Using $\mathbf{icenReg}$ for interval censored data in $\mathbf R$

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Introduction

This manual is meant to provide an introduction to using **icenReg** to analyze interval censored data. It is written with expectation that the reader is familiar with basic survival analysis methods. Familiarity with the Kaplan Meier curves and Cox proportional hazards model should be sufficient.

1.1 Interval Censoring

Interval censoring occurs when a response is known only up to an interval. A classic example is testing for diseases at a doctor's clinic; if a subject tests negative at t_1 and positive at t_2 , all that is known is that the subject acquired the disease in (t_1, t_2) , rather than an exact time. Other classic examples include examining test mice for tumors after sacrifice (results in current status or case I interval censored data, in which all observations are either left or right censored, as opposed to the more general case II), customer choice models in economics (customers are presented a price for a product and chose to purchase or not, researcher wants to know distribution of maximum spending amount; this results in current status data again), data reduction methods for sensor analyses (to reduce load on sensor system, message is intentionally surpressed if outcome is in an expected region) and data binning (responses reported only up to an interval, in some cases to keep the subjects anonymous, in some cases to reduce size of data).

Often interval censoring is ignored in analysis. For example, age is usually reported only up to the year, rather than as a continuous variable. In the case that these intervals are relatively short, the bias introduced by ignoring the interval censoring may be small enough to be safely ignored. However, in the case that the width of intervals is non trivial, statistical methods that

account for this should be used for reliable analysis.

Standard notation for interval censoring is that each observation contains a response interval $[l_i, r_i]$ such that the true event time is known to have occurred within. Note that this allows for uncensored observations $(l_i = r_i)$, right censored $(r_i = \infty)$, left censored $(l_i = 0)$ or none of the above $(0 < l_i < r_i < \infty)$.

In **icenReg**, the response value is allowed to be interval censored. If our data contains the values L and R, representing the left and right sides of the response interval, we can pass our response to a regression model using either

```
cbind(L, R)
Surv(L, R, type = "interval2")
```

It is worth nothing that other R packages, specifically for non-parametric estimation, allow you to declare whether the response intervals are open, closed or a combination of partially opened, for example $[l_i, r_i)$. In **icenReg**, it is always assumed that the intervals are closed.

1.2 Classic Estimators

The topic of interval censoring began emerging in the field of survival analysis. Although it is now considered in other fields of study (such as *tobit regression*), at this time **icenReg** focusses on survival models.

One of the earliest models is the Non-Parametric Maximum Likelihood Estimator (NPMLE), also referred to as Turnbull's Estimator. This is a generalization of the Kaplan Meier curves (which is a generalization of the empirical distribution function) that allows for interval censoring. Unlike the Kaplan Meier curves, the solution is not in closed form and several algorithms have been proposed for efficient computation. A special topic regarding the NPMLE is the bivariate NPMLE; this is for the special case of two interval censored outcomes, in which the researcher wants a non-parametric estimator of the joint distribution. This is especially computationally intense as the number of parameters can be up to n^2 .

Semi-parametric models exist in the literature as well; two classic regression models fit by **icenReg** are the Cox-PH model and the proportional odds model. The well known Cox-PH, or proportional hazards regression model, has the property that

$$h(t|X,\beta) = h_o(t)e^{X^T\beta}$$

where $h(t|X,\beta)$ is the hazard rate conditional on covariates X and regression parameters β , with h_o as the baseline hazard function. This relation is equivalent to

$$S(t|X,\beta) = S_o(t)^{e^{X^T \beta}}$$

where $S(t|X,\beta)$ is the conditional survival and $S_o(t)$ is the baseline survival function.

The less known proportional odds model can be expressed as

$$Odds(S(t|X,\beta)) = e^{X^T \beta} Odds(S_o(t))$$

or

$$\frac{S(t|X,\beta)}{1 - S(t|X,\beta)} = e^{X^T \beta} \frac{S_o(t)}{1 - S_o(t)}$$

Unlike the special example of the Cox PH model with right-censored data, the baseline parameters must be estimated concurrently with the regression parameters. The model can be kept semi-parametric (i.e. no need to decide on a parametric baseline distribution) by using the Turnbull estimator, modified to account for the given regression model, as the baseline distribution. The semi-parametric model can be computationally very difficult, as the number of baseline parameters can be quite high (up to n), which must follow shape constraints (i.e. either a set of probability masses or a cumulative hazard function, which must be strictly increasing) and there is no closed form solution to either regression or baseline parameters.

Fully parametric models exist as well and can be calculated using fairly standard algorithms. There are slight complications in that the interval censoring causes the log likelihood function to be non-concave. However, only slight modifications are required to adress this issue. In practice, fully-parametric models should be used with caution; the lack of observed values means that model inspection can be quite difficult; there are no histograms, etc., to be made. As such, even if fully parametric models are to be used for the final analysis, it is strongly encouraged to use semi-parametric models at least for model inspection. **icenReg** fits both proportional odds and proporitonal hazard models for interval censored data.

