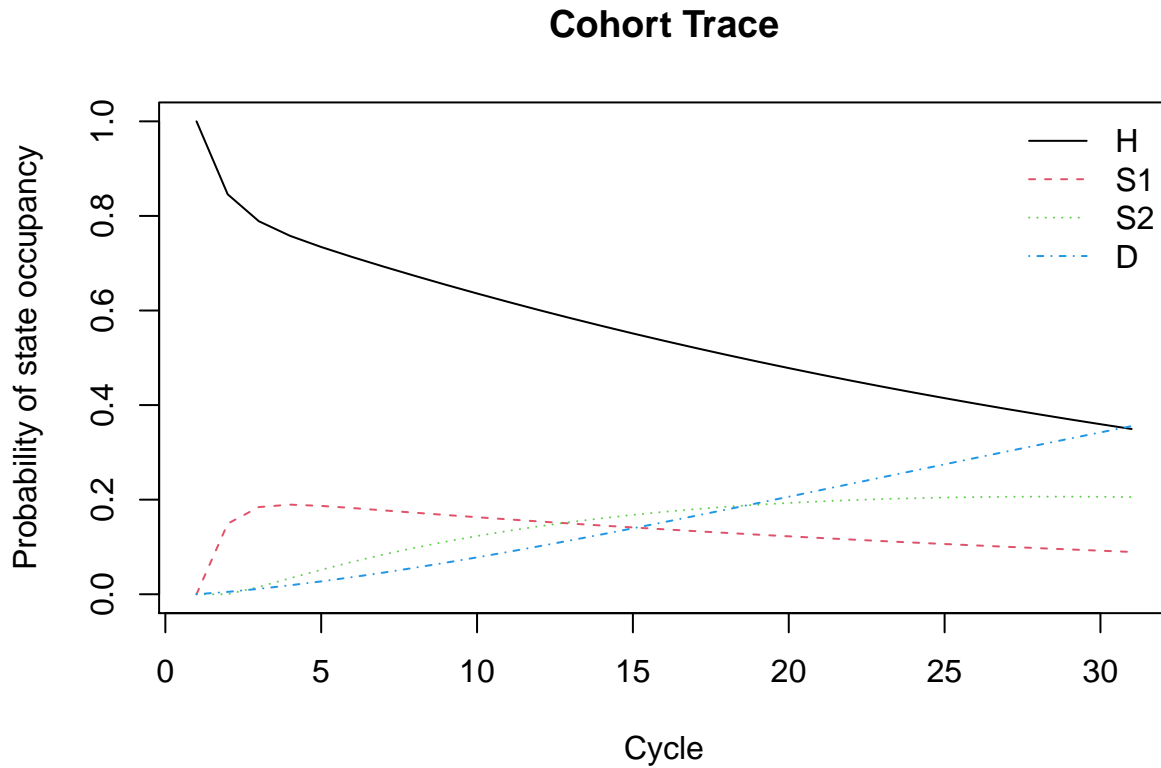


Figure 2: Cohort trace of the time-independent cSTM under no treatment.

06.2 Overall Survival (OS)

```
# calculate the overall survival (OS) probability for no treatment
v_os_notrt <- 1 - m_M_notrt[, "D"]
# alternative way of calculating the OS probability
v_os_notrt <- rowSums(m_M_notrt[, 1:3])
# create a simple plot showing the OS
plot(0:n_t, v_os_notrt, type = 'l',
     ylim = c(0, 1),
     ylab = "Survival probability",
     xlab = "Cycle",
     main = "Overall Survival")
# add grid
grid(nx = n_t, ny = 10, col = "lightgray", lty = "dotted", lwd = par("lwd"),
     equilogs = TRUE)
```

06.2.1 Life Expectancy (LE)

```
v_le <- sum(v_os_notrt) # summing probability of OS over time (i.e. life expectancy)
```

06.3 Disease prevalence

```
v_prev <- rowSums(m_M_notrt[, c("S1", "S2")]) / v_os_notrt
plot(v_prev,
     ylim = c(0, 1),
```

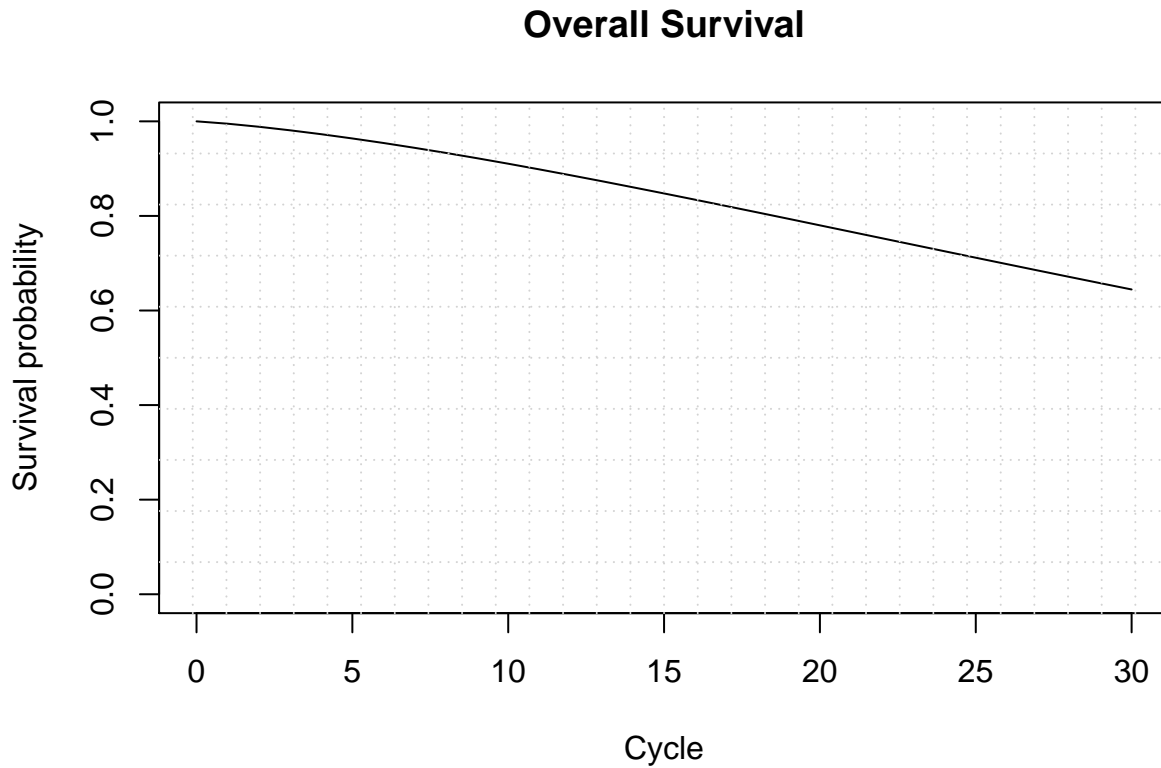


Figure 3: Overall survival of the time-independent cSTM under no treatment.

```
ylab = "Prevalence",
xlab = "Cycle",
main = "Disease prevalence")
```

06.4 Proportion of sick in S1 state

```
v_prop_S1 <- m_M_notrt[, "S1"] / v_prev
plot(0:n_t, v_prop_S1,
     xlab = "Cycle",
     ylab = "Proportion",
     main = "Proportion of sick in S1 state",
     col = "black", type = "l")
```

07 Compute Cost-Effectiveness Outcomes

```
# Vectors with costs and utilities by treatment
v_u_notrt <- c(u_H, u_S1, u_S2, u_D)
v_u_trt <- c(u_H, u_trt, u_S2, u_D)

v_c_notrt <- c(c_H, c_S1, c_S2, c_D)
v_c_trt <- c(c_H, c_S1 + c_trt, c_S2 + c_trt, c_D)
```

