Absolute risk integration using penalized logistic regression

Jesse Islam

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 - Parametric models

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- They believe the stepwise nature is the reason, as it reduces interpretability. [1]
- Want to easily model non-proportional hazards. [1]
- A streamlined approach for reaching a smooth absolute risk curve. [1]

Dr. Cox's perspective

Reid: How do you feel about the cottage industry that's grown up around it [the Cox model]?

Cox: Don't know, really. In the light of some of the further results one knows since, I think I would normally want to tackle problems parametrically, so I would take the underlying hazard to be a Weibull or something. I'm not keen on nonparametric formulations usually.

Reid: So if you had a set of censored survival data today, you might rather fit a parametric model, even though there was a feeling among the medical statisticians that that wasn't quite right.

Cox: That's right, but since then various people have shown that the answers are very insensitive to the parametric formulation of the underlying distribution [see, e.g., Cox and Oakes, Analysis of Survival Data, Chapter 8.5]. And if you want to do things like predict the outcome for a particular patient, it's much more convenient to do that parametrically.

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- Design: Prospective cohort study.
- Setting: 5 academic care centers in the United States.
- Participants: 9105 hospitalized.
- Follow-up-time: 5.56 years.
- 68% incidence rate.

SUPPORT manual imputation

Notorious for missing data

Baseline Variable	Normal Fill-in Value
Bilirubin	1.01
BUN	6.51
Creatinine	1.01
PaO2/FiO2 ratio (pafi)	333.3
Serum albumin	3.5
Urine output	2502
White blood count	9 (thousands)

 Table 1: Suggested imputation values. [Support site reference]

Mice imputation package (R)

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- 3. polyreg(Bayesian polytomous regression) For Factor Variables (>= 2 levels)

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- Previous model findings. (8)

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- Both on the scale of 180 days.

• Write out complicated model?????

• image of SPS vs APS ???????

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- Absolute Risk comparison.

1. Impute

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- 2. Compare SPS and APS over ~ 5.56 years using absolute risk curves.

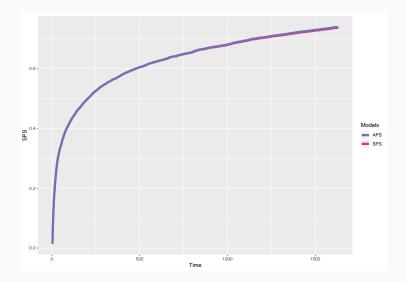
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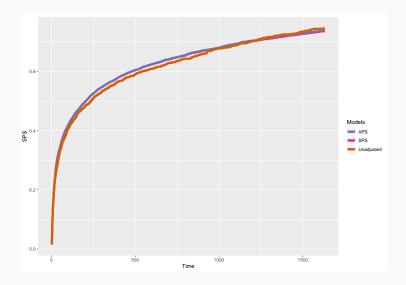
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- All models is trained on 80% of the observations.
- Remaining observations are used to generate comparative absolute risk curves.

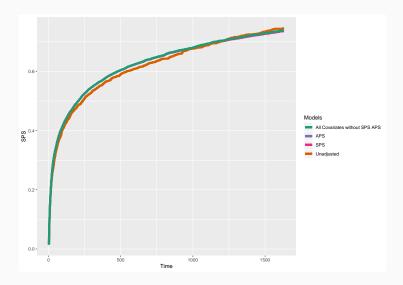
SPS vs APS



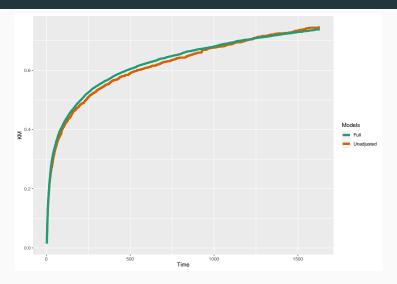
SPS vs. Kaplan-Meier



All covariates vs. physiology scores vs unadjusted



Chosen absolute risk comparisons



Conclusion:

• Linear associations without physiology scores perform similarly

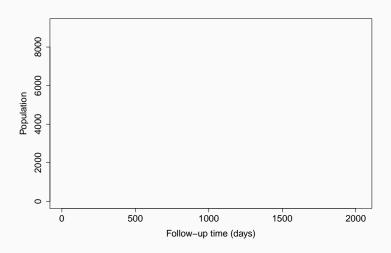
1. Clever sampling.

