Using Joint Models to Estimate Causal Effects for Salvage Therapy after Prostatectomy

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Aims, Models & Estimands

1 Background & Aim



- Setting Patients treated with surgery after diagnosis of Prostate Cancer (PCa)
 - > remain at risk of metastasis
- Follow-up
 - > PSA levels at frequent intervals

 - > ST androgen deprivation therapy, radiation therapy, chemotherapy, and combinations



- Important questions regarding Salvage Therapy

 - ▶ when to start?
 - ▷ does it work?



Quantify the amount by which Salvage Therapy reduces the risk of metastasis



University of Michigan Prostatectomy Data

- → 3634 PCa patients followed-up in 1996–2013
 - * aged 40 to 84 years with clinically localized cT1 to cT3 disease
 - * received radical prostatectomy
- baseline variables: PSA, Gleason, T-stage, age, race, gland volume, perineural invasion, planned adjuvant therapy



Challenges

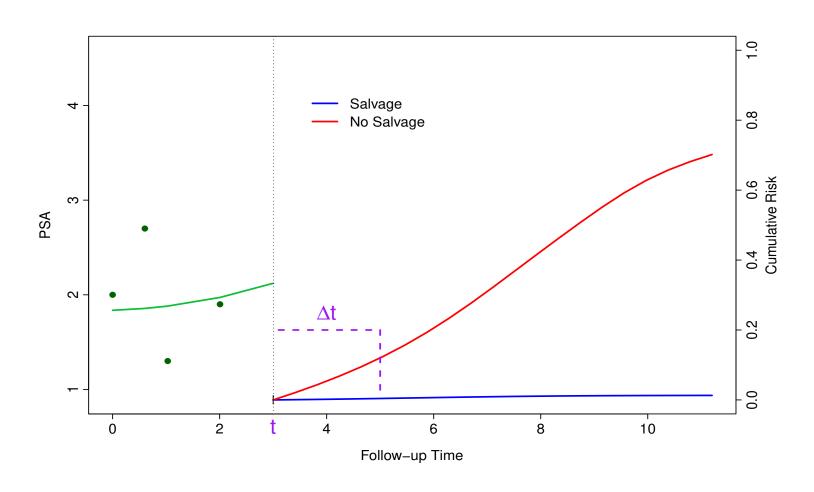
- ▷ Observational Data no RCT
 - * selection bias
 - * ascertainment bias
- ▷ Time-Varying Salvage Therapy
 - * depends on previous PSA
 - * PSA time-dependent confounder
 - * endogeneity

2 Causal ST Effects



- Standard assumptions for Causal Inference
 - Consistency: Observed outcomes equal the counterfactual outcomes for the actually assigned treatment
 - ▷ Sequential Exchangeability: The counterfactual outcomes are independent of the assigned treatment conditionally on the history of PSA measurements and baseline covariates







Which is the target group?

Notation

 $\triangleright T_m$: time to metastasis

 $\triangleright T_d$: time to death

 $\triangleright \mathcal{H}^*(t)$: a version of the PSA history up to t

 $hd T_m^{(a)}$ and $T_d^{(a)}$ counterfactual outcomes

* a = 1, ST given at t

* a=0, ST was not given in $[t,t+\Delta t]$



Marginal Salvage Therapy Effect

b we average over all PSA histories

$$ST^{M}(t + \Delta t, t) = \Pr\{T_{m}^{(1)} \le t + \Delta t \mid T_{m} > t, T_{d} > t\} - \Pr\{T_{m}^{(0)} \le t + \Delta t \mid T_{m} > t, T_{d} > t\}$$

• Notes:

 \triangleright of lesser relevance to the urologists because they decide who gets ST based on PSA \Rightarrow more bias

▷ averages over a big group of patients ⇒ smaller variance



Conditional Salvage Therapy Effect

 \triangleright we condition on the PSA history of a specific patient, i.e., $\mathcal{H}^*(t) = \mathcal{H}_i(t)$

$$ST^{C}(t + \Delta t, t) = \Pr\{T_{m}^{(1)} \le t + \Delta t \mid T_{m} > t, T_{d} > t, \mathcal{H}_{i}(t)\}$$
$$-\Pr\{T_{m}^{(0)} \le t + \Delta t \mid T_{m} > t, T_{d} > t, \mathcal{H}_{i}(t)\}$$

