

Tutorial: Bayesian variable selection for survival data

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Overview

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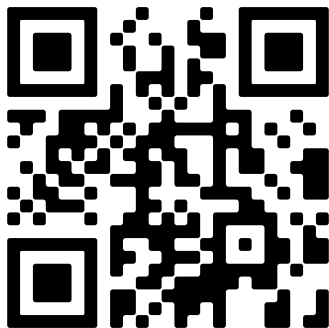


Figure: <https://github.com/FJRubio67/BVSSurv>

Survival Models

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- In some cases, we may know some additional characteristics about the individuals, meaning we have access to **covariates** $\mathbf{x}_i = (x_{i1}, \dots, x_{ip})^\top$, (age, sex, deprivation level, comorbidities, tumour stage, ...).

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- We need a Statistical Model.

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- Survival models are often formulated using the hazard function.

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$$\ell_p(\boldsymbol{\beta}) = \sum_{\delta_i=1} \mathbf{x}_i^\top \boldsymbol{\beta} - \sum_{\delta_i=1} \log \left(\sum_{k \in \mathcal{R}(t_i)} \exp \{ \mathbf{x}_k^\top \boldsymbol{\beta} \} \right),$$

where t_i , $i = 1, \dots, n$, are the survival times, $\mathcal{R}(t) = \{i : t_i \geq t\}$ denotes the risk set at time t .

- See: <https://rpubs.com/FJRubio/CPHM>

Accelerated Failure Time model

- The AFT postulates that covariates affect simultaneously the time scale and the hazard scale:

$$h_{AFT}(t \mid \mathbf{x}_i, \boldsymbol{\theta}, \boldsymbol{\beta}) = h_0 \left(t \exp \{ \mathbf{x}_i^\top \boldsymbol{\beta} \} \mid \boldsymbol{\theta} \right) \exp \{ \mathbf{x}_i^\top \boldsymbol{\beta} \} .$$

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- See:

<https://github.com/FJRubio67/ShortCourseParamSurvival>

Regression models: the likelihood function

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$$L(\beta, \theta) = \prod_{i=1}^n f_j(t_i | \mathbf{x}_i, \theta, \beta)^{\delta_i} S_j(t_i | \mathbf{x}_i, \theta, \beta)^{1-\delta_i}$$

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- This also shows that the likelihood can be characterised using the hazard function.
- Maximum Likelihood Estimators for several choices of the baseline hazard can be obtained using the R package `HazReg` (<https://github.com/FJRubio67/HazReg>).

Selection Methods

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- Forward, backward, and both strategies.
- Myriad of disadvantages: lack of control on errors, *inconsistent*, lack of uncertainty quantification about the *model selection*.

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- Extensions to survival knockoffs.

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- Bayesian methods (AFT, Cox) + Spike-Slab prior.
- We introduce the variable inclusion indicators

$$\gamma_j = \begin{cases} 1 & \text{if variable } j \text{ is included} \\ 0 & \text{otherwise} \end{cases}$$

- Discrete construction (mixture prior)

$$\beta_j \mid \gamma_j \sim (1 - \gamma_j)\delta_0(\beta_j) + \gamma_j\mathbf{N}(\mathbf{0}, \eta_j),$$

δ_0 is a mass probability at 0.

Continuous Spike-Slab priors

- Continuous construction

$$\beta_j \mid \gamma_j \sim (1 - \gamma_j)F(0, \tilde{\eta}_j) + \gamma_j N(0, \eta_j),$$

F can be a Laplace distribution (spike-and-slab LASSO), or another scale mixture of normals.

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- See: [Handbook of Bayesian Variable Selection](#) [Tadesse and Vannucci, 2021].

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Other methods

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- General message: **No Panacea.**

Bayesian Variable Selection

Model formulation: AFT model

- The AFT model postulates

$$\log(t_i) = \mathbf{x}_i^\top \boldsymbol{\beta} + \epsilon_i,$$

where ϵ_i are independent across $i = 1, \dots, n$ with mean $E(\epsilon_i) = 0$ and variance $V(\epsilon_i) = \sigma^2$ (assumed finite).

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- We will focus on the case where $\epsilon_i \sim N(0, \sigma^2)$, but other distributional assumptions are possible.

