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A proportional hazards regression model for the subdistribution with covariates adjusted censoring weight for competing risks data

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ABSTRACT. With competing risks data, one often needs to assess the treatment and covariate effects on the cumulative incidence function. Fine and Gray proposed a proportional hazards regression model for the subdistribution of a competing risk with the assumption that the censoring distribution and the covariates are independent. Covariate-dependent censoring sometimes occurs in medical studies. In this paper, we study the proportional hazards regression model for the subdistribution of a competing risk with proper adjustments for covariate-dependent censoring. We consider using a covariate-adjusted weight function by fitting the Cox model for the censoring distribution and using the predictive probability for each individual. Our simulation study shows that the covariate-adjusted weight estimator is basically unbiased when the censoring time depends on the covariates, and the covariate-adjusted weight approach works well for the variance estimator as well. We illustrate our methods with bone marrow transplant data from the Center for International Blood and Marrow Transplant Research (CIBMTR). Here cancer relapse and death in complete remission are two competing risks.

key words: competing risks; cumulative incidence function; proportional hazards model; subdistribution; inverse probability of censoring weight

1 Introduction

Biomedical research often involves competing risks in which each subject is at risk of failure from K different causes. For competing risks data, one only observes the first event to occur and precludes the occurrence of another event. Also, one often wishes to estimate and model the cumulative incidence function (CIF), which is the marginal probability of failure of a specific cause. The standard approach of modeling CIF is to model the cause-specific hazard function for all causes. Let $\lambda_k(t; \mathbf{Z})$ be the k th conditional cause-specific hazard ($k = 1, 2$ for simplicity), where \mathbf{Z} is given set of covariates. The CIF of cause 1 given by \mathbf{Z} is

$$F_1(t; \mathbf{Z}) = P(\tilde{T} \leq t, \epsilon = 1 | \mathbf{Z}) = \int_0^t \lambda_1(s; \mathbf{Z}) \exp \left[- \int_0^{s^-} \{ \lambda_1(u; \mathbf{Z}) + \lambda_2(u; \mathbf{Z}) \} du \right] ds,$$

where \tilde{T} is the failure time and ϵ indicates the type of failure. Here, all cause-specific hazards need to be modeled adequately and correctly. [Prentice et al. \(1978\)](#) and [Cheng et al. \(1998\)](#) proposed using [Cox \(1972\)](#) proportional hazards model for all causes. Alternatively, [Shen & Cheng \(1999\)](#) considered a special additive model, and [Scheike & Zhang \(2002, 2003\)](#) proposed and studied a flexible Cox-Aalen model, which allows some of the covariates to have time-varying effects. Since the cumulative incidence function of a specific cause is a function of cause-specific hazards for all causes, it is difficult to summarize the covariate effect ([Zhang & Fine, 2008](#)) and identify the covariate effect on the cumulative incidence function. Recently, some new regression methods have been developed to directly model the cumulative incidence function. [Fine & Gray \(1999\)](#) (FG) developed a regression method to directly model the CIF by modeling the subdistribution hazard function through a Cox type regression model, $\lambda_k^*(t; \mathbf{Z}) = -d \log\{1 - F_k(t; \mathbf{Z})\} / dt = \lambda_{k0}^*(t) \exp\{\boldsymbol{\beta}_k^T \mathbf{Z}\}$ based on early work by [Gray \(1988a\)](#) and [Pepe \(1991\)](#). FG proposed using an inverse probability of the censoring weighting (IPCW) technique to estimate the regression parameter $\boldsymbol{\beta}$ and cumulative baseline subdistribution hazard function $\Lambda_{k0}^*(t) = \int_0^t \lambda_{k0}^*(s) ds$. This approach has been implemented in an R-package, `cmprsk`. FG's model has been considered and used extensively in cancer studies, epidemiological studies, and many other biomedical studies ([Scrucca et al., 2007](#); [Wolbers et al., 2009](#); [Kim, 2007](#); [Lau et al., 2009](#)). Let $r(t) = I\{C \geq (\tilde{T} \wedge t)\}$ and $G_C(t) =$

$P(C > t)$, where C is the censoring time. Fine and Gray's approach is based on the fact that $E[r(t)/G_C\{(\tilde{T} \wedge C) \wedge t\} | Data] = 1$ provided that censoring time is independent of the covariates, and FG proposed using the Kaplan-Meier estimator to estimate the unknown censoring distribution G_C . However, in biomedical research studies, the censoring time may depend on some of the covariates and the treatment group. In a clinical trial, patients may be more likely to drop out with some specific value of covariate characteristics, and one treatment group may have a higher dropout rate than the others (Mai, 2008). DiRienzo & Lagakos (2001a,b) showed when the distribution of censoring depends on both treatment group and the covariates, in general the null asymptotic distribution of the score test is not centered at zero when the model is misspecified, the tests of treatment group effect can be severely biased. Heinze et al. (2003) showed that if the censoring distributions are not similar in the two comparison groups, the log-rank test and fitting a regression model, such as fitting a proportional hazards model, may not be valid. For the competing risks data, one can show that $E[r(t)/G_C\{(\tilde{T} \wedge C) \wedge t | Data\} | Data] = 1$, where $G_C\{(\tilde{T} \wedge C) \wedge t | Data\}$ is the conditional censoring distribution given by $Data$. Thus, parameter estimates using the inverse probability of censoring weighting approach with the Kaplan-Meier estimator may be biased when the censoring distribution depends on some of the covariates. To adjust the IPCW when censoring distribution depends on some of the covariates, Fine & Gray (1999) suggested using a stratified Kaplan-Meier estimator for the discrete covariates and assuming the Cox model for the continuous covariates. In this study, we considered a regression model for the censoring distribution, such as a Cox proportional hazards model, and using predicted the censoring probability for each individual subject for the weight function. With the Cox model adjusted weight, we derived an efficient variance estimator, and we performed a simulation study to examine the bias that would arise without adjusting covariates for estimating the censoring distribution, potential bias reduction and robustness of using the Cox model for the censoring distribution. Furthermore, Fine and Gray proposed using a stabilized factor $\hat{G}_C(t)$ with inverse weight $r(t)\hat{G}_C(t)/\hat{G}_C\{(\tilde{T} \wedge C) \wedge t\}$. Our simulation indicates that this stabilized weight improves the efficiency and reduces the bias, but not enough. With the Cox model adjusted weight function, we also considered using a stabilized weight $r(t)\hat{G}_C(t|\mathbf{X})/\hat{G}_C\{(\tilde{T} \wedge C) \wedge t|\mathbf{X}\}$ to improve efficiency and to reduce bias, where \mathbf{X} is the covariates, which is associated with the censoring distribution and could be a subset covariates of \mathbf{Z} .

The outline of the remainder of the paper is as follows. In Section 2 we describe the

competing risks data structure. We introduce a regression-adjusted inverse weighted estimation for the proportional subdistribution hazards model and present the asymptotic results that can be used for inference. Simulation studies are provided in Section 3. In Section 4 we analyze two real data sets, which were originally studied by [Kumar et al. \(2012\)](#) and by [Ringdén et al. \(2012\)](#) using data from the Center for International Blood and Marrow Transplant Research (CIBMTR). Concluding remarks are provided in Section 5.

2 Data and covariate adjusted censoring weight

Let \tilde{T}_i and C_i be the event time and right censoring time for i th individual, respectively. $\epsilon_i \in \{1, \dots, K\}$ indicates the cause of failure. For simplicity, we assume $K = 2$ in this study. Let $T_i = \min(\tilde{T}_i, C_i)$ and $\Delta_i = \mathcal{I}(\tilde{T}_i \leq C_i)$. We observe n independent and identically distributed (*i.i.d.*) data $\{T_i, \Delta_i, \Delta_i \epsilon_i, \mathbf{Z}_i\}$ for $i = 1, \dots, n$, where $\mathbf{Z}_i = (Z_{i1}, \dots, Z_{iq})^\top$ are associated covariates. We assume that $(\tilde{T}_i, \epsilon_i)$ are independent of C_i given covariates of \mathbf{Z}_i . We are interested in modeling the cumulative incidence function of cause 1, $F_1(t; \mathbf{Z})$. Based on [Gray \(1988b\)](#) subdistribution hazard technique, [Fine & Gray \(1999\)](#) proposed a proportional subdistribution hazards model

$$\lambda_1^*(t; \mathbf{Z}) = \frac{-d \log\{1 - F_1(t; \mathbf{Z})\}}{dt} = \lambda_{10}^*(t) \exp\{\boldsymbol{\beta}_0^\top \mathbf{Z}\}. \quad (2.1)$$

There is a direct relationship between the CIF and subdistribution hazard function:

$$F_1(t; \mathbf{Z}) = 1 - \exp\left\{-\left(\int_0^t \lambda_{10}^*(u) du\right) e^{\boldsymbol{\beta}_0^\top \mathbf{Z}}\right\}.$$