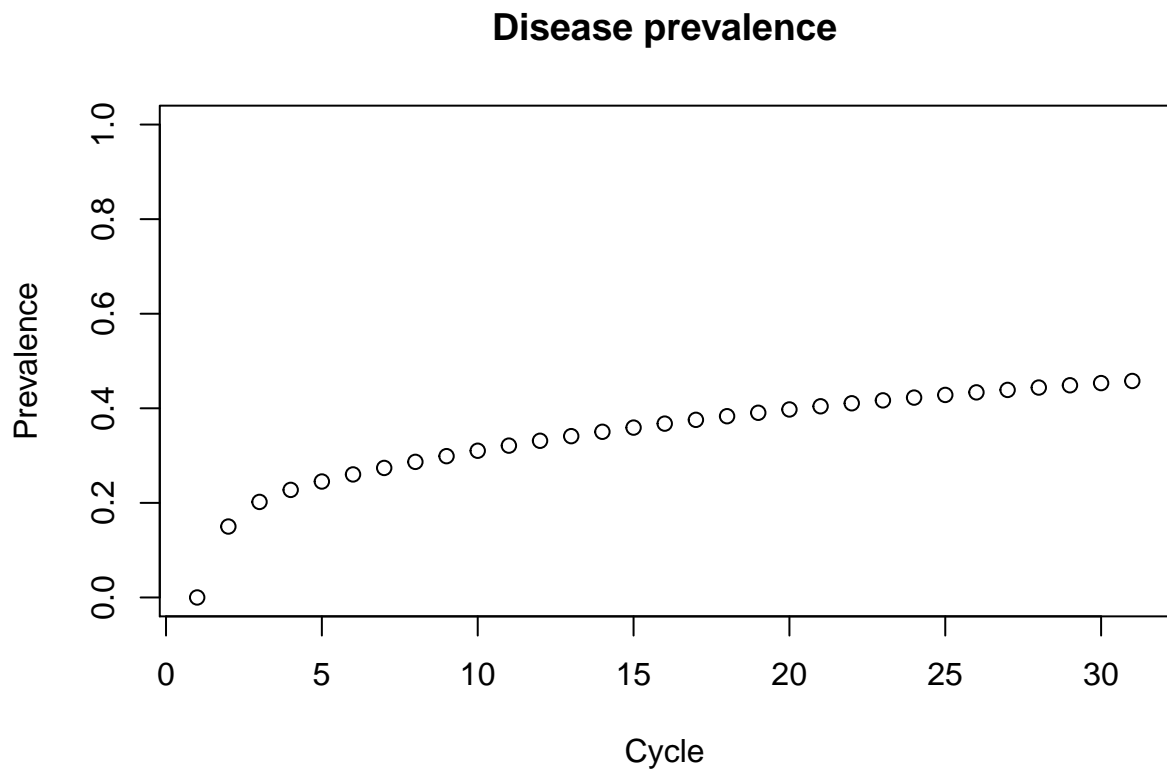


Figure 4: Disease prevalence of the time-independent cSTM under no treatment.

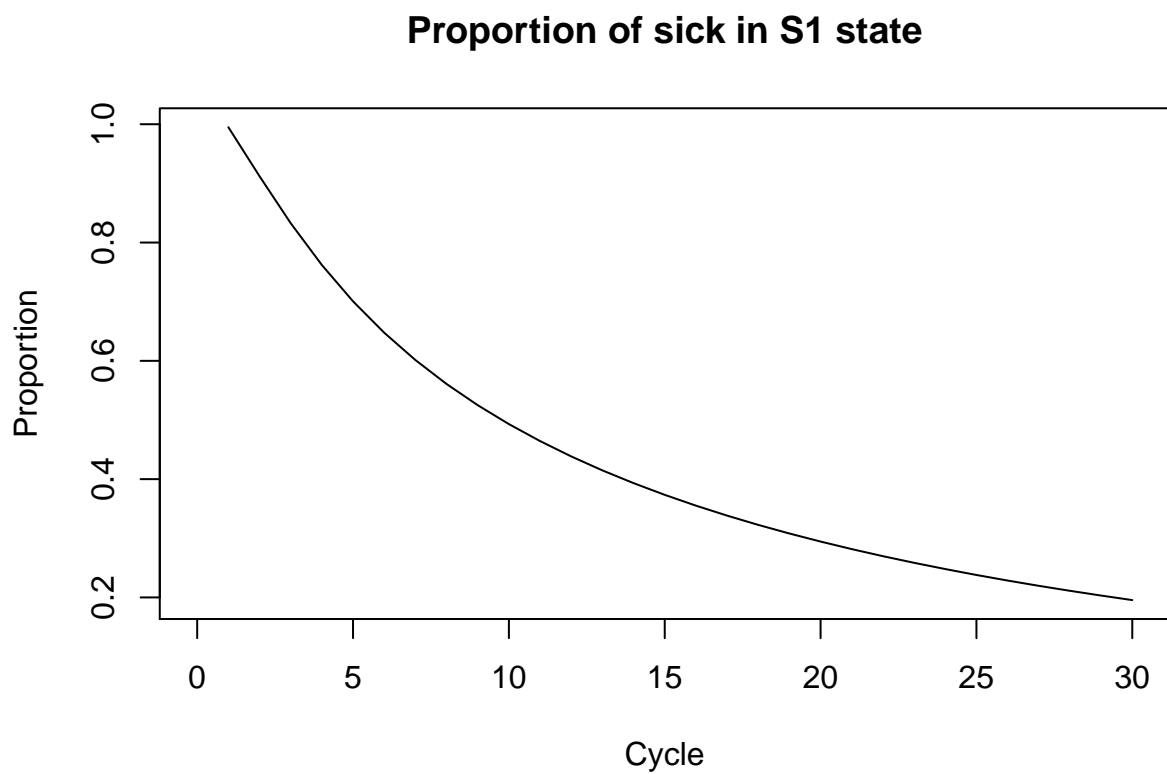


Figure 5: Proportion of sick in S1 state of the time-independent cSTM under no treatment.

07.1 Mean Costs and QALYs for Treatment and NO Treatment

```
v_tu_notrt <- m_M_notrt %>% v_u_notrt
v_tu_trt   <- m_M_trt   %>% v_u_trt

v_tc_notrt <- m_M_notrt %>% v_c_notrt
v_tc_trt   <- m_M_trt   %>% v_c_trt
```

07.2 Discounted Mean Costs and QALYs

```
tu_d_notrt <- t(v_tu_notrt) %>% v_dwe
tu_d_trt   <- t(v_tu_trt)   %>% v_dwe

tc_d_notrt <- t(v_tc_notrt) %>% v_dwc
tc_d_trt   <- t(v_tc_trt)   %>% v_dwc

# store them into a vector
v_tc_d <- c(tc_d_notrt, tc_d_trt)
v_tu_d <- c(tu_d_notrt, tu_d_trt)

# data.frame with discounted costs and effectiveness
df_ce <- data.frame(Strategy = v_names_str,
                    Cost     = v_tc_d,
                    Effect    = v_tu_d
                    )

df_ce

##           Strategy      Cost    Effect
## 1 No Treatment  75795.04 15.84802
## 2   Treatment 141511.41 16.41446
```

07.3 Compute ICERs of the Markov model

```
df_cea <- calculate_icers(cost      = df_ce$Cost,
                          effect    = df_ce$Effect,
                          strategies = df_ce$Strategy
                          )

df_cea

##           Strategy      Cost    Effect Inc_Cost Inc_Effect      ICER Status
## 1 No Treatment  75795.04 15.84802      NA      NA      NA      ND
## 2   Treatment 141511.41 16.41446 65716.37 0.5664367 116017.2      ND
```

07.4 Plot frontier of the Markov model

```
plot(df_cea, effect_units = "QALYs", xlim = c(15.6, 16.6))
```

08 Deterministic Sensitivity Analysis

08.1 List of input parameters

Create list `l_params_all` with all input probabilities, cost and utilities.

