Multistate models

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August 2019

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Multistate models

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These assumptions are true for death and many chronic diseases.

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- ▶ Aim of the modeling the transition rates between states, is to be able predict how population moves between states:
- state occupancy probabilities
- visit probability
- length of stay (sojourn time)

Multistate models (ms-Markov)

Generalization of Poisson regression to multiple disease states:

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- ▶ In the clinical literature, the term "Markov model" is often used about any type of multistate model
- ...and the Markov property is handy in probability theory

Multistate models (ms-Markov) 4/39

▶ Define the (disease) states

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- Not a trivial task do we want e.g.
 - cause of death
 - disease status at death

Times should be recorded as dates

birth date

- birth date
- entry date

- birth date
- entry date
- entry state

- birth date
- entry date
- entry state
- exit date

- birth date
- entry date
- entry state
- exit date
- death date

- birth date
- entry date
- entry state
- exit date
- death date
- state entry dates for all states

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From this each person's trajectory through states can be constructed

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$$d_{psv}\log(\lambda_{psv}) - \lambda_{psv}y_{ps},$$
 where:

- ullet λ_{psv} rate for person p in state s going to state v
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- Total log-lik is sum of terms over persons and transitions

Multistate models (ms-Markov) 7/39

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- components **not** independent, but the total likelihood is a product; hence of the same form as the likelihood of independent Poisson variates
- practical analysis is just analysis of each transition rate separately
- as long as no two rates out of the same state are modeled we can use subsets of Lexis objects

Multistate models (ms-Markov)

Multistate models with Lexis

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Example: Renal failure data from Steno

Hovind P, Tarnow L, Rossing P, Carstensen B, and Parving H-H: Improved survival in patients obtaining remission of nephrotic range albuminuria in diabetic nephropathy. *Kidney Int.*, 66(3):1180–1186, 2004.

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- ▶ 96 patients entering at nephrotic range albuminuria (NRA), i.e. U-alb> 300mg/day.
- ► Is remission from this condition (i.e return to U-alb< 300mg/day) predictive of the prognosis?
- ► Endpoint of interest: Death or end stage renal disease (ESRD), i.e. dialysis or kidney transplant.

		Remission	
	Total	Yes	No
No. patients No. events Follow-up time (years)	125 77 1084.7	32 8 259.9	93 69 824.8
Cox-model: Timescale: Time since nephrotic range albuminuria (NRA) Entry: 2.5 years of GFR-measurements after NRA Outcome: ESRD or Death			
Estimates:	RR	95% c.i.	p
Fixed covariates: Sex (F vs. M): Age at NRA (per 10 years):	0.92 1.42	(0.53,1.57) (1.08,1.87)	0.740 0.011
Time-dependent covariate: Obtained remission:	0.28	(0.13,0.59)	0.001

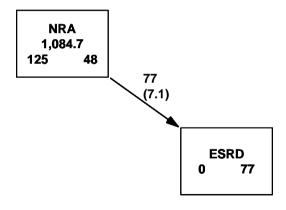
Features of the analysis

- Remission is included as a time-dependent variable.
- Age at entry is included as a fixed variable.

```
renal[1:5,]
id dob doe dor dox event
17 1967.944 1996.013 NA 1997.094 2
26 1959.306 1989.535 1989.814 1996.136 1
27 1962.014 1987.846 NA 1993.239 3
33 1950.747 1995.243 1995.717 2003.993 0
42 1961.296 1987.884 1996.650 2003.955 0
```

Note patient 26, 33 and 42 obtain remission.