- It is convenient to reparameterize $\alpha = \beta/\sigma$, and $\tau = 1/\sigma$, as then the log-likelihood is concave, provided the number of uncensored individuals is greater than the number of model parameters ($n_o \geq p$) and that X_o has full column rank (the design matrix associated to the uncensored observations).
- The log-likelihood is

$$\begin{aligned}\ell(\alpha, \tau) &= -\frac{n_o}{2} \log \left(\frac{2\pi}{\tau^2} \right) - \frac{1}{2} \sum_{\delta_i=1} (\tau y_i - \mathbf{x}_i^\top \alpha)^2 \\ &\quad + \sum_{\delta_i=0} \log \{ \Phi(\mathbf{x}_i^\top \alpha - \tau y_i) \},\end{aligned}\tag{1}$$

- Our goal is model selection, which we formalize as choosing among two possibilities

$$\gamma_j = \begin{cases} 0, & \text{if } \beta_j = 0, \\ 1, & \text{if } \beta_j \neq 0, \end{cases}$$

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corresponding to no effect, or the inclusion of a linear effect of each covariate $j = 1, \dots, p$. There are extensions to selection of non-linear effects but they are beyond the aims of this [Rossell and Rubio, 2023].

- That is, $\gamma = (\gamma_1, \dots, \gamma_p)$ determines what covariates enter the model and their effect, and there are 2^p total models to consider.

Bayesian Variable Selection

The posterior model probabilities

$$\pi(\gamma | y) = \frac{p(y | \gamma)\pi(\gamma)}{\sum_{\gamma} p(y | \gamma)\pi(\gamma)}, \quad (2)$$

where $\pi(\gamma)$ is the model prior probability, and

$$p(y | \gamma) = \int p(y | \alpha_{\gamma}, \tau) \pi(\alpha_{\gamma}, \tau | \gamma) d\alpha_{\gamma} d\tau,$$

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- when the interest is in prediction, use Bayesian model averaging where models are weighted according to $\pi(\gamma \mid y)$,
- or alternatively choosing a sparse model giving similar predictions.

Either way $\pi(\gamma \mid y)$ are critical for inference, hence the importance to understand their behavior.

Priors

Two priors: local and non-local

Idea: g-priors or non-local priors for α .

$$\pi_L(\alpha_\gamma, \tau) = \prod_{\gamma_j=1} N(\alpha_j; 0, g_L n / (x_j^\top x_j)) \pi(\tau)$$

$$\pi_M(\alpha_\gamma, \tau) = \prod_{\gamma_j=1} \frac{\alpha_j^2}{g_M} N(\alpha_j; 0, g_M) \pi(\tau).$$

Priors on the precision and the model

- By default, we consider independent Beta-Binomial priors

$$\pi(\gamma) = \text{BetaBin}(p_\gamma; p, a_1, b_1),$$

where $\text{BetaBin}(z; p, a, b)$ is Beta-Binomial distribution.

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- and $g_L, g_M, a_\tau, b_\tau \in \mathbb{R}_+$ are given dispersion parameters: Not an automatic method!

Local prior

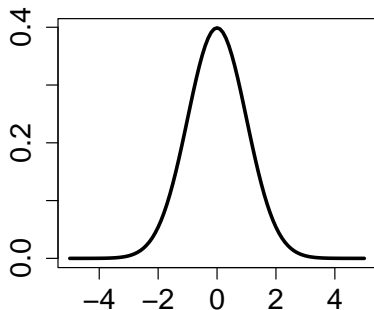


Figure: g-prior

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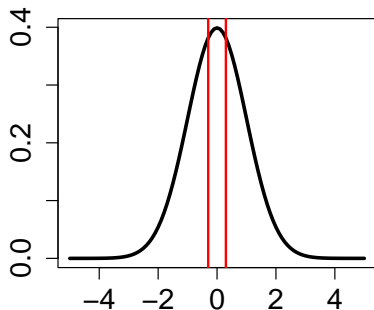


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Non-local prior

Johnson and Rossell [2010]

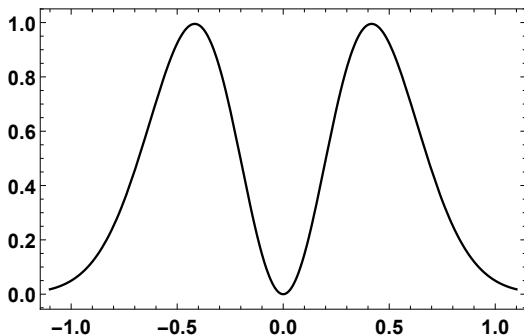


Figure: “Bayes factors that are obtained by using local alternative priors exhibit a disturbing large sample property. As the sample size n increases, they accumulate evidence much more rapidly in favour of true alternative models than in favour of true null models.”