• Notes:

 \triangleright much more relevant to the urologists \Rightarrow **less bias**

▷ averages over a narrow group of patients ⇒ larger variance



Marginal-Conditional Salvage Therapy Effect

 \triangleright consider ST for patients who had PSA levels above the threshold value c at their last visit, i.e., $\mathcal{H}^*(t) = \{Y(t): Y(t) > c\}$

$$ST^{MC}(t + \Delta t, t) = \Pr\{T_m^{(1)} \le t + \Delta t \mid T_m > t, T_d > t, \mathcal{H}^*(t)\}$$
$$-\Pr\{T_m^{(0)} \le t + \Delta t \mid T_m > t, T_d > t, \mathcal{H}^*(t)\}$$

• Notes:

- ▷ relevant to the urologists ⇒ compromised bias
- ▷ averages over a bigger group of patients ⇒ compromised variance

3 Structural Models



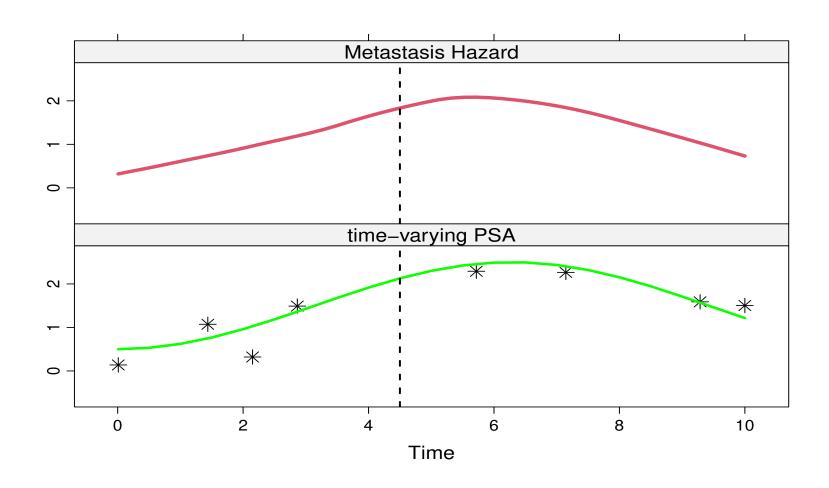
Standard Cox models not appropriate



Joint Models for Longitudinal and Time-to-Event Data

3 Structural Models (cont'd)





3 Structural Models (cont'd)



Joint models completely specify the joint distribution of PSA, time-to-metastasis & time-to-death

- Under sequential ignorability,
 - > they provide valid marginal distributions
 - > without requiring to model the treatment assignment mechanism

4 PSA Sub-Model



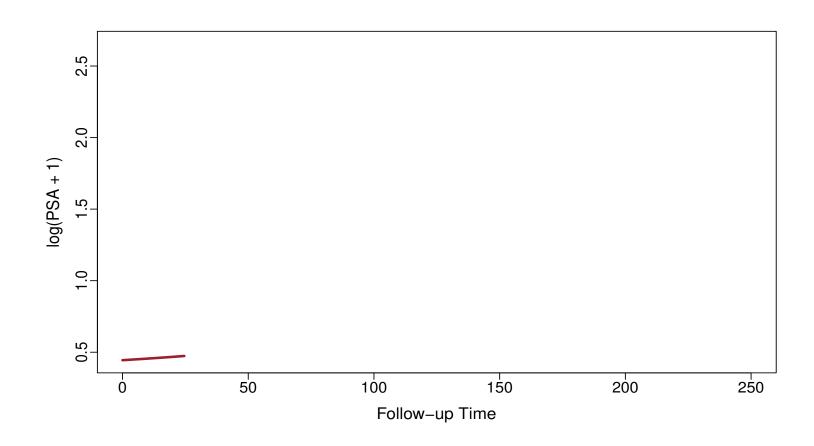
- As PSA increases, patients may receive ST
- ullet We let S_i denote the time a patient initiated ST
 - \triangleright for patients who did not initiate ST, $S_i = \infty$
- After ST, PSA levels are expected to drop
 - but may rise again before metastasis