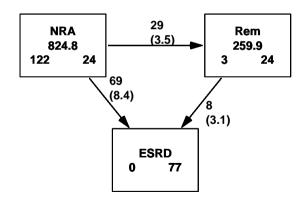
```
> Lr <- Lexis( entry = list( per=doe,
                             age=doe-dob,
+
                              tfi=0),
                exit = list( per=dox ),
         exit.status = event>0,
              states = c("NRA", "ESRD"),
                data = renal )
> summary( Lr )
Transitions:
     То
From
      NRA ESRD
               Records: Events: Risk time:
                                               Persons:
  NRA 48
            77
                     125
                                77
                                      1084.67
                                                    125
```

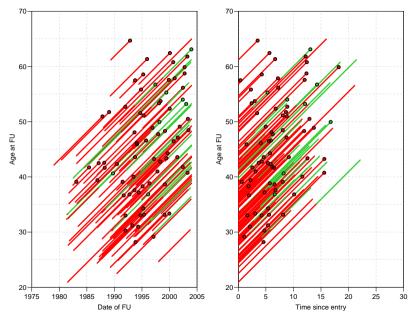


Cutting follow-up at remission: cutLexis

```
> Lc <- cutLexis( Lr, cut=Lr$dor,
+
                timescale="per",
                new.state="Rem",
         precursor.states="NRA" )
> summary( Lc )
Transitions:
     To
From
      NRA Rem ESRD
                    Records:
                              Events: Risk time:
                                                    Persons:
  NR.A
       24
           29
                69
                          122
                                    98
                                           824.77
                                                         122
           24
                           32
                                           259.90
                                                          32
  R.em
           53
                77
  Sum
       24
                          154
                                   106
                                          1084.67
                                                         125
```

Showing states and FU: boxes.Lexis





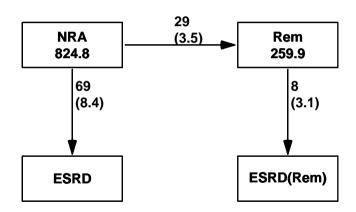
Splitting states: cutLexis

```
> Lc <- cutLexis( Lr, cut=Lr$dor,
                 timescale="per",
+
                 new.state="Rem",
         precursor.states="NRA",
              split.states=TRUE )
> summary( Lc )
Transitions:
     Tο
From
      NRA Rem ESRD ESRD(Rem)
                                Records: Events: Risk time:
                                                                Persons:
  NR.A
       24
           29
                 69
                                     122
                                                98
                                                                      122
                                                       824.77
                                      32
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                                                                      32
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           24
                                                 8
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Showing states and FU: boxes.Lexis



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- ▶ **Risk time** is the risk time in the "From" state
- **Events** are transitions to the "To" state
- All other transitions out of "From" are treated as censorings
- Possible to fit models separately for each transition

Prediction in multistate models: simLexis and renal failure

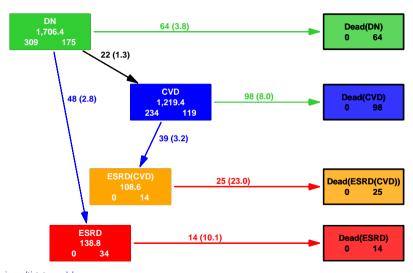
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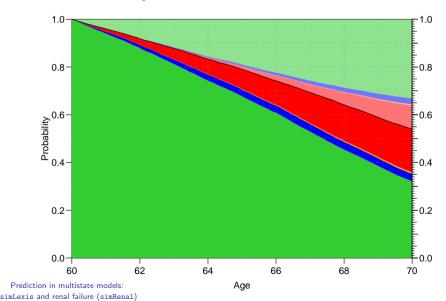
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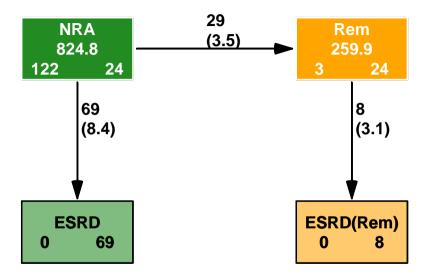
August 2019

A more complicated multistate model



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Modeling in a multistate model

Modeling in a multistate model

Each transition modeled by a model for rates (Cox-model, Poisson-model for split data, glm or gam):

... using the Lexis properties

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```
> # Rem-rate
> mr <- gam.Lexis( sLc, from="NRA", to="Rem",
                        formula = ~s(tfi. k=10) + sex)
mgcv::gam Poisson analysis of Lexis object sLc with log link:
Rates for the transition: NRA->Rem
> # ESRD-rates
> mx <- gam.Lexis( sLc. formula = ~ s(tfi,k=10) + sex +
                        I((doe-dob-40)/10) + I(lex.Cst=="Rem"))
+
mgcv::gam Poisson analysis of Lexis object sLc with log link:
Rates for transitions: NRA->ESRD, Rem->ESRD(Rem)
```