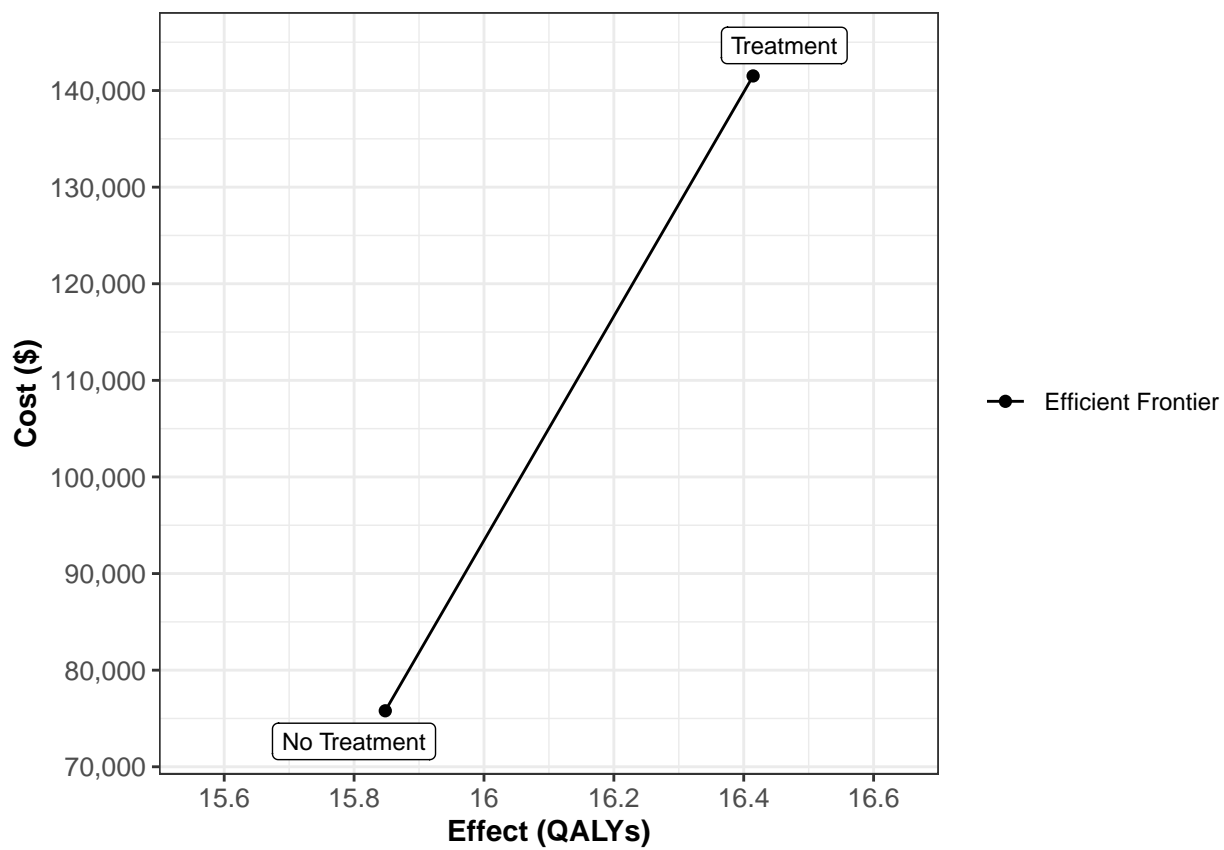


Figure 6: Cost-effectiveness efficient frontier for the time-independent Sick-Sicker model.

```

l_params_all <- as.list(data.frame(
  p_HD      = 0.005, # probability to die when healthy
  p_HS1     = 0.15,  # probability to become sick when healthy, conditional on surviving
  p_S1H     = 0.5,   # probability to become healthy when sick, conditional on surviving
  p_S1S2    = 0.105, # probability to become sicker when sick, conditional on surviving
  hr_S1     = 3,     # hazard ratio of death in sick vs healthy
  hr_S2     = 10,    # hazard ratio of death in sicker vs healthy
  c_H       = 2000,  # cost of remaining one cycle in the healthy state
  c_S1      = 4000,  # cost of remaining one cycle in the sick state
  c_S2      = 15000, # cost of remaining one cycle in the sicker state
  c_trt     = 12000, # cost of treatment(per cycle)
  c_D       = 0,     # cost of being in the death state
  u_H       = 1,     # utility when healthy
  u_S1      = 0.75,  # utility when sick
  u_S2      = 0.5,   # utility when sicker
  u_D       = 0,     # utility when dead
  u_trt     = 0.95,  # utility when treated
  d_e       = 0.03,  # discount factor for effectiveness
  d_c       = 0.03   # discount factor for costs
))

# store the parameter names into a vector
v_names_params <- names(l_params_all)

```

08.2 Load Sick-Sicker Markov model function

```

source("Functions_markov_sick-sicker_sol.R")
# Test function
calculate_ce_out(l_params_all)

```

```

##      Strategy      Cost    Effect    NMB
## 1 No Treatment  75795.04  15.84802  1509007
## 2   Treatment 141511.41  16.41446  1499935

```

08.3 One-way sensitivity analysis (OWSA)

```

options(scipen = 999) # disabling scientific notation in R
# data.frame containing all parameters, their base-case values, and the min and
# max values of the parameters of interest
df_params_owsa <- data.frame(pars = c("p_S1S2", "c_trt", "u_S1", "u_trt"),
                             min  = c(0.05, 6000, 0.65, 0.80), # min parameter values
                             max  = c(0.155, 18000, 0.85, 0.98) # max parameter values
                             )

owsa_nmb <- run_owsa_det(params_range = df_params_owsa, # data.frame with parameters for OWSA
                        params_basecase = l_params_all, # list with all parameters
                        nsamp           = 100,          # number of parameter values
                        FUN              = calculate_ce_out, # function to compute outputs
                        outcomes         = c("NMB"),      # output to do the OWSA on
                        strategies      = v_names_str,   # names of the strategies
                        n_wtp            = 120000)        # extra argument to pass to FUN

```

```
##
```

```
|
```

```
plot(owsa_nmb, txtsize = 10, n_x_ticks = 4,
     facet_scales = "free") +
  theme(legend.position = "bottom")
```

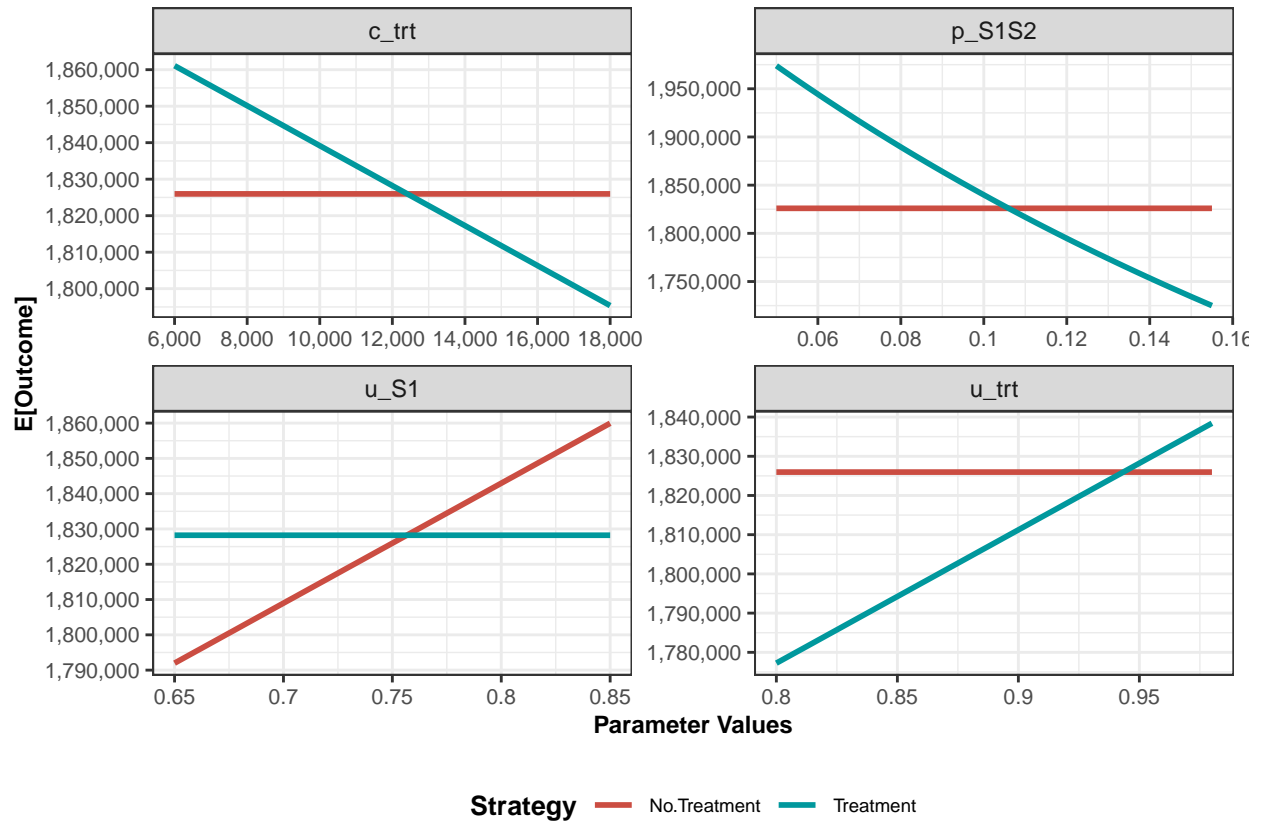


Figure 7: One-way sensitivity analysis for the time-independent Sick-Sicker model.