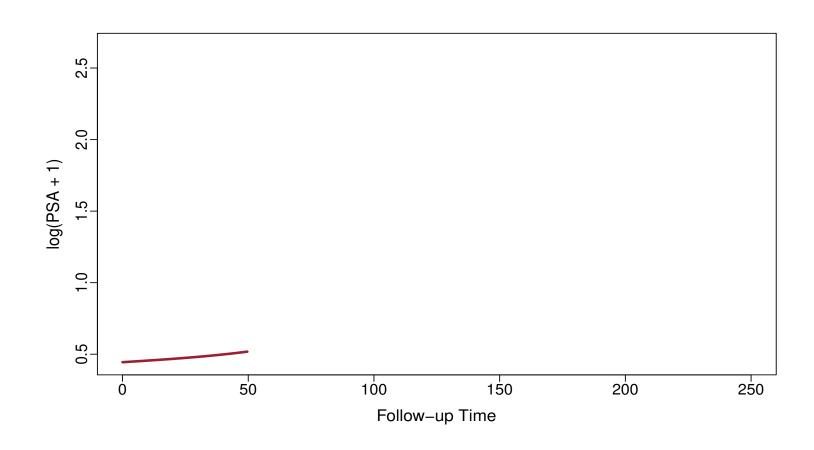
$$\log\{\mathsf{PSA}_i(t)+1\} = \begin{cases} \eta_i(t) + \varepsilon_i(t) = \boldsymbol{x}_i(t)\boldsymbol{\beta} + \boldsymbol{z}_i(t)\boldsymbol{b}_i + \varepsilon_i(t), \ t < S_i \\ \\ \tilde{\eta}_i(t) + \varepsilon_i(t) = \\ \\ \eta_i(t) + \left\{\tilde{\boldsymbol{x}}_i(\tilde{t})\tilde{\boldsymbol{\beta}} + \tilde{\boldsymbol{z}}_i(t)\tilde{\boldsymbol{b}}_i\right\} + \varepsilon_i(t), \quad t \geq S_i, \end{cases}$$

$$oldsymbol{u}_i = (oldsymbol{b}_i, ilde{oldsymbol{b}}_i) \sim \mathcal{N}(oldsymbol{0}, oldsymbol{\Omega})$$