Default is to model all transitions

How do we get from rates (Poisson-models) to probabilities:

1 Analytic calculations:

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 - conceptually simple
 - computationally not quite simple
 - easy to generalize
 - hard to get confidence intervals (bootstrap)

Simulation of a survival time

▶ For a rate function $\lambda(t)$, $\Lambda(t) = \int_0^t \lambda(s) \, \mathrm{d}s$:

$$S(t) = \exp(-\Lambda(t))$$

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$$u = S(t) \Leftrightarrow \Lambda(t) = -\log(u)$$

Simulation of a survival time

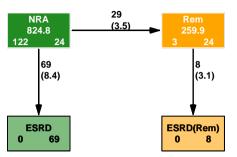
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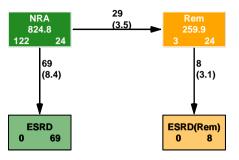
$$S(t) = \exp(-\Lambda(t))$$

▶ Simulate a survival probability $u \in [0, 1]$:

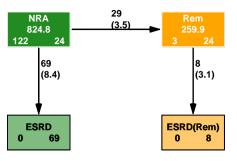
$$u = S(t) \Leftrightarrow \Lambda(t) = -\log(u)$$

- \blacktriangleright Knowledge of $\Lambda(t)$ makes it easy to find a survival time — essentially just linear interpolation.

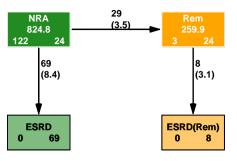




▶ Simulate a "survival time" for each transition **out** of a state.

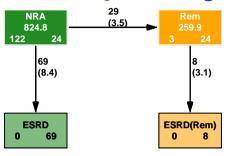


- ▶ Simulate a "survival time" for each transition **out** of a state.
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- ▶ Simulate a "survival time" for each transition **out** of a state.
- ▶ The smallest of these is the transition time.
- Choose the corresponding transition type as transition.

Transition objects are glm/gam



```
> Tr <- list( "NRA" = list( "ESRD" = mx,
+ "Rem" = mr),
+ "Rem" = list( "ESRD(Rem)" = mx))
```

Input required:

► A Lexis object representing the initial state of the persons to be simulated.

(lex.dur and lex.Xst will be ignored.)

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Output produced:

- A Lexis object with simulated event histories for may persons
- Use nState to count how many persons in each state at different times

Using simLexis I

Put one record a new Lexis object (init, say). representing a person with the desired covariates.

Must have same structure as the one used for estimation — time scales must be initiated even if not used in models

```
> init <- sLc[NULL,c(timeScales(sLc),"lex.Cst")]
> init[1,"per"] <- 1994
> init[1,"age"] <- 40
> init[1,"tfi"] <- 0
> init[1,"lex.Cst"] <- "NRA"
> init[1,"sex"] <- "M"
> init[1,"dob"] <- 1954
> init[1,"doe"] <- 1994
> init
```