Let $N_i^1(t) = I(\tilde{T}_i \leq t, \epsilon_i = 1)$ be the underlying counting process associated with cause 1. For right censored competing risks data, $N_i^1(t)$ and $Y_i^1(t) = 1 - N_i^1(t^-)$ are not fully observed. For a censored individual, it is only observed up to the censoring time C_i . Define $r_i(t) = I\{C_i \geq (\tilde{T}_i \wedge t)\}$. Then, $r_i(t)N_i^1(t)$ and $r_i(t)Y_i^1(t)$ are computable for all time t . Let

$G_C(t; \mathbf{Z}) = P(C \geq t | \mathbf{Z})$ be the conditional censoring distribution. Based on

$$\begin{aligned} E \left\{ \frac{r_i(t)N_i^1(t)}{G_C(T_i \wedge t; \mathbf{Z}_i)} \right\} &= E \left[E \left\{ \frac{r_i(t)N_i^1(t)}{G_C(T_i \wedge t; \mathbf{Z}_i)} \middle| \mathbf{Z}_i \right\} \right] \\ &= E \{ N_i^1(t) | \mathbf{Z}_i \} \frac{E\{r_i(t) | \mathbf{Z}_i\}}{G_C(T_i \wedge t; \mathbf{Z}_i)} \\ &= F_1(t; \mathbf{Z}_i) \end{aligned}$$

FG proposed using an inverse probability of the censoring weighting (IPCW) approach to fit the model (2.1) and proposed an IPCW weight function $w_i^{\text{KM}}(t) = r_i(t)\widehat{G}_C^{\text{KM}}(t)/\widehat{G}_C^{\text{KM}}(T_i \wedge t)$, where $\widehat{G}_C^{\text{KM}}(t)$ is the Kaplan-Meier estimator for the unknown censoring distribution. FG proposed estimating the unknown regression coefficient $\boldsymbol{\beta}$ by solving the score equation

$$\mathbf{U}_{\text{KM}}(\boldsymbol{\beta}) = \sum_i \int_0^\tau \left\{ \mathbf{Z}_i - \frac{\sum_j w_j^{\text{KM}}(u)Y_j^1(u)\mathbf{Z}_j \exp\{\boldsymbol{\beta}^\top \mathbf{Z}_j\}}{\sum_j w_j^{\text{KM}}(u)Y_j^1(u) \exp\{\boldsymbol{\beta}^\top \mathbf{Z}_j\}} \right\} w_i^{\text{KM}}(u)dN_i^1(u) = \mathbf{0},$$

where τ is end of the study time point, and denote the estimate as $\widehat{\boldsymbol{\beta}}_{\text{KM}}$. FG showed that under the regularity conditions and the condition that the censoring distribution is independent of covariates, $\widehat{\boldsymbol{\beta}}_{\text{KM}}$ is consistent for $\boldsymbol{\beta}_0$ and derived large sample properties for $\sqrt{n}(\widehat{\boldsymbol{\beta}}_{\text{KM}} - \boldsymbol{\beta}_0)$ and $\sqrt{n}\{\widehat{\Lambda}_{10}^{\text{KM}}(t) - \Lambda_{10}^*(t)\}$, where the cumulative baseline subdistribution hazard $\Lambda_{10}^*(t) = \int_0^t \lambda_{10}^*(u)du$ is estimated by

$$\widehat{\Lambda}_{10}^{\text{KM}}(t) = \sum_i \int_0^t \frac{w_i^{\text{KM}}(u)dN_i^1(u)}{\sum_j w_j^{\text{KM}}(u)Y_j^1(u) \exp\{\widehat{\boldsymbol{\beta}}_{\text{KM}}^\top \mathbf{Z}_j\}}.$$

It has been shown that in biomedical research studies the censoring time may depend on some of the covariates and the treatment group. To have asymptotically unbiased inferences, we needed to model the censoring distribution. In this study, as suggested by [Fine & Gray \(1999\)](#), we considered the most commonly used Cox proportional hazards model for the censoring distribution,

$$\lambda_C(t; \mathbf{X}) = \lambda_{C0}(t) \exp\{\boldsymbol{\gamma}_0^\top \mathbf{X}\},$$

where \mathbf{X} is the covariates associated with the censoring distribution, which could be a subset covariates of \mathbf{Z} . For a given \mathbf{X} , we estimated the predicted censoring survival probability

$G_C(t; \mathbf{X}) = P(C > t | \mathbf{X})$ by

$$\widehat{G}_C^{\text{COX}}(t; \mathbf{X}) = \exp \left\{ -\widehat{\Lambda}_{C0}(t) \exp \left(\widehat{\boldsymbol{\gamma}}^\top \mathbf{X} \right) \right\}, \quad (2.2)$$

where $\widehat{\boldsymbol{\gamma}}$ is a maximum partial likelihood estimate for $\boldsymbol{\gamma}_0$ and $\widehat{\Lambda}_{C0}(t)$ is a standard Nelson-Aalen type estimator for the cumulative baseline censoring hazard $\Lambda_{C0}(t) = \int_0^t \lambda_{C0}(u) du$. In this study, we considered a covariates-adjusted IPCW weight function

$$w_i^{\text{COX}}(t) = r_i(t) \widehat{G}_C^{\text{COX}}(t; \mathbf{X}_i) / \widehat{G}_C^{\text{COX}}(T_i \wedge t; \mathbf{X}_i).$$

We estimated $\boldsymbol{\beta}$ in model (2.1) by solving the score equation

$$\mathbf{U}_{\text{COX}}(\boldsymbol{\beta}) = \sum_i \int_0^\tau \left\{ \mathbf{Z}_i - \frac{\sum_j w_j^{\text{COX}}(u) Y_j^1(u) \mathbf{Z}_j \exp\{\boldsymbol{\beta}^\top \mathbf{Z}_j\}}{\sum_j w_j^{\text{COX}}(u) Y_j^1(u) \exp\{\boldsymbol{\beta}^\top \mathbf{Z}_j\}} \right\} w_i^{\text{COX}}(u) dN_i^1(u) = \mathbf{0},$$

and denoted the estimate as $\widehat{\boldsymbol{\beta}}_{\text{COX}}$. Then we estimated $\Lambda_{10}^*(t)$ by

$$\widehat{\Lambda}_{10}^{\text{COX}}(t) = \sum_i \int_0^t \frac{w_i^{\text{COX}}(u) dN_i^1(u)}{\sum_j w_j^{\text{COX}}(u) Y_j^1(u) \exp\{\widehat{\boldsymbol{\beta}}_{\text{COX}}^\top \mathbf{Z}_j\}}.$$

Under regularity conditions, it can be shown that $\sqrt{n} \left(\widehat{\boldsymbol{\beta}}_{\text{COX}} - \boldsymbol{\beta}_0 \right)$ converges in distribution to a mean zero Gaussian distribution with an asymptotic variance that can be estimated by

$$\widehat{\Sigma}_{\boldsymbol{\beta}}^{\text{COX}} = n \sum_i \left(\widehat{\mathbf{W}}_{\boldsymbol{\beta},i}^{\text{COX}} \right)^{\otimes 2} = n \left\{ \mathbf{I}_{\text{COX}} \left(\widehat{\boldsymbol{\beta}}_{\text{COX}} \right) \right\}^{-1} \left\{ \sum_i \left(\widehat{\boldsymbol{\xi}}_i^{\text{COX}} + \widehat{\boldsymbol{\psi}}_i^{\text{COX}} \right)^{\otimes 2} \right\} \left\{ \mathbf{I}_{\text{COX}} \left(\widehat{\boldsymbol{\beta}}_{\text{COX}} \right) \right\}^{-1},$$

where $\mathbf{a}^{\otimes 2} = \mathbf{a} \mathbf{a}^\top$ for a column vector \mathbf{a} , $\mathbf{I}_{\text{COX}}(\boldsymbol{\beta}) = -\partial \{ \mathbf{U}_{\text{COX}}(\boldsymbol{\beta}) \} / \partial \boldsymbol{\beta}$, and explicit expressions for $\widehat{\boldsymbol{\xi}}_i^{\text{COX}}$ and $\widehat{\boldsymbol{\psi}}_i^{\text{COX}}$ are given in the Appendix. Similarly, $\sqrt{n} \left\{ \widehat{\Lambda}_{10}^{\text{COX}}(t) - \Lambda_{10}^*(t) \right\}$ converges in distribution to a mean zero Gaussian process with asymptotic variances, which can be estimated by

$$\widehat{\Sigma}_{\Lambda_{10}}^{\text{COX}} = n \sum_i \left\{ \widehat{W}_{\Lambda,i}^{\text{COX}}(t) \right\}^2,$$

where an expression for $\widehat{W}_{\Lambda,i}^{\text{COX}}(t)$ can be found in the Appendix.