08.3.2 Optimal strategy with OWSA

```
owsa_opt_strat(owsa = owsa_nmb, txtsize = 10)
```

08.3.3 Tornado plot

```
owsa_tornado(owsa = owsa_nmb)
```

08.4 Two-way sensitivity analysis (TWSA)

```
# dataframe containing all parameters, their basecase values, and the min and
# max values of the parameters of interest
df_params_twsa <- data.frame(pars = c("c_trt", "u_trt"),
                             min = c(6000, 0.80), # min parameter values
                             max = c(18000, 0.98) # max parameter values
                             )

twsa_nmb <- run_twsa_det(params_range = df_params_twsa, # data.frame with parameters for TWSA
                        params_basecase = l_params_all, # list with all parameters
```

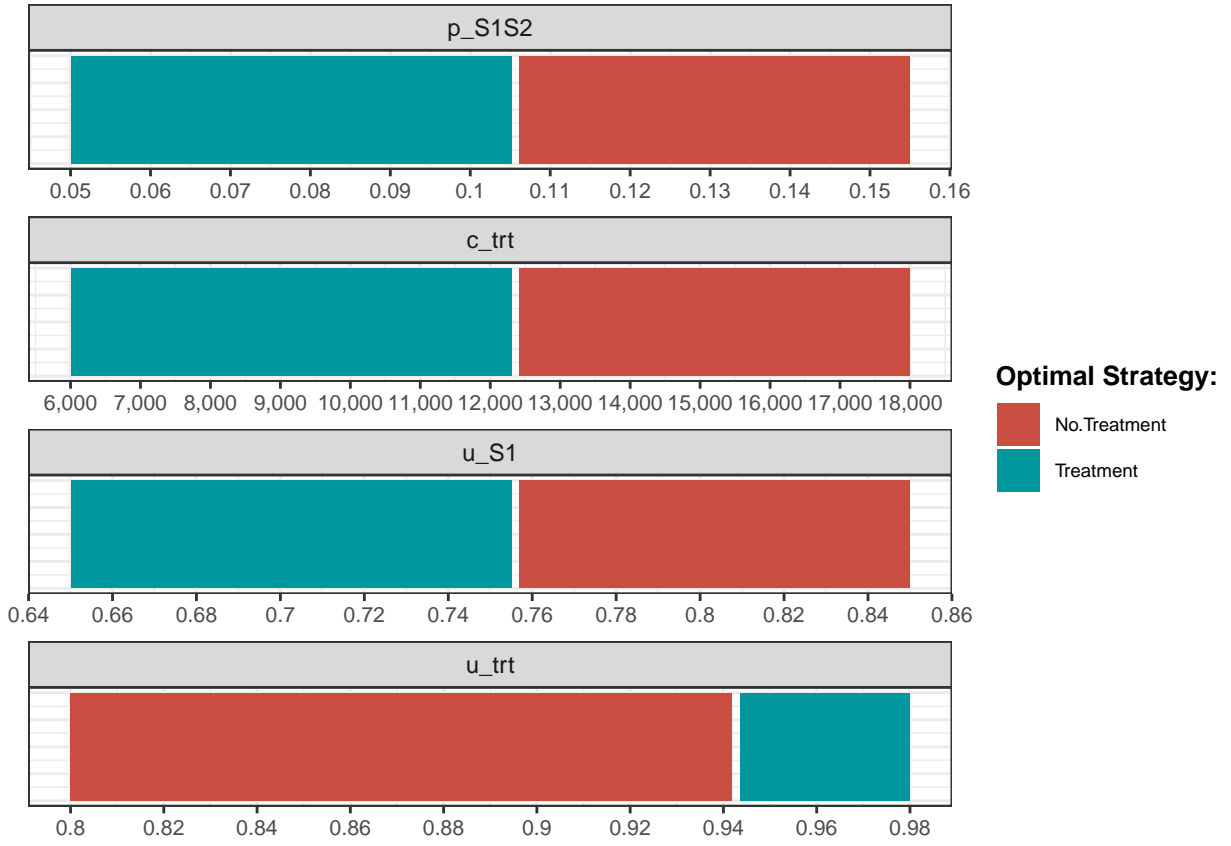


Figure 8: Optimal strategy with one-way sensitivity analysis for the time-independent Sick-Sicker model.

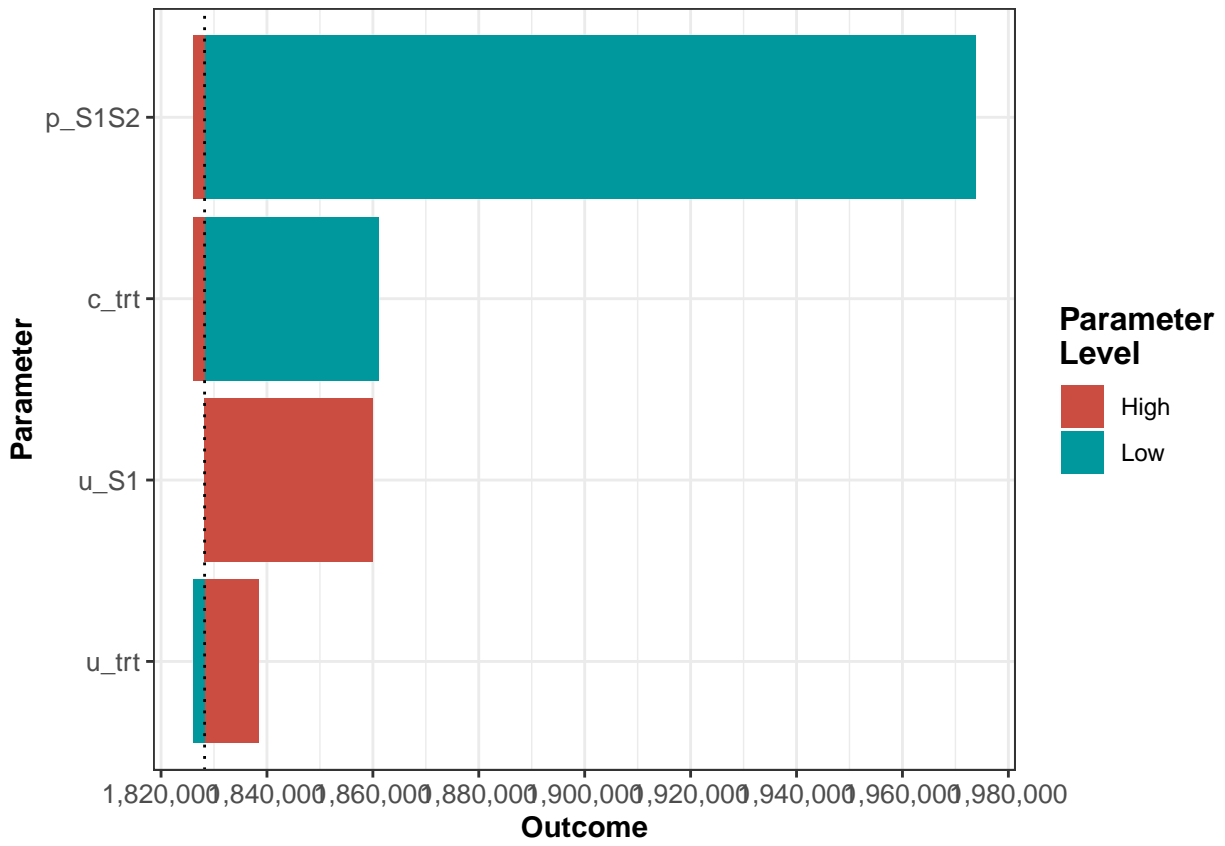


Figure 9: Tornado plot for the time-independent Sick-Sicker model.

```

nsamp      = 40,           # number of parameter values
FUN        = calculate_ce_out, # function to compute outputs
outcomes   = c("NMB"),     # output to do the TWSA on
strategies = v_names_str,   # names of the strategies
n_wtp      = 120000        # extra argument to pass to FUN