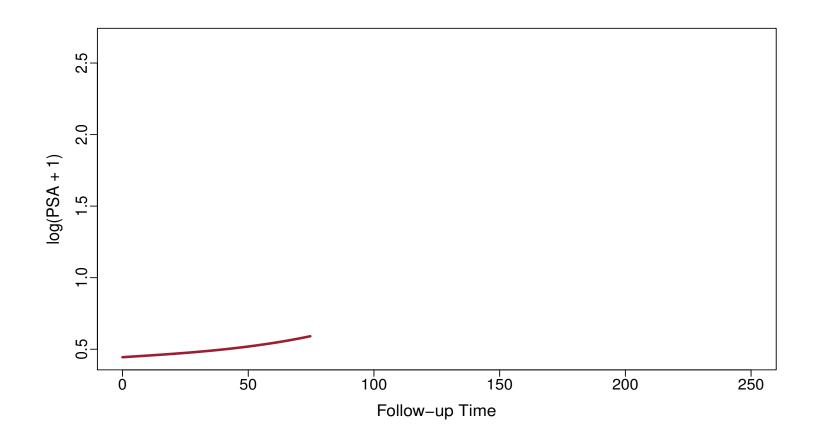




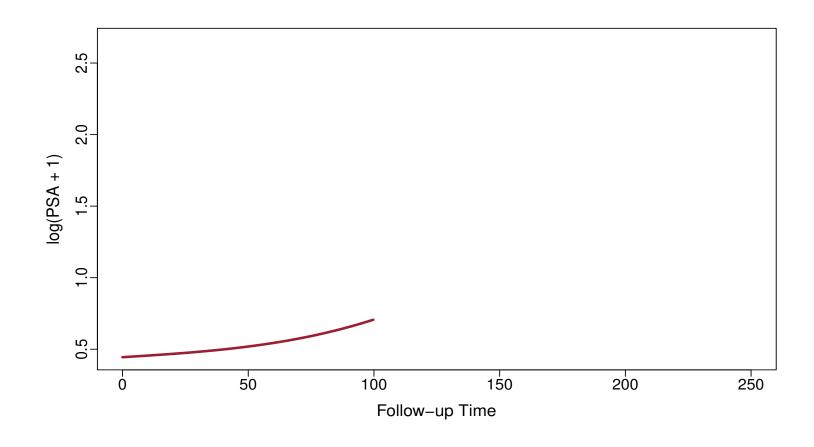




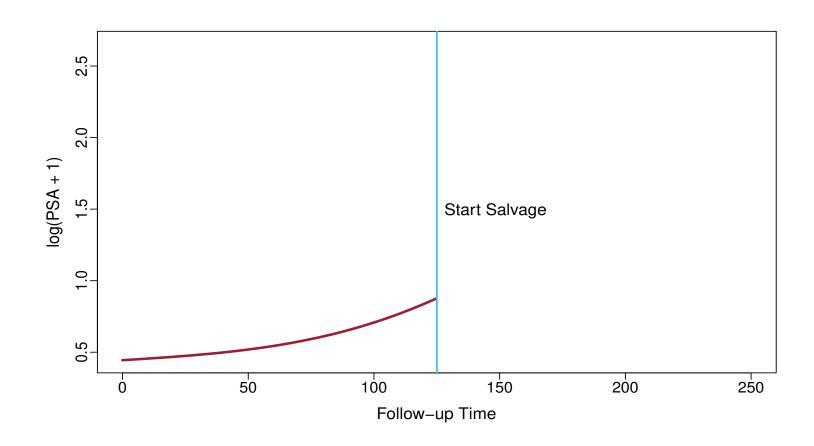




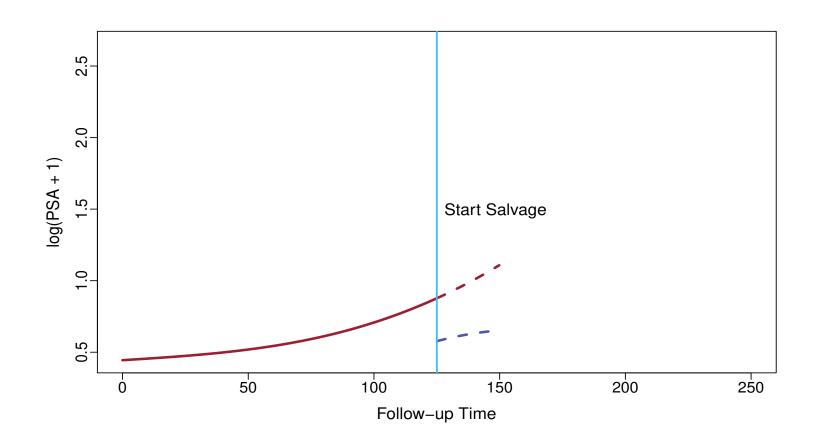




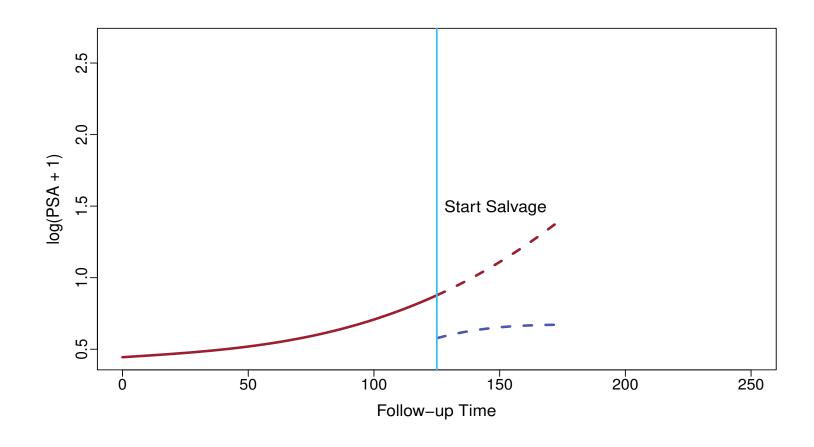




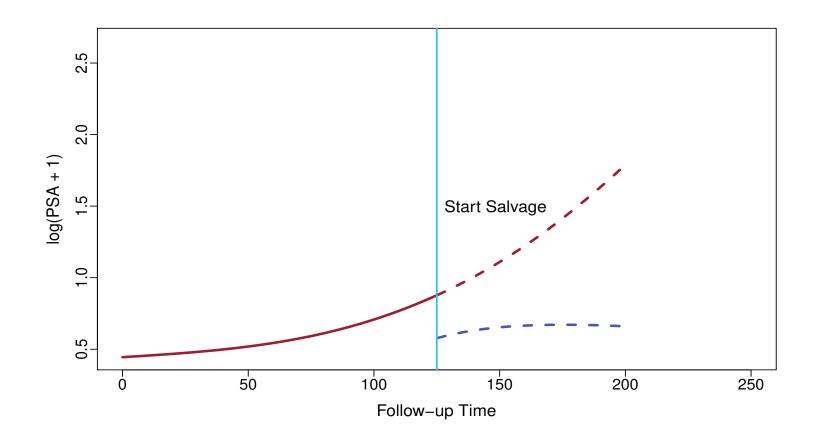