For a given set value of covariates, \mathbf{Z} , the predicted CIF of cause 1 can be estimated by

$\widehat{F}_1^{\text{KM}}(t; \mathbf{Z}) = 1 - \exp \left\{ -\widehat{\Lambda}_{10}^{\text{KM}}(t) \exp \left(\widehat{\boldsymbol{\beta}}_{\text{KM}}^{\text{T}} \mathbf{Z} \right) \right\}$ or $\widehat{F}_1^{\text{COX}}(t; \mathbf{Z}) = 1 - \exp \left\{ -\widehat{\Lambda}_{10}^{\text{COX}}(t) \exp \left(\widehat{\boldsymbol{\beta}}_{\text{COX}}^{\text{T}} \mathbf{Z} \right) \right\}$, respectively. [Fine & Gray \(1999\)](#) derived the large sample property for $\sqrt{n} \left\{ \widehat{F}_1^{\text{KM}}(t; \mathbf{Z}) - F_1(t; \mathbf{Z}) \right\}$ when the censoring distribution is independent of the covariates. When the censoring distribution depends on the covariates through a Cox model, by functional Delta method, we can show that $\sqrt{n} \left\{ \widehat{F}_1^{\text{COX}}(t; \mathbf{Z}) - F_1(t; \mathbf{Z}) \right\}$ converges in distribution to a Gaussian process with asymptotic variances, which can be estimated by

$$n \left\{ 1 - \widehat{F}_1^{\text{COX}}(t; \mathbf{Z}) \right\}^2 \sum_i \left\{ \widehat{W}_{F_1, i}^{\text{COX}}(t; \mathbf{Z}) \right\}^2,$$

where

$$\widehat{W}_{F_1, i}^{\text{COX}}(t; \mathbf{Z}) = \exp \left(\widehat{\boldsymbol{\beta}}_{\text{COX}}^{\text{T}} \mathbf{Z} \right) \left\{ \widehat{\Lambda}_{10}^{\text{COX}}(t) \left(\widehat{\mathbf{W}}_{\boldsymbol{\beta}, i}^{\text{COX}} \right)^{\text{T}} \mathbf{Z} + \widehat{W}_{\Lambda, i}^{\text{COX}}(t) \right\}.$$

Resampling techniques can be used to construct confidence bands for $\Lambda_{10}^*(t)$ and $F_1(t; \mathbf{Z})$ ([Lin et al., 1994](#); [Scheike et al., 2008](#)).

3 Simulations

We compared the finite-sample performance of the estimator using the covariate-adjusted censoring weight to the unadjusted estimator using the Kaplan-Meier estimator for the censoring distribution. Two simulation studies were considered to examine the potential bias reduction with the covariate-adjusted censoring weight estimator. For the first study, we had one binary covariate. For the second study, we considered one binary covariate and one continuous covariate. In both studies, we compared the performances of estimators using two weights, $w_i^{\text{KM}}(t)$ and $w_i^{\text{COX}}(t)$, respectively.

3.1 Study 1

The regression model below has one binary covariate Z . Given Z , the cumulative incidence functions are given by

$$F_1(t; Z) = 1 - \left\{ 1 - p \left(1 - e^{-t} \right) \right\}^{\exp(\beta Z)}$$

and

$$F_2(t; Z) = (1 - p)^{\exp(\beta Z)} \left\{ 1 - e^{-t \exp(\beta Z)} \right\},$$

where $p = F_1(\infty|Z = 0)$. We let $p = 0.66$ and Z be a Bernoulli random variable, with a value 1 for half of the sample and 0 for the other half. We set $\beta = 1$ and considered the following three simulation scenarios.

Scenario 1	Censoring times are independent of Z : Generate censoring times from an exponential distribution $\sim \exp(\lambda_C)$ Set $\lambda_C = 0.556$ for 30% censoring, $\lambda_C = 1.342$ for 50% censoring
Scenario 2	Censoring times depend on Z by a Cox model: Generate censoring times from a Cox model, $\lambda_C(t Z) = \lambda_C \exp(\beta_C Z)$ Set $\beta_C = 2.5$ and $\lambda_C = 0.137$ for 30% censoring Set $\beta_C = 2.5$ and $\lambda_C = 0.391$ for 50% censoring
Scenario 3	Censoring times depend on Z , not by a Cox model: $C \sim U(0.25, 4.00)$, if $Z = 0$, $C \sim U(0.07, 1.12)$, if $Z = 1$ for 30% censoring $C \sim U(0.25, 2.00)$, if $Z = 0$, $C \sim U(0.06, 0.46)$, if $Z = 1$ for 50% censoring

For each setting, we simulated 10,000 replicates with sample size of $n = 100$ and 300, respectively. The regression coefficient β was estimated by the methods described in Section 2. We report the average of bias (**Bias**), the sample standard deviation of $\hat{\beta}$ (**SD**), the average of estimated standard error ($\hat{\sigma}$), average of standardized bias (**Std-B** = $E\{|\hat{\beta} - \beta|/\hat{\sigma}\}$), the coverage probability of β , and mean squared error (**MSE**). Table 1 shows the simulation results. We also examined the potential bias of estimating the cumulative baseline subdistribution hazard, $\Lambda_{10}^*(t)$, using both weights at a set of time points, $t = (0.25, 0.5, 0.75, 1.00)^T$. Figure 1 shows the simulation results.

The simulation results show that when the censoring time depends on the covariate (scenario 2 and 3), the unadjusted estimator produces significant biased results, and the estimator using the covariates-adjusted censoring weight provides satisfactory results where the biases are all close to zero. Both estimators give satisfactory variance estimate and have almost identical sample standard deviations (see scenario 2 and 3 in Table 1). Regarding the cumulative subdistribution hazard estimators, estimates using the Cox model adjusted weight have smaller biases compared to those using the unadjusted Kaplan-Meier weight at almost all time points (see Figure 1). Simulation results also indicated that the estimator using the Cox model adjusted weight provides satisfactory results when the Cox model is not the true model for the censoring distribution (see scenario 3 in Table 1 and Figure 1). In scenario 1, where the censoring distribution is independent of covariate Z , both estimators provide

satisfactory results in estimating the covariate effect and cumulative baseline subdistribution hazard function. Both estimators also have almost identical sample standard deviation and similar MSE, which indicate that the potential efficiency losses are minimum when using covariate-adjusted censoring weight.

3.2 Study 2

The regression models below have one binary covariate Z_1 and one continuous covariate Z_2 . Given Z_1 and Z_2 , the cumulative incidence functions are given by

$$F_1(t; Z_1, Z_2) = 1 - \{1 - p(1 - e^{-t})\}^{\exp(\beta_1 Z_1 + \beta_2 Z_2)}$$

and

$$F_2(t; Z_1, Z_2) = (1 - p)^{\exp(\beta_1 Z_1 + \beta_2 Z_2)} \{1 - e^{-t \exp(\beta_1 Z_1 + \beta_2 Z_2)}\}.$$

We let $p = 0.66$, and Z_1 is a Bernoulli random variable, with a value 1 for half of the sample and 0 for the other half. Z_2 is a $N(0, 1)$ random variable. We set $\beta_1 = 1$, $\beta_2 = 0.5$ and considered the following four scenarios.

Scenario 1	Censoring times are independent of Z_1 and Z_2 Generate censoring times from an exponential distribution $\sim \exp(\lambda_C)$ Set $\lambda_C = 0.547$ for 30% censoring, $\lambda_C = 1.352$ for 50% censoring
Scenario 2	Censoring times depend on Z_1 by a Cox model Generate censoring times from $\lambda_C(t \mathbf{Z}) = \lambda_C \exp(\beta_{C1} Z_1)$ Set $\beta_{C1} = 2.5$. Set $\lambda_C = 0.137$ for 30% censoring, $\lambda_C = 0.397$ for 50% censoring
Scenario 3	Censoring times depend on Z_1 and Z_2 by a Cox model Generate censoring times from $\lambda_C(t \mathbf{Z}) = \lambda_C \exp(\beta_{C1} Z_1 + \beta_{C2} Z_2)$ Set $\beta_{C1} = 2.5$, $\beta_{C2} = 2.5$. Set $\lambda_C = 0.082$ for 30% censoring, $\lambda_C = 0.389$ for 50% censoring
Scenario 4	Censoring times depend on Z_1 , not by a Cox model $C \sim U(0.25, 4.00)$, if $Z_1 = 0$, $C \sim U(0.07, 1.14)$, if $Z_1 = 1$ for 30% censoring $C \sim U(0.25, 2.00)$, if $Z_1 = 0$, $C \sim U(0.06, 0.438)$, if $Z_1 = 1$ for 50% censoring

For each setting, we simulated 10,000 replicates with $n = 100$ and 300. The regression coefficients β_1 and β_2 were estimated by the methods described in Section 2. Table 2 shows the simulation results. We also examined the potential bias of estimating the cumulative baseline subdistribution hazard, $\Lambda_{10}^*(t)$, using both weights at a set of time points $t = (0.25, 0.5, 0.75, 1.00)^\top$ for selected scenarios. Figure 2 shows the simulation results.