```

```
## |
```

```
|
```

```
plot(twsa_nmb)
```

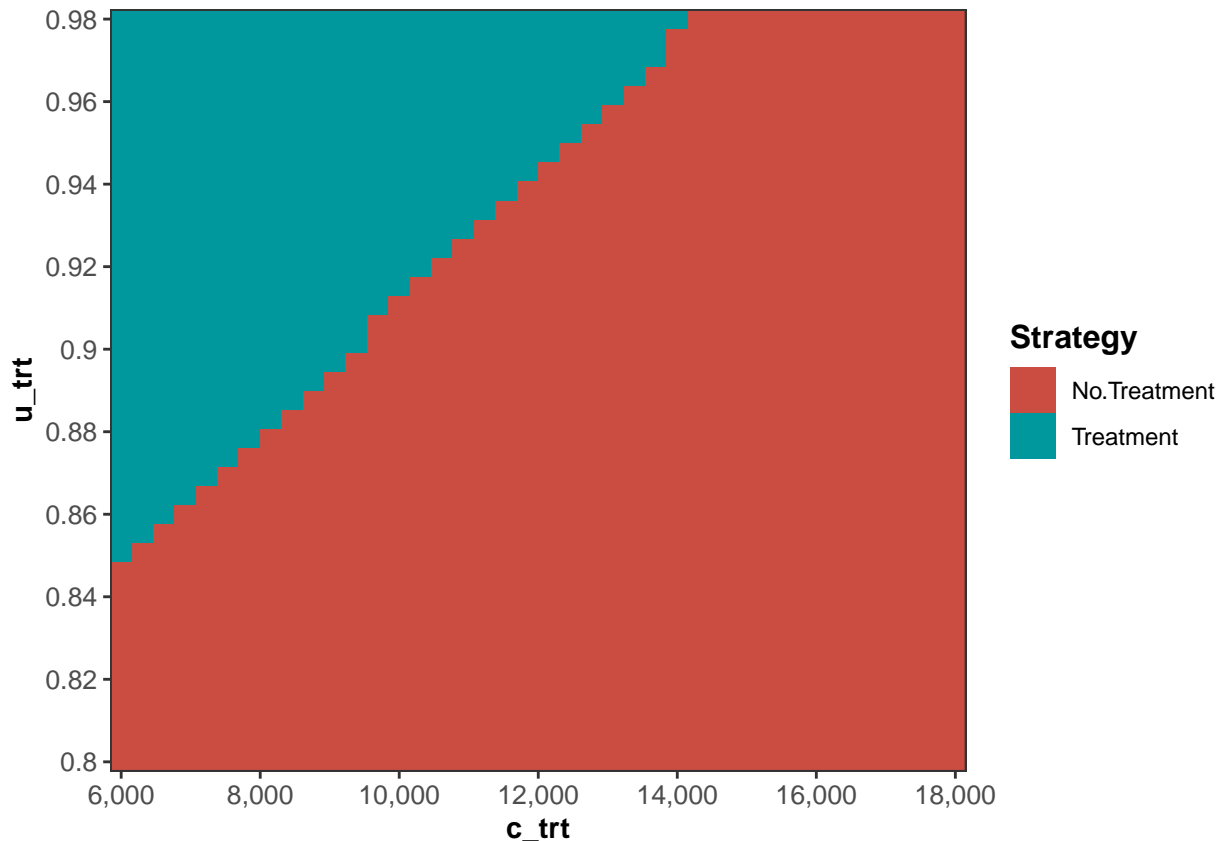


Figure 10: Two-way sensitivity analysis for the time-independent Sick-Sicker model.

09 Probabilistic Sensitivity Analysis (PSA)

```

# Function to generate PSA input dataset
gen_psa <- function(n_sim = 1000, seed = 071818){
  set.seed(seed) # set a seed to be able to reproduce the same results
  df_psa <- data.frame(
    # Transition probabilities (per cycle)
    p_HS1 = rbeta(n_sim, shape1 = 30, shape2 = 170), # probability to become sick when healthy
    p_S1H = rbeta(n_sim, shape1 = 60, shape2 = 60), # probability to become healthy when sick
    p_S1S2 = rbeta(n_sim, shape1 = 84, shape2 = 716), # probability to become sicker when sick
    p_HD = rbeta(n_sim, shape1 = 10, shape2 = 1990), # probability to die when healthy
    hr_S1 = rlnorm(n_sim, meanlog = log(3), sdlog = 0.01), # rate ratio of death in S1 vs healthy
  )
}

```

```

hr_S2 = rlnorm(n_sim, meanlog = log(10), sdlog = 0.02), # rate ratio of death in S2 vs healthy

# State rewards
# Costs
c_H = rgamma(n_sim, shape = 100, scale = 20), # cost of remaining one cycle in state H
c_S1 = rgamma(n_sim, shape = 177.8, scale = 22.5), # cost of remaining one cycle in state S1
c_S2 = rgamma(n_sim, shape = 225, scale = 66.7), # cost of remaining one cycle in state S2
c_Trt = rgamma(n_sim, shape = 73.5, scale = 163.3), # cost of treatment (per cycle)
c_D = 0, # cost of being in the death state

# Utilities
u_H = rbeta(n_sim, shape1 = 200, shape2 = 3), # utility when healthy
u_S1 = rbeta(n_sim, shape1 = 130, shape2 = 45), # utility when sick
u_S2 = rbeta(n_sim, shape1 = 230, shape2 = 230), # utility when sicker
u_D = 0, # utility when dead
u_Trt = rbeta(n_sim, shape1 = 300, shape2 = 15), # utility when being treated
d_e = 0.03, # discount factor for effectiveness
d_c = 0.03 # discount factor for costs
)
return(df_psa)
}
# Try it
gen_psa(10)

```

```

##      p_HS1    p_S1H    p_S1S2    p_HD    hr_S1    hr_S2    c_H
## 1  0.15132349 0.4820445 0.08485456 0.003973762 3.055410 10.148768 1887.449
## 2  0.12522484 0.4392546 0.11683397 0.004813328 3.005252 9.928044 1966.291
## 3  0.15673720 0.5311020 0.11925841 0.004491843 2.982392 10.021998 2101.059
## 4  0.11276858 0.4493088 0.09393848 0.010646056 2.981918 10.167486 1728.664
## 5  0.14292946 0.5825087 0.11628065 0.007045338 3.024975 10.006505 2160.895
## 6  0.20088601 0.4580278 0.10403415 0.004058637 2.971345 10.012715 1969.304
## 7  0.14347796 0.5492245 0.09294048 0.005658464 2.989181 10.001860 1641.192
## 8  0.13149270 0.4899896 0.12838861 0.007354969 2.984080 9.919332 2276.032
## 9  0.15593267 0.5705602 0.09237907 0.004162148 2.988974 10.142331 1717.867
## 10 0.09824029 0.5127637 0.10683148 0.002141140 2.968692 9.749869 2132.549
##      c_S1    c_S2    c_Trt c_D    u_H    u_S1    u_S2 u_D    u_Trt
## 1  4173.600 14607.88 10673.74  0 0.9829120 0.7252701 0.4757751  0 0.9435773
## 2  4188.264 14714.84 13380.61  0 0.9693926 0.7610392 0.4591563  0 0.9501528
## 3  4578.682 16664.54 12522.58  0 0.9910033 0.7076433 0.5032719  0 0.9670124
## 4  3395.922 15669.87 13833.65  0 0.9928844 0.7124119 0.5132492  0 0.9649128
## 5  3878.562 15302.70 13154.65  0 0.9748972 0.7344718 0.4895693  0 0.9325969
## 6  3970.522 15303.55 13914.95  0 0.9832065 0.7169406 0.4628316  0 0.9456881
## 7  3680.588 15378.20 12117.86  0 0.9917677 0.7851270 0.4504530  0 0.9093339
## 8  3781.956 15614.73 12070.99  0 0.9727485 0.7582961 0.4727426  0 0.9390084
## 9  3448.146 15401.50 11760.17  0 0.9836544 0.7648816 0.4851046  0 0.9537789
## 10 4656.666 15198.97 10216.98  0 0.9915345 0.7751203 0.5311969  0 0.9577708
##      d_e d_c
## 1  0.03 0.03
## 2  0.03 0.03
## 3  0.03 0.03
## 4  0.03 0.03
## 5  0.03 0.03
## 6  0.03 0.03
## 7  0.03 0.03

```



```

## 8 0.03 0.03
## 9 0.03 0.03
## 10 0.03 0.03

# Number of simulations
n_sim <- 1000

# Generate PSA input dataset
df_psa_input <- gen_psa(n_sim = n_sim)
# First six observations
head(df_psa_input)

##      p_HS1      p_S1H      p_S1S2      p_HD      hr_S1      hr_S2      c_H
## 1 0.1513235 0.5617628 0.1033817 0.006269864 2.962948 9.820157 1658.447
## 2 0.1252248 0.4380343 0.1152594 0.005371105 3.014068 10.100813 2050.281
## 3 0.1567372 0.5458102 0.1274505 0.004301015 3.018588 9.825985 2094.425
## 4 0.1127686 0.5106953 0.1015742 0.005319105 2.979709 9.697122 2135.411
## 5 0.1429295 0.5277308 0.1020384 0.004225209 3.025490 10.254387 1665.257
## 6 0.2008860 0.5252364 0.1030676 0.008319206 2.979175 10.366400 2049.794
##      c_S1      c_S2      c_Trtr c_D      u_H      u_S1      u_S2 u_D      u_Trtr
## 1 4191.109 15165.80 10292.050 0 0.9816602 0.7395884 0.5106656 0 0.9496993
## 2 4133.823 15042.66 10779.006 0 0.9906084 0.7240259 0.4927761 0 0.9731185
## 3 3567.033 15052.49 12811.443 0 0.9834960 0.6910869 0.5024542 0 0.9598494
## 4 3726.226 16084.60 10072.741 0 0.9919197 0.7065183 0.5118351 0 0.9357411
## 5 3654.486 15201.84 14917.228 0 0.9953850 0.6838064 0.5021483 0 0.9635901
## 6 3927.001 17060.57 9581.097 0 0.9813330 0.7766346 0.5035428 0 0.9506926
##      d_e d_c
## 1 0.03 0.03
## 2 0.03 0.03
## 3 0.03 0.03
## 4 0.03 0.03
## 5 0.03 0.03
## 6 0.03 0.03

# Histogram of parameters
ggplot(melt(df_psa_input, variable.name = "Parameter"), aes(x = value)) +
  facet_wrap(~Parameter, scales = "free") +
  geom_histogram(aes(y = ..density..)) +
  scale_x_continuous(breaks = scales::pretty_breaks(n = 3)) +
  theme_bw(base_size = 16) +
  theme(axis.text = element_text(size=6))

# Initialize dataframes with PSA output
# Dataframe of costs
df_c <- as.data.frame(matrix(0,
                             nrow = n_sim,
                             ncol = n_str))
colnames(df_c) <- v_names_str
# Dataframe of effectiveness
df_e <- as.data.frame(matrix(0,
                             nrow = n_sim,
                             ncol = n_str))
colnames(df_e) <- v_names_str