Using simLexis II

```
per age tfi lex.Cst sex dob doe
1 1994 40
                   NR.A
           0
                         M 1954 1994
> system.time(
+ sim1 <- simLexis( Tr, init, N=10000 ) )
        system elapsed
  user
 48.655 52.448 36.065
> summary(sim1)
Transitions:
     To
From
          Rem ESRD ESRD(Rem) Records:
                                         Events: Risk time:
                                                             Persons:
  NRA 293 1797 7910
                                  10000
                                            9707
                                                   74766.04
                                                                10000
          874
                          923
                                   1797
                                             923 19852.42
                                                                 1797
  R.em
  Sum 293 2671 7910
                          923
                                  11797
                                           10630 94618.45
                                                                10000
```

Using a simulated Lexis object — pState I

```
> NN < - nState(sim1, at = seq(0,15,0.1),
                  from = 0,
            time.scale = "tfi" )
> head( NN )
    State
when
       NR.A
            Rem
                ESRD ESRD(Rem)
     10000
 0.1 9955 20 25
 0.2 9889 46 65
 0.3 9837 75 88
 0.4 9779 104 117
 0.5 9732
            127
                  140
> nw1 <- pState(NN, perm = c(1,2,4,3))
> head( nw1. 3 )
```

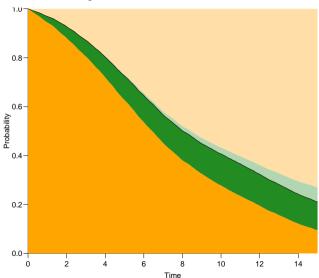
Using a simulated Lexis object — pState II

```
State
when
        NRA Rem ESRD(Rem) ESRD
     1.0000 1.0000
                  1.0000
 0.1 0.9955 0.9975 0.9975 1
 0.2 0.9889 0.9935 0.9935
> tail( nw1, 3 )
     State
when
         NR.A
             Rem ESRD(Rem) ESRD
 14.8 0.1004 0.2165 0.2751
 14.9 0.0969 0.2128 0.2717
 15 0.0953 0.2101 0.2701
> par(mar=c(3,3,0.1,0.1), mgp=c(3,1,0)/1.6, las=1)
> plot(nw1, col=clr[c(2.1.4.3)])
> lines( as.numeric(rownames(nw1)), nw1[,2] )
```

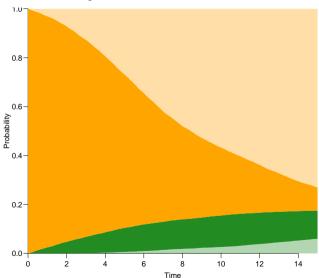
Using a simulated Lexis object — pState III

```
> nw2 < - pState(NN, perm = c(4,2,1,3))
> head( nw2, 3 )
    State
when ESRD(Rem) Rem NRA ESRD
            0 0.0000 1.0000
         0 0.0020 0.9975
 0.2
            0 0.0046 0.9935
> tail( nw2, 3 )
     State
when
      ESRD(Rem) Rem NRA ESRD
 14.8 0.0586 0.1747 0.2751
 14.9 0.0589 0.1748 0.2717
 15 0.0600 0.1748 0.2701
> par(mar=c(3,3,0.1,0.1), mgp=c(3,1,0)/1.6, las=1)
> plot(nw2, col=clr[c(4,1.2.3)])
```

Simulated probabilities



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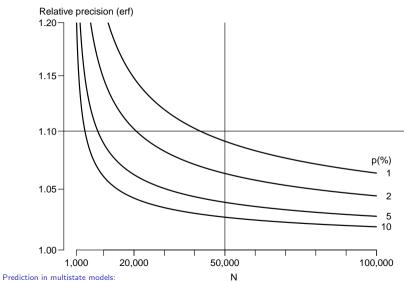
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▶ So c.i. of the form $p \stackrel{\times}{\div} \operatorname{erf}$ where:

$$erf = \exp(1.96 \times (1-p)/\sqrt{Np(1-p)})$$

Precision of simulated probabilities



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- Simulation allows estimation of cumulative probabilities:
 - Estimate transition rates (as usual)
 - Simulate probabilities (not as usual)