This simulation study shows similar results as in study 1. The unadjusted estimator produces biased results when the censoring distribution depends on the covariates (scenario 2 to 4), and the estimator using the Cox model adjusted weight provides a good bias reduction. Both estimators give satisfactory variance estimates for both parameters. Regarding the cumulative baseline subdistribution hazard estimates, estimates using the Cox-adjusted weight have smaller biases at almost all points (see Figure 2).

Both simulation studies show that the unadjusted estimator produces significant biased results when the censoring time depends on the covariates and the proposed estimator using covariate adjusted weight works well in bias reduction.

4 Real data examples

4.1 Example 1

We considered data from multiple myeloma patients treated with allogeneic stem cell transplantation from the Center for International Blood and Marrow Transplant Research (CIBMTR) (Kumar et al., 2012). The CIBMTR is comprised of clinical and basic scientists who share data on their blood and bone marrow transplant patients with the CIBMTR Data Collection Center located at the Medical College of Wisconsin. The CIBMTR has a repository of information regarding the results of transplants at more than 450 transplant centers worldwide. The data used in this paper consist of patients transplanted from 1995 to 2005, and we compared the outcomes between transplant periods: 2001-2005 (N=488) versus 1995-2000 (N=375) (Kumar et al., 2012). Two competing events are multiple myeloma relapse and treatment-related mortality (TRM) defined as death without relapse. The CIBMTR study (Kumar et al., 2012) identified that donor type and prior autologous transplantation were associated with relapse or TRM. The variables considered in this example are transplant time period (GP (main interest of the study): 1 for transplanted in 2001-2005 versus 0 for transplanted in 1995-2000), donor type (DNR: 1 for Unrelated or other related donor (N=280)

versus 0 for HLA-identical sibling (N=584)), and prior autologous transplant (PREAUTO: 1 for Auto+Allo transplant (N=399) versus 0 for allogeneic transplant alone (N=465)).

First, we fit a Cox model for the censoring distribution where relapsed or dead individuals are considered as censoring subjects. The hazard ratios (HR) are: HR(GP)=6.42 ($P < 0.0001$); HR(DNR)=0.48 ($p = 0.0018$); HR(PREAUTO)=1.73 ($p = 0.0013$). These results indicate that the censoring distribution depends on the transplant time period, donor type and prior autologous transplantation. Next, we fit a proportional subdistribution hazards model (2.1) with the Kaplan-Meier estimated unadjusted weight and the Cox model adjusted weight, and we computed the predicted cumulative incidence probability for a patient who received an HLA-identical sibling donor allogeneic transplantation in 1995-2000 or in 2001-2005 (see results in Table 3-4 and Figure 3). Both weights give similar estimates for TRM. However, for cancer relapse, the regression estimate of the main treatment effect are $\hat{\beta} = 0.38$ and $\hat{\beta} = 0.54$ by unadjusted weight and Cox model adjusted weight, respectively. At three years after transplant, the differences in cumulative incidence of relapse between late and early transplant (TX) patients are 0.09 (CIF=0.34 for the late TX versus CIF=0.25 for the early TX) and 0.13 (CIF=0.35 for the late TX versus CIF=0.22 for the early TX) by unadjusted weight and Cox model adjusted weight, respectively. The unadjusted weight underestimates the effect size of CIF of relapse by 4% compared to the point estimate using the Cox model adjusted weight (Table 4). Underestimated effect size counts about 14% ($0.04/((0.22+0.35)/2)$) of estimated average CIF, which leads to quite a large relative bias.

4.2 Example 2

We considered another CIBMTR study data set (Ringdén et al., 2012) that consists of 177 myeloma patients who received a reduced-intensity conditioning allogeneic transplantation. Cancer relapse and TRM were two competing risks in this study. 105 patients received prior autologous transplant, and 72 patients received allogeneic transplant alone. We were interested in transplant type effect on relapse and TRM. Let PREAUTO be the indicator of transplant type (1 for Auto+Allo transplant versus 0 for Allogeneic transplant alone). Here the censoring distribution depends on the transplant type ($p = 0.0047$). We fit a proportional subdistribution hazards model (2.1) for PREAUTO with unadjusted weight and Cox model adjusted weight, respectively. For relapse, we have $\hat{\beta}_{\text{COX}} = -0.34(\hat{\sigma} = 0.25)$; $\exp(\hat{\beta}_{\text{COX}}) = 0.71$ and $\hat{\beta}_{\text{KM}} = -0.41(\hat{\sigma} = 0.25)$; $\exp(\hat{\beta}_{\text{KM}}) = 0.66$. Here the Cox model adjusted weight

reduces a relative bias of 17% $((0.41 - 0.34)/0.41)$.

5 Concluding remarks

We have shown that the estimator using the Kaplan-Meier estimated unadjusted inverse probability of censoring weight is not asymptotically unbiased when the censoring distribution depends on the covariates and the biases could be significant for fixed sample sizes. We considered a regression model for the censoring distribution, and we considered using the Cox proportional hazards model and predicted censoring weight for each individual. We have illustrated that the Cox model adjusted weight works well when censoring distribution depends on the covariates, and potential efficiency losses are minimal for both independent and dependent censoring cases. With the transplant data, we determined that the covariate-adjusted weight can be adopted to reduce bias. We are working on an R package, which will be available to the public.

In this study, we only considered using the most common Cox proportional hazards model for the censoring distribution. The Cox model requires a proportional effect (constant effect) for each covariate. However, the proportionality assumption may not be true for some of the covariates. When the Cox model does not fit the data well, one may consider alternative regression models for the censoring distribution. An alternative model-based weight function needs to be considered, an efficient variance estimator needs to be derived, potential bias reduction needs to be studied, and a computing package needs to be further developed as well.

Recently, the inverse probability of censoring weighting (IPCW) technique (Robins & Rotnitzky, 1992) has been used extensively for right-censored survival data and, specifically, for competing risks data. It has been shown that regression modeling of the censoring distribution can be used to improve the efficiency of the IPCW technique (Bickel et al., 1993; Van der Laan & Robins, 2003; Scheike et al., 2008) even if the censoring distribution is independent of the covariates. In this study, we showed that the covariate-adjusted IPCW technique can be used to reduce bias for modeling the subdistribution hazard function when censoring depends on the covariates. In general, the covariate-adjusted IPCW technique should be considered to improve efficiency and reduce bias.

6 Appendix

Here we give a brief derivation for the variance estimation for $\sqrt{n}(\widehat{\boldsymbol{\beta}}_{\text{COX}} - \boldsymbol{\beta}_0)$ and $\sqrt{n}(\widehat{\Lambda}_{10}^{\text{COX}}(t) - \Lambda_{10}^*(t))$, and give explicit expressions for $\widehat{\boldsymbol{\xi}}_i^{\text{COX}}$, $\widehat{\boldsymbol{\psi}}_i^{\text{COX}}$ and $\widehat{W}_{\Lambda,i}^{\text{COX}}(t)$. Let $M_i^1(t) = N_i^1(t) - \int_0^t Y_i^1(u) \exp(\boldsymbol{\beta}_0^\top \mathbf{Z}_i) d\Lambda_{10}^*(u) du$, which is a zero mean martingale for complete data. Assuming the censoring distribution depends on covariates \mathbf{X} through a Cox proportional hazards model where \mathbf{X} could be a subset covariates of \mathbf{Z} ,

$$\lambda_{\text{C}}(t; \mathbf{X}) = \lambda_{\text{C0}}(t) \exp\{\boldsymbol{\gamma}_0^\top \mathbf{X}\}.$$

By Taylor's approximation,

$$\sqrt{n}(\widehat{\boldsymbol{\beta}}_{\text{COX}} - \boldsymbol{\beta}_0) = \sqrt{n} \left\{ \mathbf{I}_{\text{COX}}(\widehat{\boldsymbol{\beta}}_{\text{COX}}) \right\}^{-1} \{ \mathbf{U}_{\text{COX}}(\boldsymbol{\beta}_0) \} + o_p(1), \quad (6.1)$$

where

$$\begin{aligned} \mathbf{U}_{\text{COX}}(\boldsymbol{\beta}_0) &\approx_p \sum_i \int_0^\tau \{ \mathbf{Z}_i - \mathbf{E}_{\text{COX}}(\boldsymbol{\beta}_0, u) \} w_i^{\text{COX}}(u) dM_i^1(u) \\ &= \sum_i \int_0^\tau \{ \mathbf{Z}_i - \mathbf{E}_{\text{COX}}(\boldsymbol{\beta}_0, u) \} r_i(u) \frac{G_{\text{C}}(u; \mathbf{X}_i)}{G_{\text{C}}(T_i \wedge u; \mathbf{X}_i)} dM_i^1(u) \end{aligned} \quad (6.2)$$

$$+ \sum_i \int_0^\tau \left(\frac{\widehat{G}_{\text{C}}^{\text{COX}}(u; \mathbf{X}_i)}{\widehat{G}_{\text{C}}^{\text{COX}}(T_i \wedge u; \mathbf{X}_i)} - \frac{G_{\text{C}}(u; \mathbf{X}_i)}{G_{\text{C}}(T_i \wedge u; \mathbf{X}_i)} \right) \{ \mathbf{Z}_i - \mathbf{E}_{\text{COX}}(\boldsymbol{\beta}_0, u) \} r_i(u) dM_i^1(u) \quad (6.3)$$

$$\mathbf{I}_{\text{COX}}(\boldsymbol{\beta}) = -\partial \{ \mathbf{U}_{\text{COX}}(\boldsymbol{\beta}) \} / \partial \boldsymbol{\beta} \quad (6.4)$$

and

$$\begin{aligned} \mathbf{S}_{\text{COX}}^{(k)}(\boldsymbol{\beta}, u) &= \sum_i w_i^{\text{COX}}(u) Y_i^1(u) \mathbf{Z}_i^{\otimes k} \exp\{\boldsymbol{\beta}^\top \mathbf{Z}_i\}, \text{ for } k = 0, 1, 2 \\ \mathbf{E}_{\text{COX}}(\boldsymbol{\beta}, u) &= \frac{\mathbf{S}_{\text{COX}}^{(1)}(\boldsymbol{\beta}, u)}{S_{\text{COX}}^{(0)}(\boldsymbol{\beta}, u)}. \end{aligned}$$