```

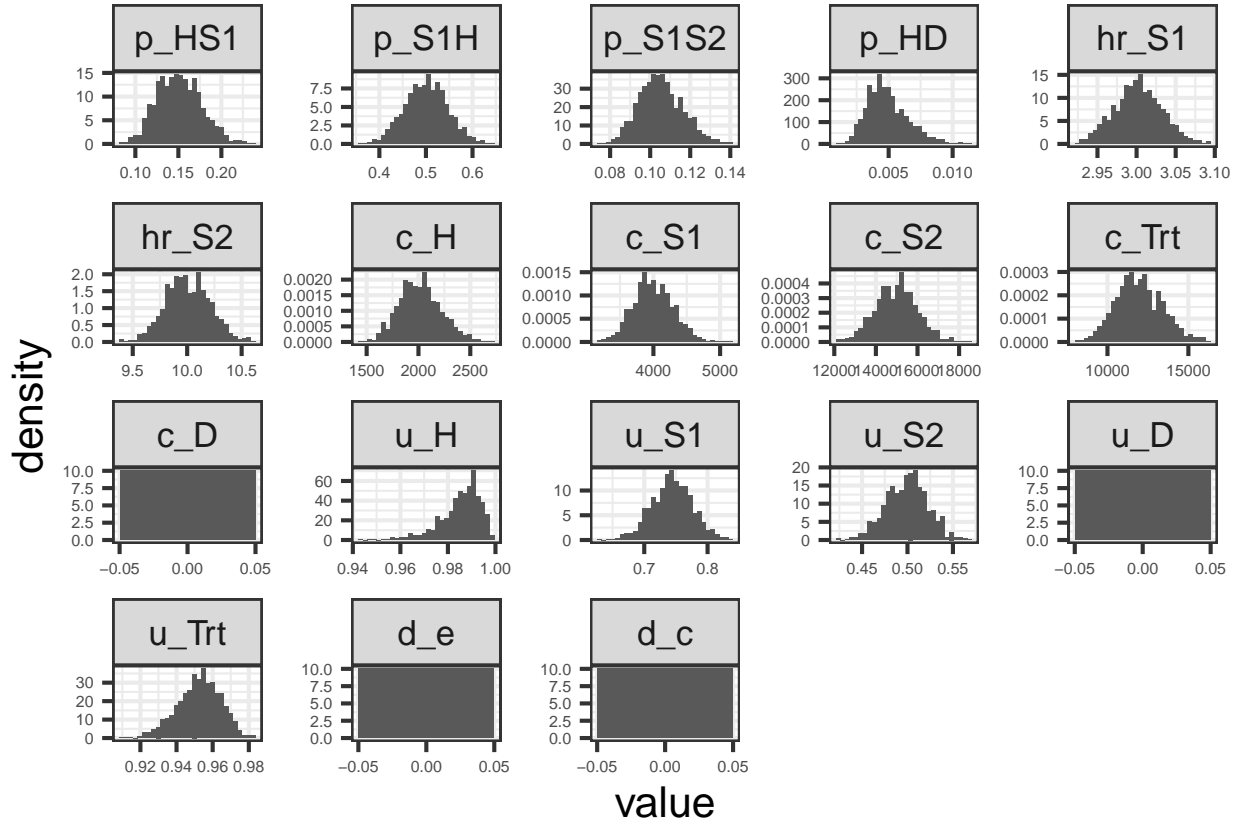


Figure 11: Histogram of parameters distributions for the time-independent Sick-Sicker model.

09.1 Conduct probabilistic sensitivity analysis

```
# Run Markov model on each parameter set of PSA input dataset
for(i in 1:n_sim){
  l_out_temp <- calculate_ce_out(df_psa_input[i, ])
  df_c[i, ] <- l_out_temp$Cost
  df_e[i, ] <- l_out_temp$Effect
  # Display simulation progress
  if(i/(n_sim/10) == round(i/(n_sim/10), 0)) { # display progress every 10%
    cat('\r', paste(i/n_sim * 100, "% done", sep = " "))
  }
}
```

```
## 10 % done 20 % done 30 % done 40 % done 50 % done 60 % done 70 % done 80 % done 90 % done 100 % done
```

09.2 Create PSA object for dampack

```
l_psa <- make_psa_obj(cost      = df_c,
                     effectiveness = df_e,
                     parameters  = df_psa_input,
                     strategies   = v_names_str)
l_psa$strategies <- v_names_str
colnames(l_psa$effectiveness)<- v_names_str
colnames(l_psa$cost)<- v_names_str
```

09.2.1 Save PSA objects

```
save(df_psa_input, df_c, df_e, v_names_str, n_str, l_psa,
     file = "markov_sick-sicker_PSA_dataset.RData")
```

09.3 Create probabilistic analysis graphs

```
load(file = "markov_sick-sicker_PSA_dataset.RData")
```

Vector with willingness-to-pay (WTP) thresholds.

```
v_wtp <- seq(0, 200000, by = 10000)
```

09.3.1 Cost-Effectiveness scatter plot

09.4 Conduct CEA with probabilistic output

```
# Compute expected costs and effects for each strategy from the PSA
df_out_ce_psa <- summary(l_psa)
```

```
# Calculate incremental cost-effectiveness ratios (ICERs)
df_cea_psa <- calculate_icers(cost      = df_out_ce_psa$meanCost,
                             effect     = df_out_ce_psa$meanEffect,
                             strategies = df_out_ce_psa$Strategy)
df_cea_psa
```

##	Strategy	Cost	Effect	Inc_Cost	Inc_Effect	ICER	Status
## 1	No Treatment	75955.61	15.64787	NA	NA	NA	ND
## 2	Treatment	143251.54	16.29156	67295.93	0.6436874	104547.5	ND

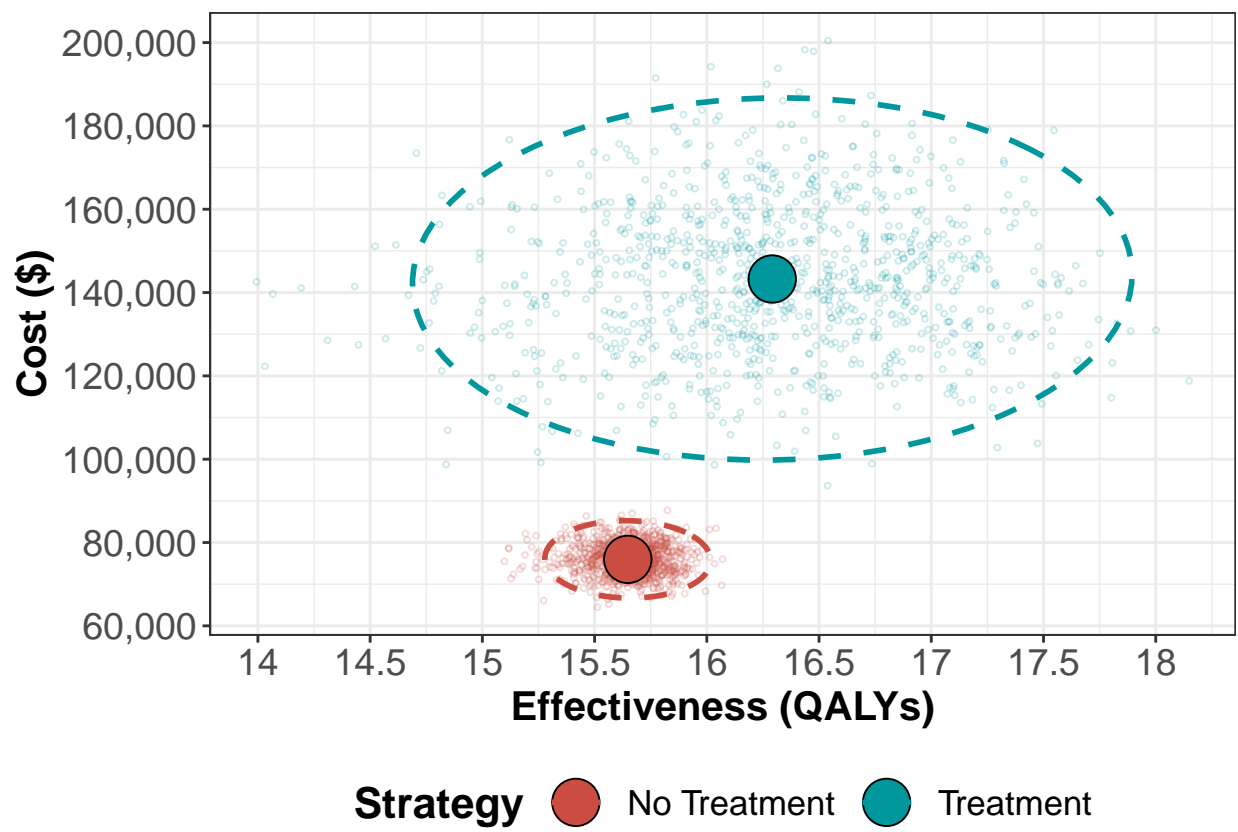


Figure 12: Cost-effectiveness scatter plot.

```

# Save CEA table with ICERs
# As .RData
save(df_cea_psa,
     file = "markov_sick-sicker_probabilistic_CEA_results.RData")
# As .csv
write.csv(df_cea_psa,
         file = "markov_sick-sicker_probabilistic_CEA_results.csv")

