# **Bayesian survival time prediction**

# for high-dimensional causal effects estimation using regression trees

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### Problem

Estimating causal effects for survival data in settings with many more predictors than observations (p > n) is challenging.

We require a method to predict the survival times:

 $\mathbb{E}[T \mid A = a, X = x],$ 

given A (treatment) and  $X \in \mathbb{R}^p$  (covariates) that captures nonlinear relationships and interaction effects. Additionally, the method should adapt to different levels of sparsity while retaining all variables in the model.

# **Results**

We performed a simulation study with the following settings:

Sample size of n = 100 observations 

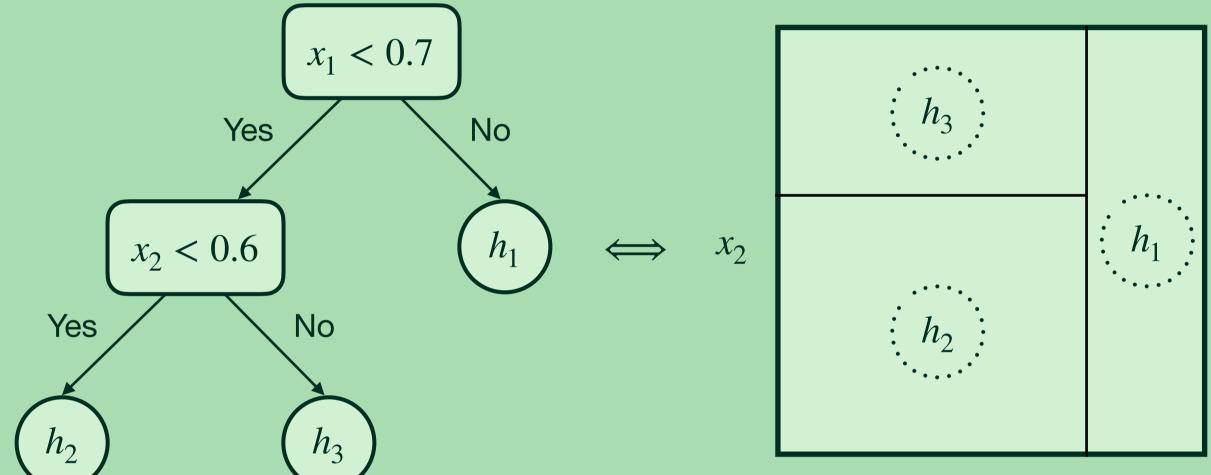
#### Model

The survival time is modelled using an *accelerated failure time (AFT*) model:

$$\log(T) = \sum_{j=1}^{m} f(X, a; \theta_j) + \epsilon,$$

with a sum-of-trees regression function and  $\epsilon \sim \mathcal{N}(0, \sigma^2)$ .

Each tree recursively partitions the covariate space, assigning a parameter value to each region. A two-dimensional example of a single tree:



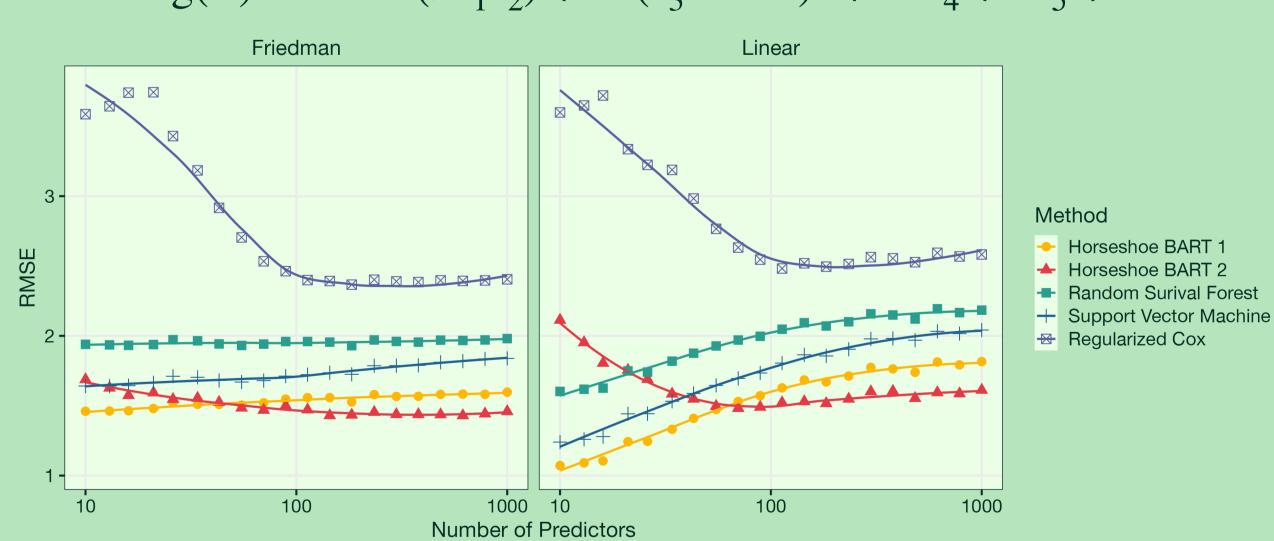
- **Censoring rate** is ca. 30%
- Noise level is ca. 20%
- Covariates  $X \in \mathbb{R}^p$  are drawn independently from U[0,1]

We consider two data-generating processes:

1. Linear model with 10% of nonzero coefficients:

 $\log(T) = \beta^{\top} X + \epsilon \,.$ 

2. Nonlinear model with interactions terms (Friedman function):



### $\log(T) = 10\sin(\pi x_1 x_2) + 20(x_3 - 1/2)^2 + 10x_4 + 5x_5 + \epsilon.$

 $x_1$ 

For each tree, we model the step heights h using a horseshoe prior, which provides adaptive shrinkage. This prior is placed on the distribution of h as follows:

> $h_j \mid \lambda_j^2, \tau^2, \sigma^2 \sim \mathcal{N}(0, \lambda_j^2 \tau^2 \sigma^2),$  $\lambda_j^2 \sim C^+(0, 1),$  $\tau^2 \sim C^+(0, 1),$

where  $C^+$  denotes the half-Cauchy distribution.

This allows for global shrinkage via  $\tau$  and for local signals to be picked up via the  $\lambda$ 's.

We use a non-informative inverse gamma prior for  $\sigma^2$  while the trees follow a BART prior.



Methods are compared using the root mean square error (RMSE), evaluated on a held-out set of size 1000.

Posterior samples are drawn using an efficient reversible jump Markov chain Monte Carlo (RJ-MCMC) algorithm.



## Conclusion

Our method effectively captures nonlinearity, interactions, and adapts to sparsity, outperforming alternatives in high-dimensional survival prediction while retaining all variables for causal effect estimation.



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