It has been shown that for given covariates \mathbf{X}_i ([Andersen & Gill, 1982](#)),

$$\begin{aligned}
\widehat{G}_C^{\text{COX}}(t; \mathbf{X}_i) - G_C(t; \mathbf{X}_i) &\approx_p -G_C(t; \mathbf{X}_i) \left\{ e^{(\widehat{\boldsymbol{\gamma}}^\top \mathbf{X}_i)} \widehat{\Lambda}_{C0}(t) - e^{(\boldsymbol{\gamma}_0^\top \mathbf{X}_i)} \Lambda_{C0}(t) \right\} \\
&\approx_p -\widehat{G}_C^{\text{COX}}(t; \mathbf{X}_i) \sum_j \widehat{W}_{\text{COX},j}^C(t; \mathbf{X}_i)
\end{aligned}$$

where

$$\begin{aligned}
\widehat{W}_{\text{COX},j}^C(t; \mathbf{X}_i) &= \widehat{h}(t; \mathbf{X}_i)^\top \{ \mathbf{I}_C(\widehat{\boldsymbol{\gamma}}) \}^{-1} \int_0^\tau \{ \mathbf{X}_j - \mathbf{E}_C(\widehat{\boldsymbol{\gamma}}, u) \} d\widehat{M}_{\text{COX},j}^C(u) \\
&\quad + \int_0^t e^{(\widehat{\boldsymbol{\gamma}}^\top \mathbf{X}_i)} \frac{d\widehat{M}_{\text{COX},j}^C(u)}{S_C^{(0)}(\widehat{\boldsymbol{\gamma}}, u)}
\end{aligned}$$

and

$$\begin{aligned}
\mathbf{S}_C^{(k)}(\boldsymbol{\gamma}, u) &= \sum_j Y_j(u) \mathbf{X}_j^{\otimes k} \exp\{\boldsymbol{\gamma}^\top \mathbf{X}_j\}, \text{ for } k = 0, 1, 2 \\
\mathbf{I}_C(\boldsymbol{\gamma}) &= -\frac{\partial \mathbf{U}_C(\boldsymbol{\gamma})}{\partial \boldsymbol{\gamma}} = \sum_j \int_0^\tau \left\{ \frac{\mathbf{S}_C^{(2)}(\boldsymbol{\gamma}, u)}{S_C^{(0)}(\boldsymbol{\gamma}, u)} - \mathbf{E}_C(\boldsymbol{\gamma}, u)^{\otimes 2} \right\} dN_j^C(u) \\
\mathbf{E}_C(\boldsymbol{\gamma}, u) &= \mathbf{S}_C^{(1)}(\boldsymbol{\gamma}, u) / S_C^{(0)}(\boldsymbol{\gamma}, u) \\
\widehat{h}(t; \mathbf{X}_i) &= \int_0^t e^{(\widehat{\boldsymbol{\gamma}}^\top \mathbf{X}_i)} \{ \mathbf{X}_i - \mathbf{E}_C(\widehat{\boldsymbol{\gamma}}, u) \} d\widehat{\Lambda}_{C0}(u) \\
d\widehat{M}_{\text{COX},j}^C(t) &= dN_j^C(t) - Y_j(t) \exp\{\widehat{\boldsymbol{\gamma}}^\top \mathbf{X}_j\} d\widehat{\Lambda}_{C0}(t)
\end{aligned}$$

Furthermore, we have that

$$\begin{aligned}
\frac{\widehat{G}_C^{\text{COX}}(t; \mathbf{X}_i)}{\widehat{G}_C^{\text{COX}}(T_i \wedge t; \mathbf{X}_i)} &= \frac{G_C(t; \mathbf{X}_i)}{G_C(T_i \wedge t; \mathbf{X}_i)} \\
&= \frac{I(T_i < t)}{\widehat{G}_C^{\text{COX}}(T_i; \mathbf{X}_i) G_C(T_i; \mathbf{X}_i)} \left[G_C(T_i; \mathbf{X}_i) \left\{ \widehat{G}_C^{\text{COX}}(t; \mathbf{X}_i) - G_C(t; \mathbf{X}_i) \right\} \right. \\
&\quad \left. - G_C(t; \mathbf{X}_i) \left\{ \widehat{G}_C^{\text{COX}}(T_i; \mathbf{X}_i) - G_C(T_i; \mathbf{X}_i) \right\} \right] \\
&\approx_p -I(T_i < t) \left\{ \frac{\widehat{G}_C^{\text{COX}}(t; \mathbf{X}_i)}{\widehat{G}_C^{\text{COX}}(T_i; \mathbf{X}_i)} \right\} \sum_j \left\{ \widehat{W}_{\text{COX},j}^C(t; \mathbf{X}_i) - \widehat{W}_{\text{COX},j}^C(T_i; \mathbf{X}_i) \right\}
\end{aligned}$$

Now, it follows that Equation (6.2) can be approximated by $\sum_i \widehat{\boldsymbol{\xi}}_i^{\text{COX}}$, where

$$\begin{aligned}\widehat{\boldsymbol{\xi}}_i^{\text{COX}} &= \int_0^\tau \left\{ \mathbf{Z}_i - \mathbf{E}_{\text{COX}}(\widehat{\boldsymbol{\beta}}_{\text{COX}}, u) \right\} w_i^{\text{COX}}(u) d\widehat{M}_{\text{COX},i}^1(u) \\ d\widehat{M}_{\text{COX},i}^1(t) &= dN_i^1(t) - Y_i^1(t) \exp \left\{ \left(\widehat{\boldsymbol{\beta}}_{\text{COX}} \right)^\top \mathbf{Z}_i \right\} d\widehat{\Lambda}_{10}^{\text{COX}}(t)\end{aligned}$$

and for Equation (6.3), it follows that

$$\begin{aligned}(6.3) &\approx_p \sum_i \int_0^\tau \left\{ \mathbf{Z}_i - \mathbf{E}_{\text{COX}}(\boldsymbol{\beta}_0, t) \right\}^\top w_i^{\text{COX}}(t) dM_i^1(t) I(T_i < t) \sum_j \left\{ \widehat{W}_{\text{COX},j}^{\text{C}}(T_i; \mathbf{X}_i) - \widehat{W}_{\text{COX},j}^{\text{C}}(t; \mathbf{X}_i) \right\} \\ &\approx_p \sum_i \left(\sum_j \left[\int_0^\tau \left\{ \mathbf{Z}_j - \mathbf{E}_{\text{COX}}(\widehat{\boldsymbol{\beta}}_{\text{COX}}, t) \right\}^\top \left\{ \widehat{W}_{\text{COX},i}^{\text{C}}(T_j; \mathbf{X}_j) - \widehat{W}_{\text{COX},i}^{\text{C}}(t; \mathbf{X}_j) \right\} \right. \right. \\ &\quad \left. \left. \times I(T_j < t) w_j^{\text{COX}}(t) d\widehat{M}_{\text{COX},j}^1(t) \right] \right) \\ &= \sum_i \widehat{\boldsymbol{\psi}}_i^{\text{COX}}\end{aligned}$$