```

09.4.1 Plot cost-effectiveness frontier

```
plot(df_cea_psa)
```

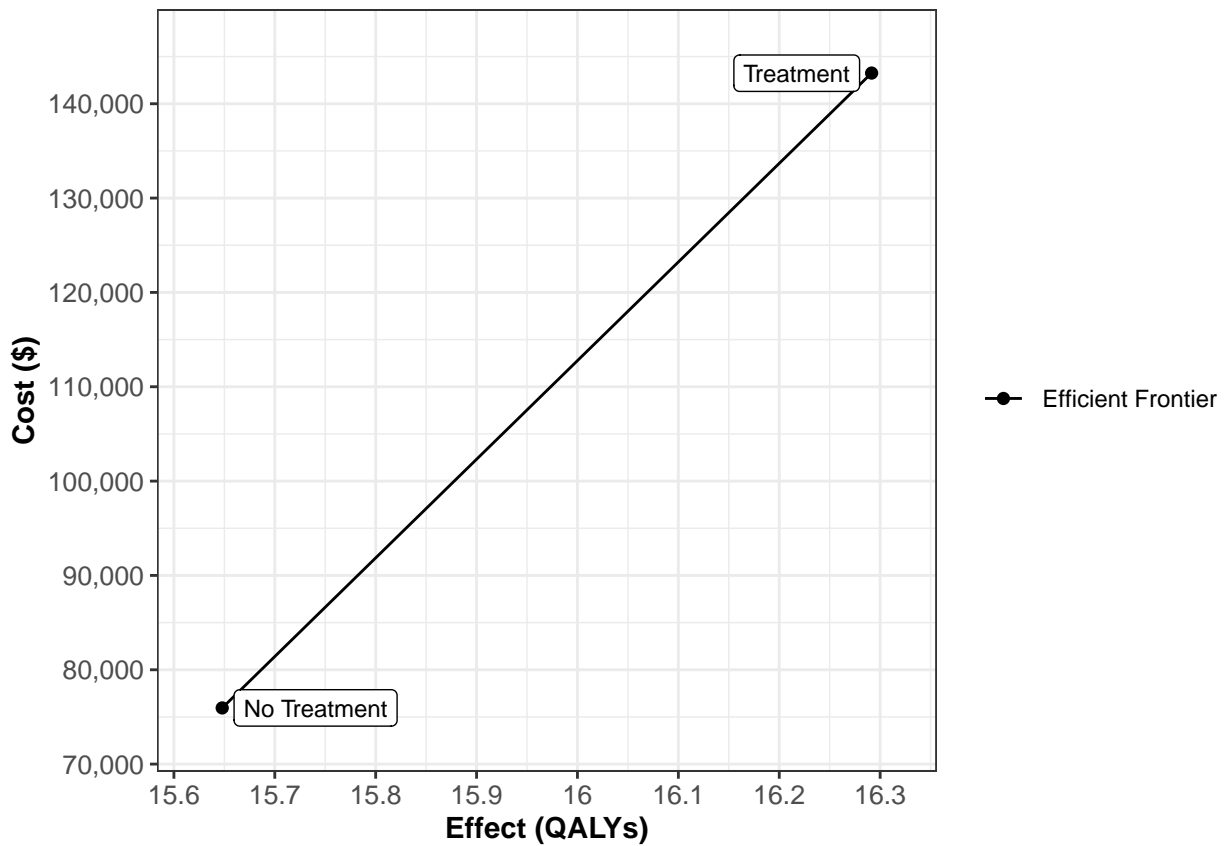


Figure 13: Cost-effectiveness efficient frontier from probabilistic outputs for the time-independent Sick-Sicker model.

09.4.2 Cost-effectiveness acceptability curves (CEACs) and frontier (CEAF)

```

##   range_min range_max cost_eff_strat
## 1      0      110000   No Treatment
## 2    110000    200000     Treatment

```

09.4.3 Expected Loss Curves (ELCs)

The expected loss is the the quantification of the foregone benefits when choosing a suboptimal strategy given current evidence.

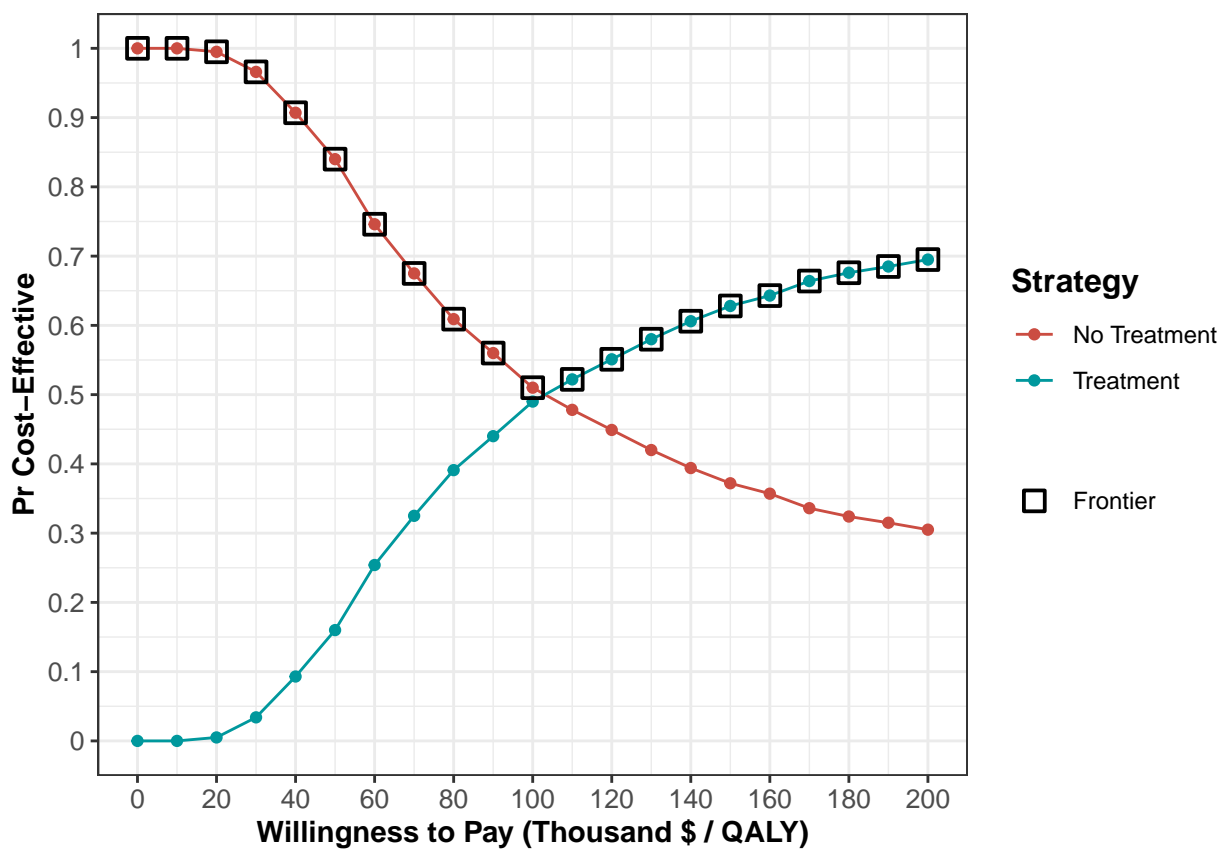


Figure 14: Cost-effectiveness acceptability curves (CEACs) and frontier (CEAF).