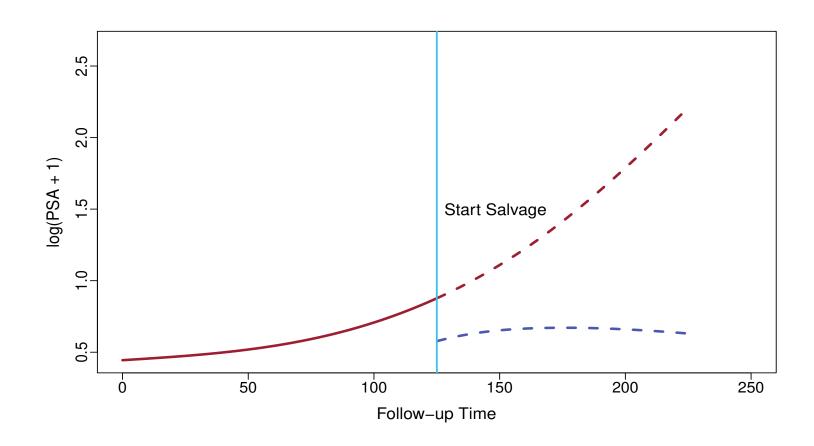




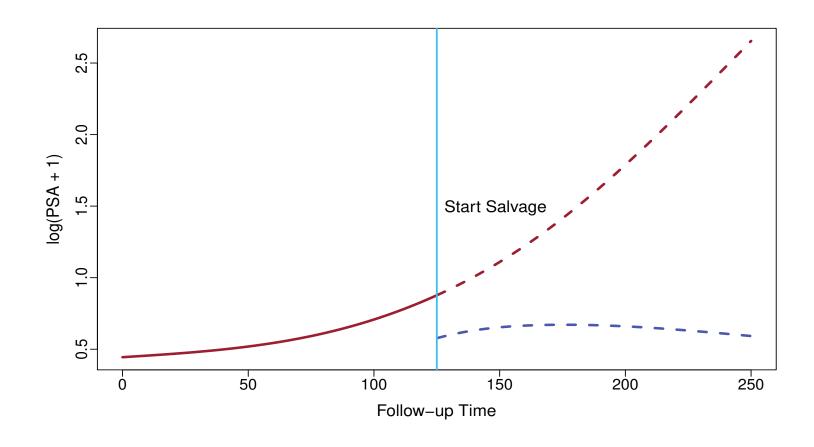












5 Metastasis and Death Sub-Models



- Metastasis and Death treated as *Competing Risks*
- Separate hazard models for metastasis and death