Thus,

$$\begin{aligned}\sqrt{n} \left(\widehat{\boldsymbol{\beta}}_{\text{COX}} - \boldsymbol{\beta}_0 \right) &\approx_p \sqrt{n} \left\{ \mathbf{I}_{\text{COX}} \left(\widehat{\boldsymbol{\beta}}_{\text{COX}} \right) \right\}^{-1} \mathbf{U}_{\text{COX}}(\boldsymbol{\beta}_0) \\ &\approx_p \sqrt{n} \left\{ \mathbf{I}_{\text{COX}} \left(\widehat{\boldsymbol{\beta}}_{\text{COX}} \right) \right\}^{-1} \sum_i \left(\widehat{\boldsymbol{\xi}}_i^{\text{COX}} + \widehat{\boldsymbol{\psi}}_i^{\text{COX}} \right)\end{aligned}$$

where $\widehat{\boldsymbol{\xi}}_i^{\text{COX}}$ is the major term in the variance estimation. Next,

$$\begin{aligned}
\sqrt{n} \left\{ \widehat{\Lambda}_{10}^{\text{COX}}(t) - \Lambda_{10}^*(t) \right\} &= \sqrt{n} \int_0^t \left\{ \frac{\sum_i w_i^{\text{COX}}(u) dN_i^1(u)}{S_{\text{COX}}^{(0)}(\widehat{\beta}_{\text{COX}}, u)} - \frac{\sum_i w_i^{\text{COX}}(u) dN_i^1(u)}{S_{\text{COX}}^{(0)}(\beta_0, u)} \right\} \\
&+ \sqrt{n} \int_0^t \left\{ \frac{\sum_i w_i^{\text{COX}}(u) dN_i^1(u)}{S_{\text{COX}}^{(0)}(\beta_0, u)} - d\Lambda_{10}^*(u) \right\} \\
&\approx_p -\sqrt{n} \int_0^t \mathbf{E}_{\text{COX}}(\beta_0, u)^\top \frac{\sum_i w_i^{\text{COX}}(u) dN_i^1(u)}{S_{\text{COX}}^{(0)}(\beta_0, u)} (\widehat{\beta}_{\text{COX}} - \beta_0) \\
&+ \sqrt{n} \int_0^t \frac{\sum_i w_i^{\text{COX}}(u) dM_i^1(u)}{S_{\text{COX}}^{(0)}(\beta_0, u)} \\
&\approx_p \sqrt{n} \sum_i \left\{ \widehat{W}_{\Lambda,1,i}^{\text{COX}}(t) - \widehat{W}_{\Lambda,2,i}^{\text{COX}}(t) \right\} \\
&= \sqrt{n} \sum_i \widehat{W}_{\Lambda,i}^{\text{COX}}(t),
\end{aligned}$$

where

$$\begin{aligned}
\widehat{W}_{\Lambda,1,i}^{\text{COX}}(t) &= \int_0^t \frac{w_i^{\text{COX}}(u) d\widehat{M}_{\text{COX},i}^1(u)}{S_{\text{COX}}^{(0)}(\widehat{\beta}_{\text{COX}}, u)}, \\
\widehat{W}_{\Lambda,2,i}^{\text{COX}}(t) &= \sum_j \int_0^t \mathbf{E}_{\text{COX}}(\widehat{\beta}_{\text{COX}}, u)^\top \frac{w_j^{\text{COX}}(u) dN_j^1(u)}{S_{\text{COX}}^{(0)}(\widehat{\beta}_{\text{COX}}, u)} \left\{ \mathbf{I}_{\text{COX}}(\widehat{\beta}_{\text{COX}}) \right\}^{-1} \left(\widehat{\xi}_i^{\text{COX}} + \widehat{\psi}_i^{\text{COX}} \right).
\end{aligned}$$

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Table 1: Simulation results for bias reduction with a single binary covariate ($\beta = 1$).

Scenario	N	Cens.	β	Unadjusted weight			Cox model adjusted weight				
				Bias (Std-B)	$\widehat{\sigma}$ (SD)	Coverage	MSE	Bias (Std-B)	$\widehat{\sigma}$ (SD)	Coverage	MSE
1	100	30%	β	0.0081 (0.0291)	0.2759 (0.2798)	0.9494	0.0783	0.0089 (0.0319)	0.2757 (0.2797)	0.9495	0.0783
		50%	β	0.0145 (0.0430)	0.3329 (0.3384)	0.9488	0.1147	0.0153 (0.0450)	0.3331 (0.3387)	0.9489	0.1149
	300	30%	β	0.0010 (0.0061)	0.1584 (0.1586)	0.9499	0.0252	0.0013 (0.0079)	0.1580 (0.1583)	0.9512	0.0251
		50%	β	0.0030 (0.0155)	0.1899 (0.1909)	0.9498	0.0365	0.0033 (0.0174)	0.1898 (0.1908)	0.9503	0.0364
2	100	30%	β	-0.1119 (0.3592)	0.3041 (0.3116)	0.9278	0.1096	0.0050 (0.0160)	0.3080 (0.3123)	0.9487	0.0976
		50%	β	-0.1162 (0.2719)	0.4075 (0.4271)	0.9359	0.1959	0.0175 (0.0398)	0.4217 (0.4388)	0.9501	0.1929
	300	30%	β	-0.1244 (0.7109)	0.1741 (0.1750)	0.8865	0.0461	0.0042 (0.0236)	0.1765 (0.1762)	0.9503	0.0311
		50%	β	-0.1336 (0.5596)	0.2306 (0.2346)	0.9063	0.0729	0.0055 (0.0226)	0.2393 (0.2409)	0.9511	0.0581
3	100	30%	β	-0.1020 (0.3383)	0.2937 (0.3016)	0.9312	0.1013	0.0045 (0.0145)	0.3036 (0.3096)	0.9503	0.0958
		50%	β	-0.0988 (0.2507)	0.3797 (0.3941)	0.9350	0.1650	0.0168 (0.0401)	0.4057 (0.4198)	0.9487	0.1765
	300	30%	β	-0.1099 (0.6491)	0.1685 (0.1692)	0.8927	0.0407	0.0026 (0.0147)	0.1748 (0.1749)	0.9535	0.0306
		50%	β	-0.1088 (0.5009)	0.2152 (0.2173)	0.9152	0.0591	0.0072 (0.0310)	0.2309 (0.2332)	0.9508	0.0544

Bias=Average of bias of $\widehat{\beta}$; SD=Sample standard deviation of $\widehat{\beta}$; $\widehat{\sigma}$ =Average of estimated standard error; Std-B = $E\{|\widehat{\beta} - \beta|/\widehat{\sigma}\}$;
MSE=Mean squared error.

Table 2: Simulation results for biases using 1 binary and 1 continuous covariate ($\beta_1 = 1, \beta_2 = 0.5$).