Another common regression model for survival data is the accelerated failure time model (AFT). At this time, this option is not available for

interval censored data. However, this model can be fit for interval censored data using **survival**'s **survreg** function.

1.3 Models fit with icenReg

The author's motivation for building **icenReg** was to build a set of fast, reliable tools for analysts' to use on applied data. At this time, the following set of models can be fit (name in paratheses is function call in **icenReg**):

- NPMLE (ic_sp can be used to fit univariate NPMLE, alternatively ICNPMLE can be used to fit univariate or bivariate NPMLE)
- Semi-parametric model (ic_sp, with options model = "ph" for porportional hazards, "po" for proportional odds)
- Fully parametric model (ic_par, in addition to model option, also have a choice of dist, with options "exponential", "gamma", "weibull", "lnorm", "loglogistic" and "generalgamma")

In addition, icenReg includes various diagnostic tools. These include

- Plots for diagnosising baseline distribution (diag_baseline)
- Plots for diagnosising covariate effects (diag_covar)
- Cross validation via multiple imputations (icenReg_cv)

1.4 Data Examples in icenReg

The package includes 4 sources of example data: two functions that simulate data and two sample data sets. The simulation functions are $simIC_weib$, which simulates interval censored regression data with a Weibull baseline distribution and simBVCen, which simulates bivariate interval censored data. The sample data sets are miceData, which contains current status data regarding lung tumors from two groups of mice and essIncData, which includes data from the European Social Survey. In this case, wages were only recorded up to an interval to protect the identity of the subjects. The dataset $essIncData_small$ is a smaller subset of essIncData (n = 500 instead of 6,712), which is used in many of the examples only so that CRAN's testing of the package runs quicker. In practice, using all these models on n = 6,712 is trival to do, rarely taking more than a few seconds even on a slower laptop.

Fitting Models using icenReg

An important note about **icenReg** is that in all models, it is assumed that the response interval is **closed**, i.e. the event is known to have occurred within $[t_1, t_2]$, compared with $[t_1, t_2)$, (t_1, t_2) , etc. This is of no consequence for fully parametric models, but does mean the solutions may differ somewhat in comparison with semi- and non-parametric models that allow differnt configurations of open and closed response intervals.

2.1 Non-parametric models

As noted earlier, for univariate interval censored data, the model may be fit with either ic_sp or ICNPMLE. For large datasets (i.e. n > 50,000), ic_sp will become faster than ICNPMLE. In addition, ic_sp can readily be provided to the plot method. For bivariate data, ICNPMLE is currently the only choice.

If the data set is relatively small and the user is interested in non-parametric tests, such as the log-rank statistic, we actually advise using the **interval** package, as this provides several testing functions. However, **icenReg** is several fold faster than **interval**, so if large datasets are used (i.e. n > 1,000), the user may have no choice but to use **icenReg**.

To fit an NPMLE model for interval censored data, we will consider the miceData provided in icenReg. This dataset contains three variables: 1, u and grp. 1 and u represent the left and right side of the interval containing the event time (note: data is current status) and grp is a group indicator with two categories.

If we separate the data into two datasets, i.e.

```
ge.data <- miceData[miceData$grp == "ge", ]</pre>
```

```
ce.data <- miceData[miceData$grp == "ce", ]</pre>
```

We can then fit the NPMLE by calling the interval censored semiparametric model, but supplying no covariates. This can be done by

```
ge.fit <- ic_sp(cbind(1, u) ~ 0, data = ge.data)
ce.fit <- ic_sp(cbind(1, u) ~ 0, data = ce.data)</pre>
```

Because the objects returned by ic_sp are intended to describe semiparametric models and thus focus on the regression parameters. For the NPMLE, we need the information about the survival curve. We can extract the estimated survival curves by getSCurves

```
ge.sc <- getSCurves(ge.fit)
ce.sc <- getSCurves(ce.fit)</pre>
```

We can then plot and examine the NPMLE's for the two different groups using plot and lines

```
plot(ge.sc, xlab = 'Time',
   ylab = 'Estimated Survival', col = 'blue')
lines(ce.sc, col = 'red')
legend('bottomleft', legend = c('ge', 'ce'),
   col = c('blue', 'red'), lty = 1)
```

Looking at figure 2.1, we can see a unique feature about the NPMLE for interval censored data. That is, there are *two* lines used to represent the survival curve. This is because with interval censored data, the NPMLE is not always unique (in fact, it usually is not); any curve that lies between the two lines has the same likelihood. For example, any curve that lies between the two blues lines in figure 2.1 maximizes the likelihood associated with "ge" group of mice.