 - ▷ baseline covariates

5 Metastasis and Death Sub-Models (cont'd)



• Metastasis Sub-Model linked to baseline covariates, Salvage and PSA

$$h_i^m(t) = \begin{cases} h_0^m(t) \exp\left(\boldsymbol{\psi}_m^{\top} \boldsymbol{w}_i + \boldsymbol{\alpha}_m^{\top} f\{\eta_i(t)\}\right), & t < S_i \\ h_0^m(t) \exp\left(\boldsymbol{\psi}_m^{\top} \boldsymbol{w}_i + \gamma_m(t - S_i) + \boldsymbol{\xi}_m^{\top} g\{\tilde{\eta}_i(t)\}\right), & t \ge S_i \end{cases}$$

5 Metastasis and Death (cont'd)



• Death Sub-Model linked to baseline covariates, Salvage but not PSA

$$h_i^d(t) = \begin{cases} h_0^d(t) \exp(\boldsymbol{\psi}_d^{\top} \boldsymbol{w}_i), & t < S_i \\ h_0^d(t) \exp(\boldsymbol{\psi}_d^{\top} \boldsymbol{w}_i + \gamma_d), & t \ge S_i \end{cases}$$

6 Causal Effect Estimation



• From the joint model, we can obtain the conditional causal effect

$$\Pr\{T_{mi}^{(a)} \leq t + \Delta t \mid T_{mi} > t, T_{di} > t, \mathcal{H}_{i}(t), \mathcal{X}_{i}\} =$$

$$\int \int \Pr\{T_{mi}^{(a)} \leq t + \Delta t \mid T_{mi} > t, T_{di} > t, \boldsymbol{u}_{i}, \mathcal{X}_{i}, \boldsymbol{\theta}\}$$

$$\times p\{\boldsymbol{u}_{i} \mid T_{mi} > t, T_{di} > t, \mathcal{H}_{i}(t), \mathcal{X}_{i}, \boldsymbol{\theta}\} \ p(\boldsymbol{\theta} \mid \mathcal{D}) \ d\boldsymbol{u}_{i} d\boldsymbol{\theta}$$

6 Causal Effect Estimation (cont'd)



- ullet Monte Carlo scheme to estimate $\mathrm{ST}_i^C(t+\Delta t,t)$
 - riangle sample $reve{m{ heta}}^{(l)}$ from the posterior of the parameters $[m{ heta} \mid \mathcal{D}]$
 - ightharpoonup sample $m{ar{u}}_i^{(l)}$ from the posterior of the random effects $[m{u}_i \mid T_{mi} > t, T_{di} > t, \mathcal{H}_i(t), \mathcal{X}_i, m{ar{ heta}}^{(l)}]$

$$ho$$
 calculate $\pi_i^{(l)}(t + \Delta t \mid t, a) = \Pr\{T_{mi}^{(a)} \leq t + \Delta t \mid T_{mi} > t, T_{di} > t, \boldsymbol{\check{u}}_i^{(l)}, \boldsymbol{\mathcal{X}}_i, \boldsymbol{\check{\theta}}^{(l)}\}$

ullet We repeat L times and get

$$\widehat{\mathsf{ST}}_{i}^{C}(t + \Delta t, t) = \frac{1}{L} \sum_{l=1}^{L} \pi_{i}^{(l)}(t + \Delta t \mid t, a = 1) - \pi_{i}^{(l)}(t + \Delta t \mid t, a = 0)$$