Scenario	N	Cens.	β	Unadjusted weight				Cox model adjusted weight			
				Bias (Std-B)	$\hat{\sigma}(SD)$	Coverage	MSE	Bias (Std-B)	$\hat{\sigma}(SD)$	Coverage	MSE
1	100	30%	β_1	0.0148 (0.0521)	0.2793 (0.2849)	0.9462	0.0814	0.0154 (0.0542)	0.2793 (0.2847)	0.9463	0.0813
			β_2	0.0130 (0.0873)	0.1424 (0.1483)	0.9381	0.0222	0.0130 (0.0876)	0.1424 (0.1482)	0.9394	0.0221
	50%	β_1	0.0293 (0.0847)	0.3353 (0.3463)	0.9474	0.1207	0.0299 (0.0863)	0.3355 (0.3464)	0.9468	0.1209	
		β_2	0.0143 (0.0812)	0.1676 (0.1763)	0.9362	0.0313	0.0146 (0.0827)	0.1676 (0.1764)	0.9362	0.0313	
	300	30%	β_1	0.0039 (0.0242)	0.1595 (0.1608)	0.9498	0.0259	0.0041 (0.0254)	0.1592 (0.1602)	0.9495	0.0257
			β_2	0.0042 (0.0521)	0.0807 (0.0813)	0.9494	0.0066	0.0043 (0.0535)	0.0806 (0.0811)	0.9485	0.0066
5%0	β_1	β_1	0.0062 (0.0327)	0.1898 (0.1883)	0.9525	0.0355	0.0063 (0.0333)	0.1897 (0.1881)	0.9529	0.0354	
		β_2	0.0063 (0.0661)	0.0941 (0.0958)	0.9448	0.0092	0.0065 (0.0674)	0.0941 (0.0957)	0.9438	0.0092	
2	100	30%	β_1	-0.938 (0.2949)	0.3036 (0.3180)	0.9243	0.1099	0.0119 (0.0370)	0.3101 (0.3220)	0.9421	0.1038
			β_2	0.0273 (0.1778)	0.1462 (0.1536)	0.9355	0.0243	0.0160 (0.1037)	0.1468 (0.1540)	0.9357	0.0240
	50%	β_1	-0.955 (0.2241)	0.4031 (0.4262)	0.9325	0.1908	0.0179 (0.0409)	0.4181 (0.4385)	0.9414	0.1926	
		β_2	0.0247 (0.1349)	0.1740 (0.1831)	0.9333	0.0341	0.0142 (0.0771)	0.1748 (0.1839)	0.9354	0.0340	
	300	30%	β_1	-1.089 (0.6286)	0.1734 (0.1733)	0.9002	0.0419	0.0049 (0.0277)	0.1771 (0.1769)	0.9514	0.0313
			β_2	0.0174 (0.2056)	0.0827 (0.0845)	0.9409	0.0074	0.0050 (0.0589)	0.0831 (0.0848)	0.9428	0.0072
50%	β_1	β_1	-1.136 (0.4956)	0.2271 (0.2291)	0.9153	0.0654	0.0049 (0.0205)	0.2363 (0.2380)	0.9516	0.0566	
		β_2	0.0142 (0.1418)	0.0979 (0.1002)	0.9407	0.0102	0.0027 (0.0273)	0.0984 (0.1004)	0.9434	0.0101	
3	100	30%	β_1	-0.299 (0.0945)	0.3061 (0.3168)	0.9418	0.1012	0.0145 (0.0459)	0.3070 (0.3149)	0.9466	0.0994
			β_2	-0.527 (0.2665)	0.1898 (0.1979)	0.9224	0.0419	0.0014 (0.0069)	0.1934 (0.1992)	0.9422	0.0397
	50%	β_1	-0.692 (0.1711)	0.3872 (0.4042)	0.9366	0.1682	0.0163 (0.0398)	0.3935 (0.4090)	0.9441	0.1676	
		β_2	-0.963 (0.3695)	0.2501 (0.2605)	0.9087	0.0771	-0.0007 (0.0025)	0.2630 (0.2728)	0.9380	0.0744	
	300	30%	β_1	-0.473 (0.2687)	0.1750 (0.1762)	0.9363	0.0333	0.0018 (0.0102)	0.1749 (0.1758)	0.9486	0.0309
			β_2	-0.631 (0.5767)	0.1079 (0.1094)	0.8987	0.0159	-0.0016 (0.0148)	0.1097 (0.1101)	0.9494	0.0121
50%	β_1	β_1	-0.874 (0.3920)	0.2196 (0.2231)	0.9266	0.0574	0.0046 (0.0206)	0.2230 (0.2251)	0.9489	0.0507	
		β_2	-1.035 (0.7300)	0.1404 (0.1418)	0.8698	0.0308	0.0022 (0.0146)	0.1488 (0.1504)	0.9451	0.0226	
4	100	30%	β_1	-0.773 (0.2578)	0.2934 (0.2999)	0.9358	0.0959	0.0149 (0.0481)	0.3035 (0.3089)	0.9476	0.0956
			β_2	0.0251 (0.1642)	0.1441 (0.1526)	0.9346	0.0239	0.0146 (0.0952)	0.1444 (0.1529)	0.9361	0.0236
	50%	β_1	-0.731 (0.1868)	0.3799 (0.3913)	0.9402	0.1584	0.0265 (0.0636)	0.4040 (0.4163)	0.9494	0.1740	
		β_2	0.0248 (0.1390)	0.1708 (0.1783)	0.9361	0.0324	0.0150 (0.0840)	0.1713 (0.1787)	0.9366	0.0322	
	300	30%	β_1	-0.924 (0.5450)	0.1672 (0.1695)	0.9076	0.0372	0.0055 (0.0314)	0.1735 (0.1750)	0.9485	0.0306
			β_2	0.0142 (0.1703)	0.0816 (0.0833)	0.9439	0.0071	0.0028 (0.0330)	0.0818 (0.0834)	0.9462	0.0070
50%	β_1	β_1	-0.877 (0.4128)	0.2149 (0.2124)	0.9327	0.0528	0.0101 (0.0446)	0.2290 (0.2269)	0.9544	0.0516	
		β_2	0.0123 (0.1255)	0.0960 (0.0977)	0.9434	0.0097	0.0023 (0.0232)	0.0963 (0.0982)	0.9455	0.0096	

Bias=Average of bias of $\hat{\beta}$; SD=Sample standard deviation of $\hat{\beta}$; $\bar{\sigma}$ =Average of estimated standard error; Std-B = $E\{|\hat{\beta} - \beta|/\hat{\sigma}\}$;
MSE=Mean squared error.

Table 3: Fit a proportional subdistribution hazards model.

	Unadjusted weight	Cox model adjusted weight
Variable	$\hat{\beta}$; $\exp(\beta)$ (95% CI); P	$\hat{\beta}$; $\exp(\beta)$ (95% CI); P
RELAPSE		
GP	0.38; 1.47(1.16-1.86); 0.0017	0.54; 1.71(1.34-2.20); < 0.0001
DNR	0.39; 1.48(1.18-1.86); 0.0007	0.35; 1.42(1.13-1.78); 0.0027
PREAUTO	0.41; 1.51(1.19-1.91); 0.0007	0.42; 1.53(1.21-1.93); 0.0004
TRM		
GP	-0.59; 0.55(0.42-0.73); < 0.0001	-0.56; 0.57(0.43-0.75); < 0.0001
DNR	0.57; 1.76(1.38-2.25); < 0.0001	0.55; 1.73(1.35-2.20); < 0.0001
PREAUTO	-0.38; 0.68(0.51-0.91); 0.0099	-0.37; 0.69(0.52-0.92); 0.0117

Table 4: Predicted CIF of relapse and TRM for a patient who received an HLA-identical sibling donor and allogeneic along transplantation

	Unadjusted Weight			Cox model adjusted Weight		
	1995-2000	2001-2005		1995-2000	2001-2005	
Time	\hat{F}_1 (95% CI)	\hat{F}_2 (95% CI)	$ \hat{F}_1 - \hat{F}_2 $	\hat{F}_1 (95% CI)	\hat{F}_2 (95% CI)	$ \hat{F}_1 - \hat{F}_2 $
RELAPSE						
1 Year	0.16 (0.13-0.19)	0.23 (0.18-0.27)	0.07	0.15 (0.13-0.17)	0.24 (0.18-0.30)	0.09
3 Year	0.25 (0.20-0.29)	0.34 (0.28-0.40)	0.09	0.22 (0.20-0.25)	0.35 (0.28-0.42)	0.13
5 Year	0.29 (0.24-0.34)	0.40 (0.33-0.46)	0.11	0.26 (0.24-0.30)	0.41 (0.33-0.49)	0.15
TRM						
1 Year	0.38 (0.32-0.43)	0.23 (0.18-0.28)	0.15	0.37 (0.34-0.41)	0.23 (0.17-0.29)	0.14
3 Year	0.42 (0.37-0.48)	0.26 (0.20-0.32)	0.16	0.42 (0.38-0.46)	0.27 (0.20-0.33)	0.15
5 Year	0.44 (0.38-0.49)	0.27 (0.21-0.33)	0.17	0.43 (0.39-0.47)	0.27 (0.21-0.34)	0.16

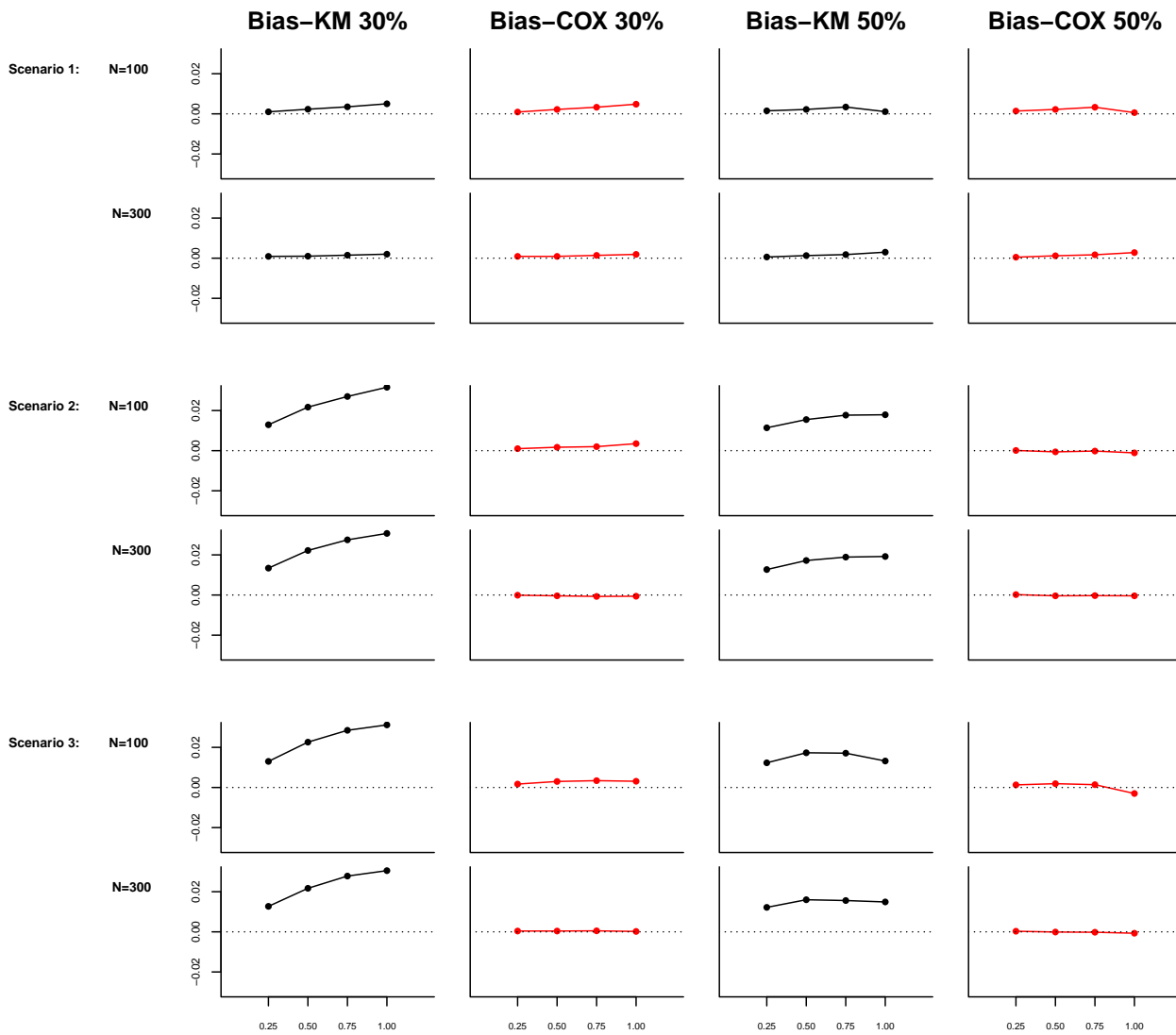


Figure 1: Simulation results (1 covariate) for biases of cumulative baseline subdistribution hazards at $t = (0.25, 0.5, 0.75, 1)^T$.