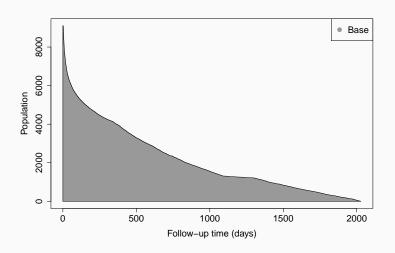
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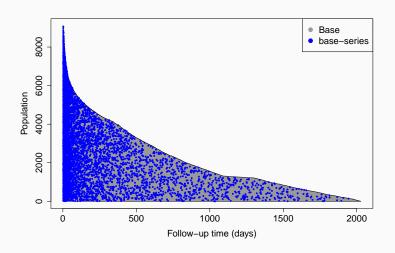
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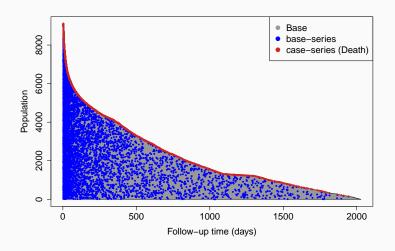
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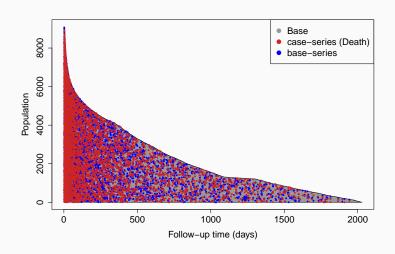
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 - Casebase is parametric, and allows different parametric fits by incorporation of the time component.
- Package contains an implementation for generating population-time plots.







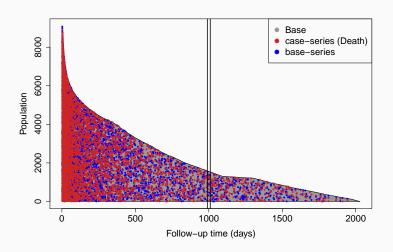




$$e^{L} = \frac{Pr(Y = 1|x, t)}{Pr(Y = 0|x, t)} = \frac{h(x, t) * B(x, t)}{b[B(x, t)/B]} = \frac{h(x, t) * B}{b}$$

* $L = \beta X$ * b = base-series. * B = Base. * B(x,t) = Risk-set for survival time t.

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log-odds = log hazard

$$e^{L} = \frac{h(x, t) * B}{b}$$
$$\frac{b * e^{L}}{B} = \hat{h}(x, t)$$
$$log(\hat{h}(x, t)) = L + log(\frac{b}{B})$$

Maximum log-likelihood, with regularization

$$I(\beta) = \sum_{i=1}^{N} \left(\sum_{k=0}^{K} x_{ik} \beta_k \right) - n_i log \left(1 + e^{\sum_{k=0}^{K} x_{ik} \beta_k} \right)$$

Maximum log-likelihood, with offset

$$I(\beta) = \sum_{i=1}^{N} \left(\sum_{k=0}^{K} x_{ik} \beta_k + \frac{b}{B} \right) - n_i \log \left(1 + e^{\sum_{k=0}^{K} x_{ik} \beta_k + \frac{b}{B}} \right)$$

Maximum log-likelihood, with offset and lasso

$$I(\beta) = \sum_{i=1}^{N} \left(\sum_{k=0}^{K} x_{ik} \beta_k + \frac{b}{B} \right) - n_i \log \left(1 + e^{\sum_{k=0}^{K} x_{ik} \beta_k + \frac{b}{B}} \right) - \lambda ||\beta||$$

Casebase: Parametric families

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$$log(h(t;\alpha,\beta)) = g(t;\alpha) + \beta X$$

Casebase: Parametric families

• We can now fit models of the form:

$$log(h(t; \alpha, \beta)) = g(t; \alpha) + \beta X$$

• By changing the function $g(t; \alpha)$, we can model different parametric families easily:

Casebase: Parametric models

Exponential: $g(t; \alpha)$ is equal to a constant

casebase::fitSmoothHazard(status ~ X1 + X2)

Gompertz: $g(t; \alpha) = \alpha t$

casebase::fitSmoothHazard(status ~ time + X1 + X2)

Weibull: $g(t; \alpha) = \alpha log(t)$

casebase::fitSmoothHazard(status ~ log(time) + X1 + X2)

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- h(x,u)= Hazard function
- Lets use the weibull hazard

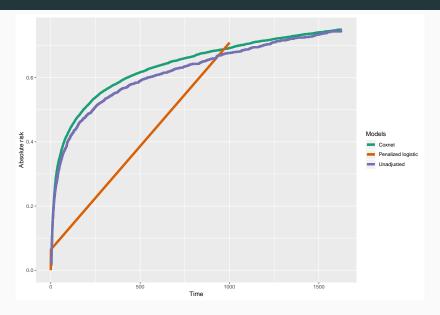
casebase surv weibull-> LASSO

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Survival comparison



Brier score equation

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- Calibration and discrimination
- IPA score equation
- In progress

Future work

Survival GWAS

References 1

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- 2.Saarela, Olli, and Elja Arjas. 2015. "Non-Parametric Bayesian Hazard Regression for Chronic Disease Risk Assessment." Scandinavian Journal of Statistics 42 (2). Wiley Online Library: 609–26.
- 3.Saarela, Olli. 2015. "A Case-Base Sampling Method for Estimating Recurrent Event Intensities." *Lifetime Data Analysis*. Springer, 1–17

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Czepiel, S. A. (2002). Maximum likelihood estimation of logistic regression models: theory and implementation. Available at czep. net/stat/mlelr. pdf, 1825252548-1564645290. ????

Tutorial and Slides

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\begin{center} Tutorial:
http://sahirbhatnagar.com/casebase/
Slides:
https://github.com/Jesse-
Islam/ATGC_survival_presentation_Feb.27.2020
Questions? \end{center}
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