6 Causal Effect Estimation (cont'd)

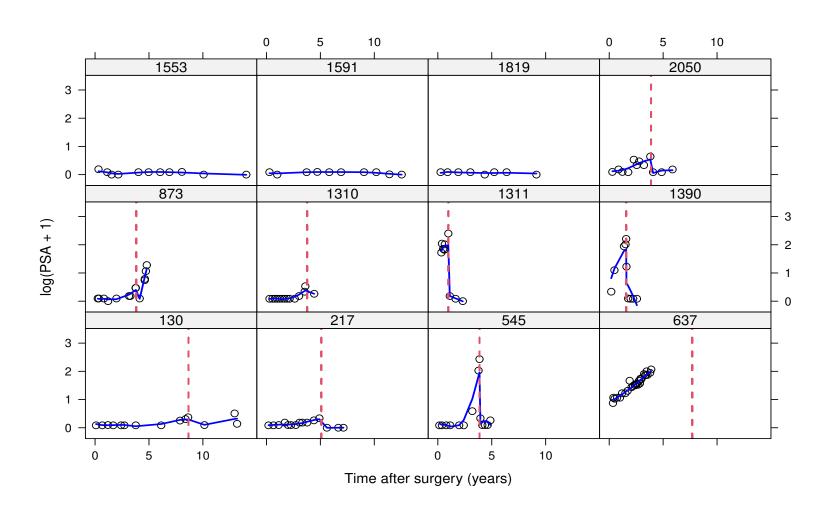


- ullet Estimation of $\mathrm{ST}^M(t+\Delta t,t)$ and $\mathrm{ST}^{MC}(t+\Delta t,t)$ proceeds by averaging the conditional effects over the respective groups of patients
- ullet For example, for $\mathrm{ST}^M(t+\Delta t,t)$
 - $\triangleright \mathcal{R}(t)$ the subset of patients at risk at time t
 - ho for each patient in $\mathcal{R}(t)$, we calculate $\widehat{\mathsf{ST}}^C_i(t+\Delta t,t)$

$$\widehat{\mathsf{ST}}^M(t+\Delta t,t) = n_r^{-1} \sum_{i:i \in R(t)} \widehat{\mathsf{ST}}^C_i(t+\Delta t,t),$$

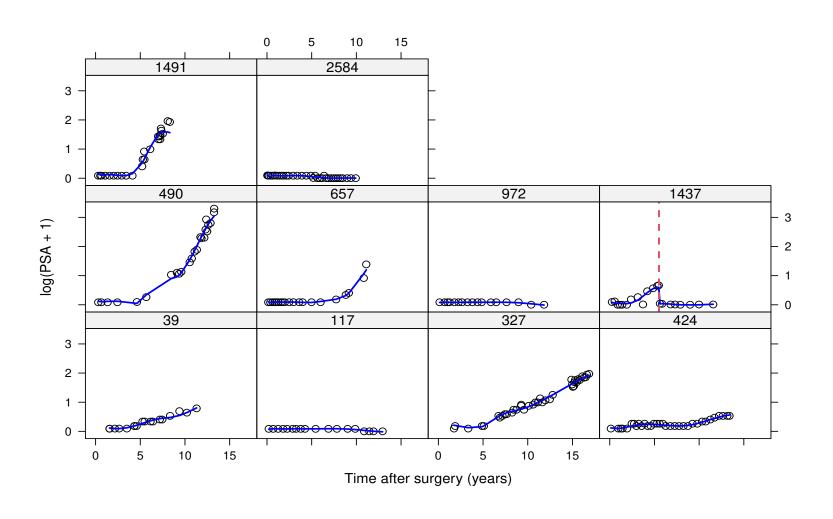
7 Results





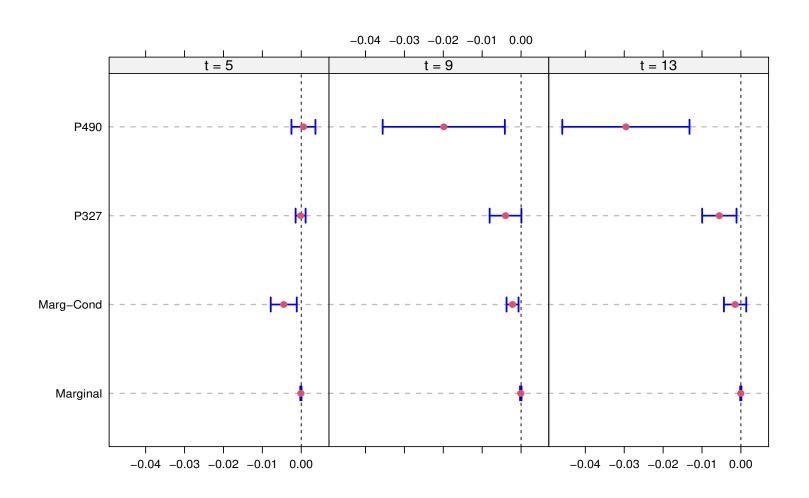
7 Results (cont'd)





7 Results (cont'd)





7 Software (cont'd)



- Implementation available in JMbayes2
 - > predict() cumulative incidence risks
- Shiny app...

Thank for your attention!

https://www.drizopoulos.com/