Computations

Computations: Approximations and Model Space Exploration

- We approximate the marginal likelihood $p(y \mid \gamma)$ using a **Laplace approximation** (tricks to recycle and make more efficient some calculations).

$$\hat{p}(y \mid \gamma) = \exp\{\ell(\tilde{\eta}_\gamma) + \log \pi(\tilde{\eta}_\gamma)\} (2\pi)^{d_\gamma/2} |H(\tilde{\eta}_\gamma) + \nabla^2 \log \pi(\tilde{\eta}_\gamma)|^{-1/2},$$

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- Not covered: Approximate Laplace Approximations [Rossell et al., 2021].

Approximations and Model Space Exploration

- **Model exploration:** Gibbs sampling (MCMC in general). Active research area.

Approximations and Model Space Exploration

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 - 1 Initialise $\gamma = \gamma_0$.
 - 2 Update $\gamma_j^{(t)} = 1$ with probability

$$\frac{p(\gamma_1^{(t)}, \dots, \gamma_{j-1}^{(t)}, \gamma_j^{(t-1)}, \gamma_{j+1}^{(t)}, \dots, \gamma_p^{(t)} \mid \mathbf{y})}{\sum_{\gamma_j=0}^1 p(\gamma_1^{(t)}, \dots, \gamma_{j-1}^{(t)}, \gamma_j, \gamma_{j+1}^{(t)}, \dots, \gamma_p^{(t)} \mid \mathbf{y})}$$

Theory

To interpret the results in the M-Open scenario, we need to define the expected log-likelihood under the data-generating F_0 .

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Under minimal conditions, $M(\eta_\gamma)$ has a unique maximiser, denoted by $\eta_\gamma^* = (\alpha_\gamma^*, \tau_\gamma^*)$. M is affected by both, **the survival process and the censoring mechanism**.

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- This implies that the highest posterior probability model consistently selects γ^* , and that including covariates with marginal posterior probability $\pi(\gamma_j^* | y) > t$, for any fixed threshold t , also leads to consistent selection.
- Bayesian model selection in the AFT model asymptotically returns the smallest γ^* that minimises the KL divergence between the true model and the AFT model.

Examples

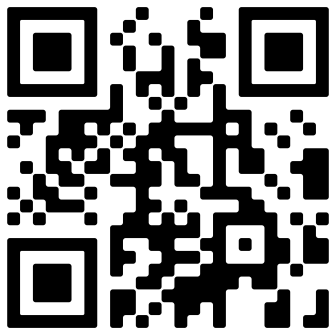


Figure: <https://github.com/FJRubio67/BVSSurv>

Example 1: Simulated data

- Simulated data from an accelerated failure time model
- $n = 250$, $p = 10$, 47% censoring.
- R code.

<https://rpubs.com/FJRubio/BVSSurvExample1>

- Sensitivity analyses: Nikooienejad et al. [2020], Simon et al. [2011], Yi et al. [2019]

Example 2: flchain data set

- This is a stratified random sample containing 1/2 of the subjects from a study of the relationship between serum free light chain (FLC) and mortality.
- $n = 6524$, $p = 5$, 70% censoring.
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- Homework: answer the open question explained in the help file.

Example 3: colon cancer data set

- Association of 172 genes + TFG-B (growth factor) + tumour stage with colon cancer survival
- $n = 260$, $p = 175$, 80% censoring.
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<https://rpubs.com/FJRubio/BVSSurvExample3>

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- Homework: Stratified analysis.

Example 4: nki70 data set

- 144 lymph node positive breast cancer patients on metastasis-free survival, 5 clinical risk factors, and gene expression measurements of 70 genes found to be prognostic for metastasis-free survival in an earlier study.
- $n = 144$, $p = 75$, 66% censoring.
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<https://rpubs.com/FJRubio/BVSSurvExample4>

- Small sample, moderate dimension, high censoring.

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- Small sample, moderate dimension, high censoring.
- Homework: Reflect on reasons for differences (prior calibration? Model misspecification?, All?).
- Homework 2: correct data preparation (using factors and dummy variables instead of numeric).

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- Censoring also has an effect in the finite-sample scenario: should we increase the sample size or the follow-up to improve power of BVS?
- Bayesian variable selection is not automatic: prior calibration is crucial in the finite sample scenario.

Additional software tools

- BVS for Cox model and non-local prior: `BVSNLP` R package.
- Spike and Slab LASSO (only posterior modes): `BhGLM` R package.
- BART: `BART` R package.
- Random Survival Forests: `randomForestSRC` R package.
- Cox-LASSO: `glmnet` R package.

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