# Report for the Markov model (cohort state-transition model)

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### Results from the paper

These two tables are from the result part of the paper by Yaqin Si.

Table 1: Increased QALY with no screening

	est	LB	UB
strategy1	498	103	894
strategy2	691	233	194
strategy3	654	105	1108

 Table 2: Prevent CVD events

 est
 LB
 UB

 strategy1
 298
 155
 441

 strategy2
 374
 181
 567

 strategy3
 346
 154
 538

## Part I. General population (without CVD)

#### Markov model

% latex table generated in R 3.6.3 by x table 1.8-4 package % Sat Dec 26 18:59:02 2020

Table 3: Data from Global Health Data Exchange

	Index	sex	$rate\_incidence\_CVD$	$rate\_death\_CVD$	$rate\_death\_nonCVD$
1	40	male	0.003888	0.000819	0.002494
2	45	$_{\mathrm{male}}$	0.006729	0.001340	0.003399
3	50	$_{\mathrm{male}}$	0.010564	0.002302	0.004951
4	55	male	0.015291	0.003665	0.007282
5	60	male	0.022078	0.006404	0.011159
6	65	male	0.030980	0.011155	0.016946
7	70	male	0.043589	0.019978	0.026305
8	40	female	0.004545	0.000351	0.001137
9	45	female	0.007094	0.000643	0.001620
10	50	female	0.010133	0.001206	0.002475
11	55	female	0.013734	0.002014	0.003705
12	60	female	0.018272	0.003872	0.005850
13	65	female	0.023744	0.006996	0.009060
_14	70	female	0.033907	0.013398	0.014907

```
## General setup
source("./function/transform_func.R")
rate_data <- rate_data[1:7,]</pre>
n t <- 10 # time horizon, number of cycles
# S1: live; S2: cvd; S3: cvdth; S4: oth death
v_names_states <- c("S1", "S2", "S3", "S4")</pre>
n_states <- length(v_names_states) # number of health states</pre>
v_names_str <- c("Strategy0", "Strategy1", "Strategy2", "Strategy3") # store the strategy names
           <- length(v names str)</pre>
                                         # number of strategies
# Utilities: for calculation of QALY
out_cvd_free <- 1  # utility when being S1</pre>
out_cvd <- 0.9 # utility when being S2</pre>
out_dth <- 0 # utility when being S3 and S4 together
out_trans_to_cvd <- -0.038
                             # TODO
p_live_oth_death <- rate_to_prob(r=rate_data$rate_death_nonCVD,t = 1)</pre>
p_live_cvd <- rate_to_prob(r=rate_data$rate_incidence_CVD,t = 1)</pre>
p_live_cvdth <- rate_to_prob(r=rate_data$rate_death_CVD,t = 1)</pre>
# transition probability from S2 to S3
p ccvd acvd <- rate to prob(rate data$rate incidence CVD*HR cvdhistory cvd,t=5)
p_ccvd_cvdth <- rate_to_prob(rate_data$rate_death_CVD*HR_cvdhistory_cvdth,t=5)</pre>
```

#### Component 1: A transition probability matrix $P_t$

$$P_t = \begin{cases} p_{[1,1,t]} & p_{[1,2,t]} & p_{[1,3,t]} & p_{[1,4,t]} \\ p_{[2,1,t]} & p_{[2,2,t]} & p_{[2,3,t]} & p_{[2,4,t]} \\ p_{[3,1,t]} & p_{[3,2,t]} & p_{[3,3,t]} & p_{[3,4,t]} \\ p_{[4,1,t]} & p_{[4,2,t]} & p_{[4,3,t]} & p_{[4,4,t]} \end{cases}$$

For example, the transition matrix for the cohort is:

$$P_1 = \begin{cases} p_{[1,1,t]} & p\_live\_cvd & p\_live\_cvdth & p\_live\_oth\_death \\ 0 & p\_ccvd\_acvd & p\_ccvd\_cvdth & p\_live\_oth\_death \\ 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 \end{cases}$$

```
#### Create transition arrays ####
a_P \leftarrow array(0, dim = c(n_states, n_states, n_t),
               dimnames = list(v_names_states, v_names_states, 0:(n_t - 1)))
## From S1
a_P["S1", "S1", 0:6] <- (1-(p_live_cvd[1] +
                           p_live_cvdth[1] +
                           p_live_oth_death[1]))
a_P["S1", "S2", 0:6] <- p_live_cvd[1]
a_P["S1", "S3", 0:6] <- p_live_cvdth[1]
a_P["S1", "S4", 0:6] <- p_live_oth_death[1]</pre>
## From S2
a_P["S2", "S1", 0:6] <- 0
a_P["S2", "S2", 0:6] <- p_ccvd_acvd[1]
a_P["S2", "S3", 0:6] <- p_ccvd_cvdth[1]
# TODO: cvd people to death of other reasons (P3)
a_P["S2", "S4", 0:6] <- 1 - (p_ccvd_acvd[1] + p_ccvd_cvdth[1])
## From S3
a_P["S3", "S3", 0:6] <- 1
## From S4
a_P["S4", "S4", 0:6] <- 1
# From S1
a_P["S1", "S1", 6:10] <- (1-(p_live_cvd[1+1] +
                           p live cvdth[1+1] +
                           p_live_oth_death[1+1]))
a_P["S1", "S2", 6:10] <- p_live_cvd[1+1]
a_P["S1", "S3", 6:10] <- p_live_cvdth[1+1]
a_P["S1", "S4", 6:10] <- p_live_oth_death[1+1]
## From S2
a_P["S2", "S1", 6:10] <- 0
a_P["S2", "S2", 6:10] <- p_ccvd_acvd[1+1]
a_P["S2", "S3", 6:10] <- p_ccvd_cvdth[1+1]
# TODO: cvd people to death of other reasons
# p_live_oth_death != p_cvd _oth_death (P3)
a_P["S2", "S4", 6:10] <-1 - (p_ccvd_acvd[1+1] + p_ccvd_cvdth[1+1])
## From S3
a_P["S3", "S3", 6:10] <- 1
## From S4
a_P["S4", "S4", 6:10] <- 1
```

### Component 2: A cohort trace matrix M

```
#### Run Markov model ####
## Initial state vector
# All starting healthy
v_s_{init} \leftarrow c(state0 = 1, state1 = 0, state2 = 0, state3 = 0)
## Initialize cohort trace for cSTM
m_M <- matrix(0,nrow</pre>
                         = (n_t + 1), ncol = n_states,
              dimnames = list(0:n_t, v_names_states))
m_M[1, ] <- v_s_init</pre>
## Iterative solution of cSTM
for(t in 1:n_t){
  ## Fill in cohort trace
 m_M[t + 1, ]
                                      %*% a_P[, , t]
                     <- m_M[t, ]
plot_trace(m_M)
```

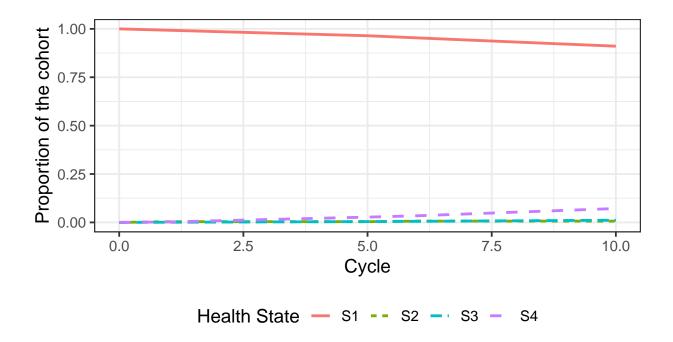


Figure 1: For the group male aging from 40-50

	Ta	able 4: Incedence rate	
Item		CVD incidence(HR)	CVD cause-specific mortality (HR)
Strategy 1	Low risk	0.63	1
	Medium risk	1.56	1
	High risk	1.6	1.7
Strategy 2	Low risk	0.43	1
	Medium risk	0.97	1
	High risk	2.06	1.7
Strategy 3	Low risk	0.63	1
	Medium risk	1.09	1
	High risk	2.11	1.7
Intervention	Weight control	0.93	0.93
	Smoke cession	0.85	0.72
	Salt reduction	0.81	0.66
Medication	Statin and antihypertensiv	re	

```
HR_l_stg1 <- 0.63
HR_m_stg1
           <- 1.56
HR_h_stg1
           <- 1.6
HR 1 stg2
           <- 0.43
HR_m_stg2
           <- 0.97
HR_h_stg2
           <- 2.06
HR_1_stg3
           <- 0.45
HR_m_stg3
           <- 1.09
HR_h_stg3
           <- 2.11
HR_smk_cvd <- 0.85
HR_smk_cvdth <- 0.72
HR_salt_cvd <- 0.81
HR_salt_cvdth <- 0.66</pre>
HR_wtc_cvd <- 0.93
HR_wtc_dth <- 0.93
HR_hpt_lip_cvd <- 0.7</pre>
HR_hpt_lip_cvdth <- 0.82</pre>
# transition probability from S1 to S2
p_live_cvd_l <- ProbFactor(p_live_cvd, HR_l_stg1)</pre>
p_live_cvd_m <- ProbFactor(p_live_cvd, HR_m_stg1)</pre>
p_live_cvd_h <- ProbFactor(p_live_cvd, HR_h_stg1)</pre>
# transition probability from S1 to S3
p_live_cvdth_l <- ProbFactor(p_live_cvdth,1) # equal</pre>
# p_live_cvdth_m <- ProbFactor(p_live_cvdth,1.7) # intervention etc.</pre>
# p_live_cvdth_h <- ProbFactor(p_live_cvdth,1) # intervention etc.</pre>
```