##	WTP	Strategy	Expected_Loss	On_Frontier
## 1	0	No Treatment	0.00000	TRUE
## 2	0	Treatment	67295.92714	FALSE
## 3	10000	No Treatment	0.00000	TRUE
## 4	10000	Treatment	60859.05317	FALSE
## 5	20000	No Treatment	35.95817	TRUE
## 6	20000	Treatment	54458.13738	FALSE
## 7	30000	No Treatment	322.38988	TRUE
## 8	30000	Treatment	48307.69512	FALSE
## 9	40000	No Treatment	1357.10595	TRUE
## 10	40000	Treatment	42905.53722	FALSE
## 11	50000	No Treatment	3297.41004	TRUE
## 12	50000	Treatment	38408.96734	FALSE
## 13	60000	No Treatment	6248.02097	TRUE
## 14	60000	Treatment	34922.70430	FALSE
## 15	70000	No Treatment	10123.60585	TRUE
## 16	70000	Treatment	32361.41521	FALSE
## 17	80000	No Treatment	14668.48694	TRUE
## 18	80000	Treatment	30469.42234	FALSE
## 19	90000	No Treatment	19729.40367	TRUE
## 20	90000	Treatment	29093.46510	FALSE
## 21	100000	No Treatment	25199.14250	TRUE
## 22	100000	Treatment	28126.32997	FALSE
## 23	110000	No Treatment	30950.62051	FALSE
## 24	110000	Treatment	27440.93400	TRUE
## 25	120000	No Treatment	36908.01787	FALSE
## 26	120000	Treatment	26961.45740	TRUE
## 27	130000	No Treatment	43029.50480	FALSE
## 28	130000	Treatment	26646.07036	TRUE
## 29	140000	No Treatment	49318.83174	FALSE
## 30	140000	Treatment	26498.52333	TRUE
## 31	150000	No Treatment	55747.56036	FALSE
## 32	150000	Treatment	26490.37798	TRUE
## 33	160000	No Treatment	62263.42425	FALSE
## 34	160000	Treatment	26569.36791	TRUE
## 35	170000	No Treatment	68869.16074	FALSE
## 36	170000	Treatment	26738.23043	TRUE
## 37	180000	No Treatment	75547.14077	FALSE
## 38	180000	Treatment	26979.33649	TRUE
## 39	190000	No Treatment	82267.39582	FALSE
## 40	190000	Treatment	27262.71757	TRUE
## 41	200000	No Treatment	89028.28011	FALSE
## 42	200000	Treatment	27586.72789	TRUE

09.4.4 Expected value of perfect information (EVPI)

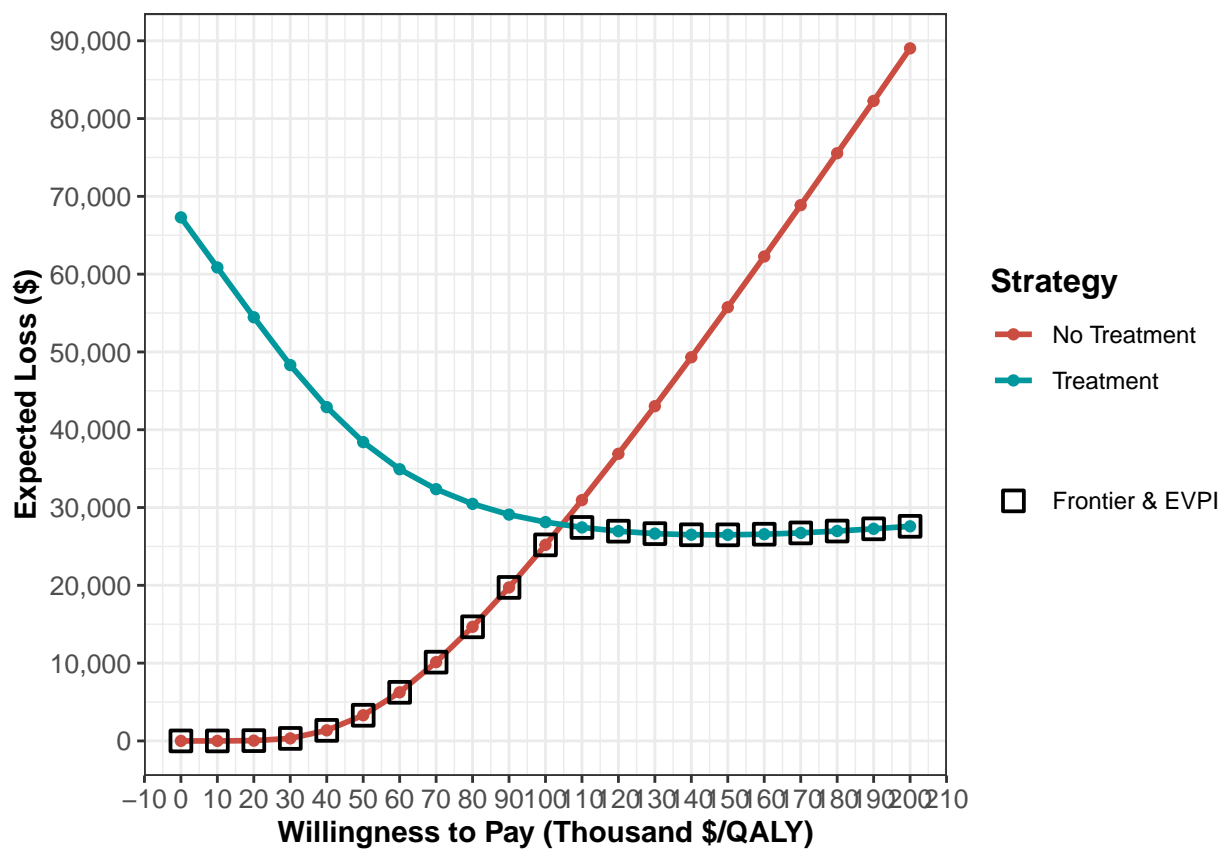


Figure 15: Expected loss curves (ELCs).

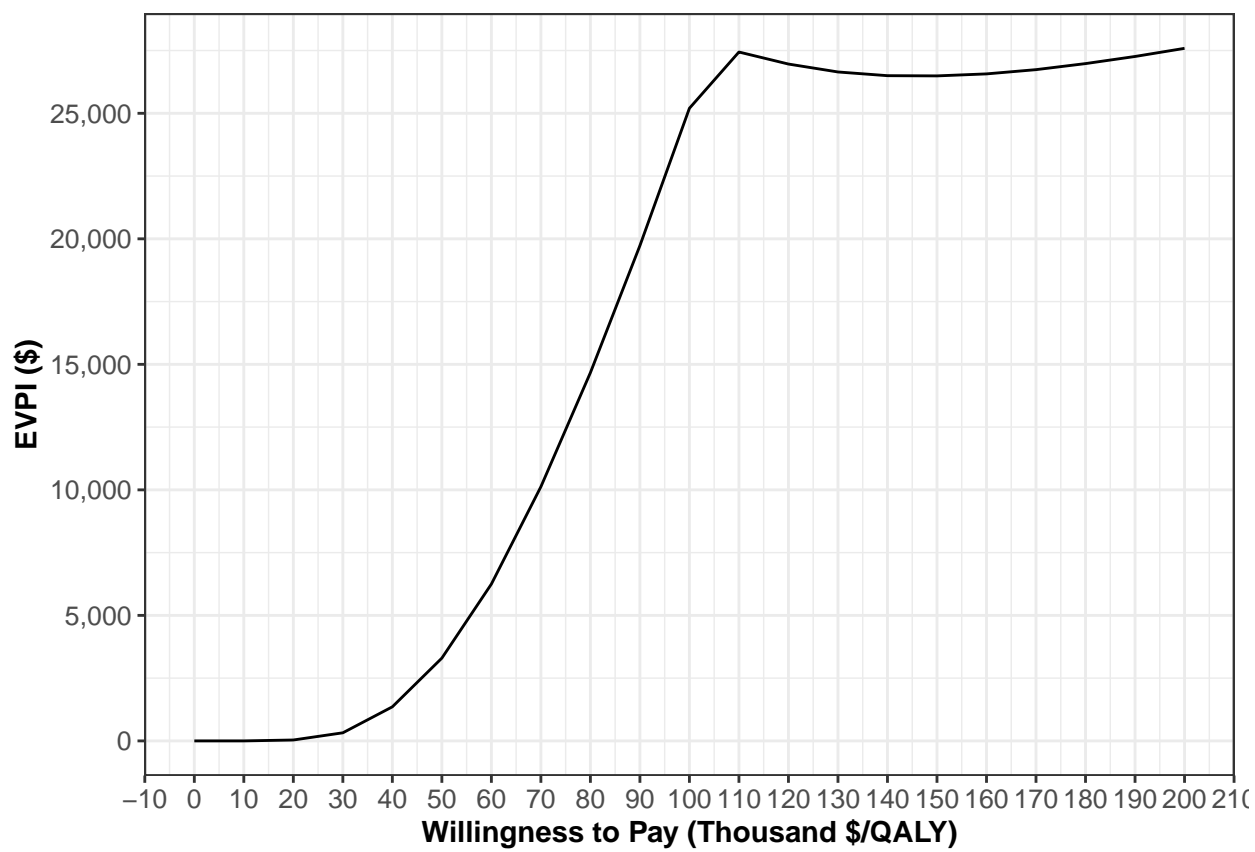


Figure 16: Expected value of perfect information (EVPI).