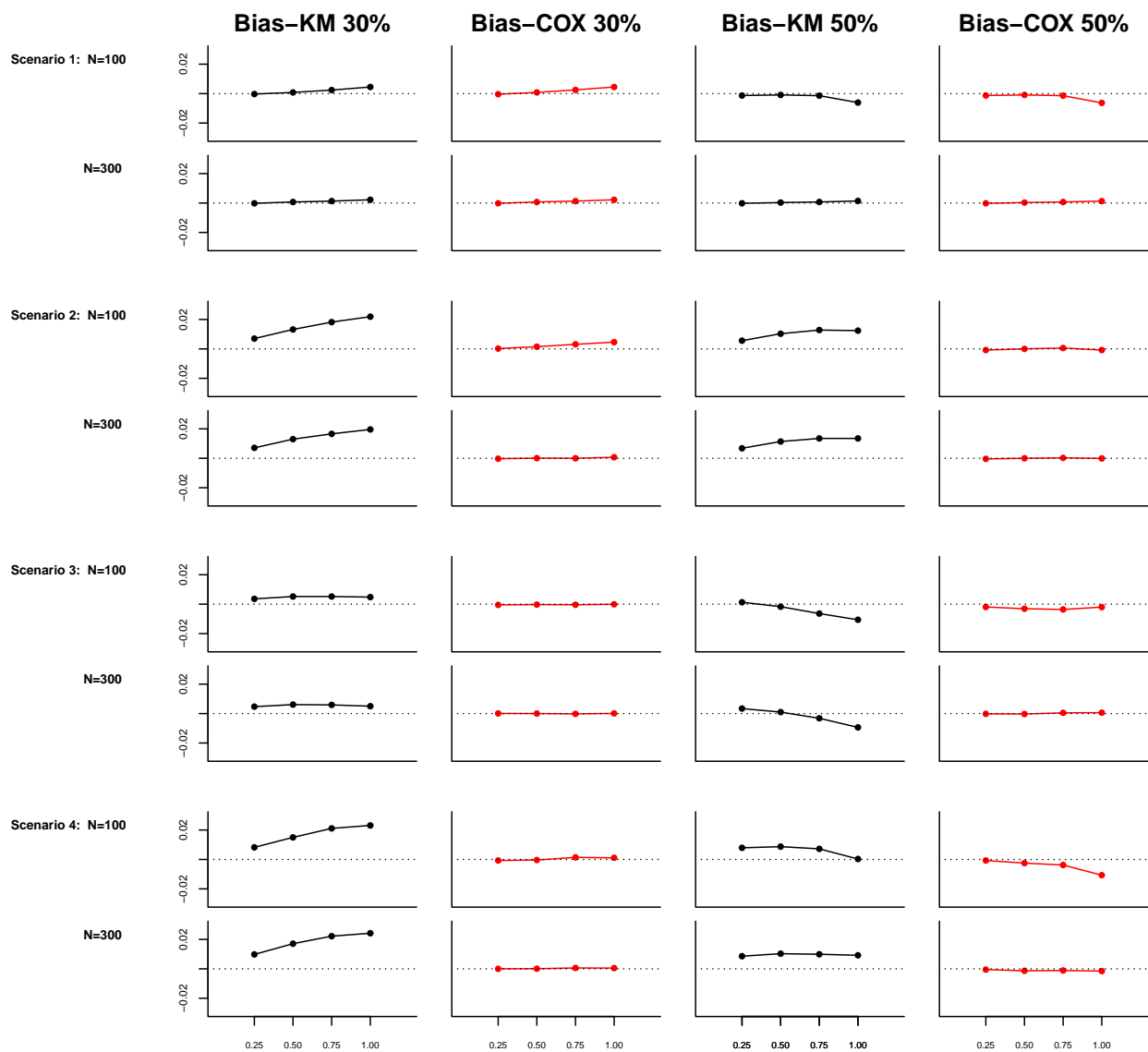


Figure 2: Simulation results (2 covariates) for biases of cumulative baseline subdistribution hazards at $t = (0.25, 0.5, 0.75, 1.00)^T$.

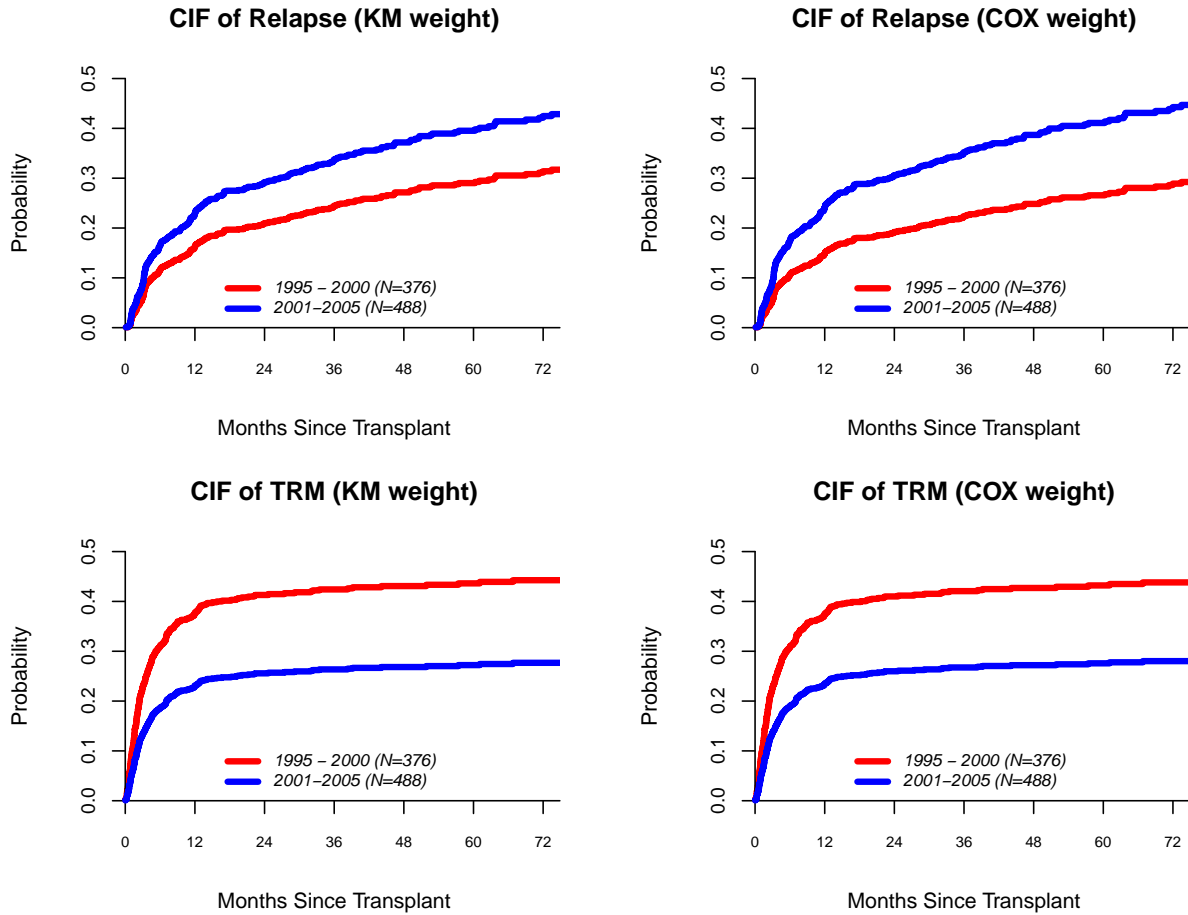


Figure 3: Predicted cumulative incidence probability of relapse and TRM for a patient who received an HLA-identical sibling donor allogeneic transplantation.