Formal statistical tests using the NPMLE are not currently supported by **icenReg**. We recommend using the **interval** package for this.

As noted earlier, ICNPMLE can be used to fit the univariate or bivariate NPMLE. However, there currently are no methods for plotting or testing on these fits. Because of this, we will not cover the use of this function here. The curious user is welcome to view ?ICNPMLE.

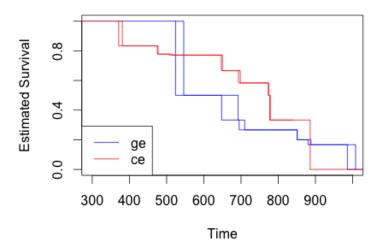


Figure 2.1: NPMLE's for both groups in miceData

2.2 Semi-parametric models

Semi-parametric models can be fit with ic_sp function. This function follows standard regression syntax. As an example, we will fit the essIncData dataset. In this dataset, have income from the European Social Survey, which includes income reported up to an interval (to preserve the identity of the subjects). Within each country, the intervals are disjoint, but between countries there is penty of overlap.

We fit the model below. Note that this may be time consuming, as the semi-parametric model is somewhat computationally intense and we are taking bs_samples bootstrap samples of the estimator.

```
fit_ph <- ic_sp(cbind(inc_1, inc_u) ~ cntry + eduLevel,
    bs_samples = 500, data = essIncData)

fit_po <- ic_sp(cbind(inc_1, inc_u) ~ cntry + eduLevel,
    bs_samples = 500, model = "po", data = essIncData)</pre>
```

The first model by default fits a Cox-PH model, while the second fits

a proportional odds model. We can look at the results using either the summary function, or just directly looking at the results (the output is the same).

```
> fit_po
```

```
Model: Proportional Odds
Baseline: semi-parametric
Call: ic_sp(formula = cbind(inc_l, inc_u) ~ cntry + eduLevel,
    data = essIncData, model = "po", bs_samples = 500)
                 Estimate Exp(Est) Std.Error z-value p
cntryPoland
                    2.6190
                             13.730
                                      0.05724
                                               45.760 0
cntryRussia
                   0.7940
                              2.212
                                      0.05450
                                               14.570 0
cntrySlovakia
                   0.6351
                              1.887
                                      0.06766
                                                9.387 0
eduLevel[12,16)
                   1.0050
                              2.732
                                      0.05186
                                               19.380 0
eduLevel[16,Inf)
                              6.390
                                               31.570 0
                   1.8550
                                      0.05875
```

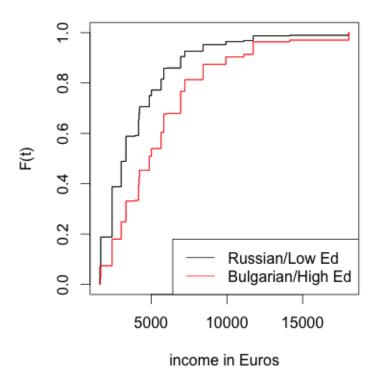
final llk = -12483.03 Iterations = 18 Bootstrap Samples = 500

For the semi-parametric models, bootstrap samples are used for inference on the regression parameters. The reason for this is that as far as we know, the limiting distribution of the baseline distribution is currently undefined. In fact, to our knowledge, even using the bootstrap error estimates for the baseline distribution is not valid. Because the regression parameters cannot be seperated in the likelihood function, using the negative inverse of the Hessian for the regression standard errors is not generally valid. However, it has been shown that using the bootstrap for inference on the regression parameters leads to valid inference.

We can use these fits to create plots as well. The plot function will plot the estimated survival curves or CDF for subjects with the set of covariates provided in the newdata argument. If newdata is left equal to NULL, the baseline survival function will be plotted.

If we wanted to plot the estimated CDF for an individual with between zero and 12 years of school from Russia against the estimated CDF for an individual from Bulgaria with 16+ years of school, this can be done with

```
newdata <- data.frame(cntry = c("Russia", "Bulgaria"),</pre>
```



```
eduLevel = c("[0,12)", "[16,Inf)") )

rownames(newdata) <- c("Russian/Low Ed",
    "Bulgarian/High Ed")

plot(fit_po, newdata, fun = "cdf",
    lgdLocation = "bottomright", xlab = "income in Euros")</pre>
```

2.3 Parametric Models

We can fit parametric models in **icenReg** using the **ic_par** function. The syntax is essentially the same as above, expect that the user needs to spec-

ify dist, the parametric family that the baseline distribution belongs to. The current choices are "exponential", "weibull" (default), "gamma", "lnorm", "loglogistic" and "generalgamma" (generalized gamma distribution). The user must also select model = "ph" or "po", just as in the semi-parametric model.

It is not necessary to specify bs_samples for parametric models, as inference is done using the asymptotic normality of the estimators. Fitting a parametric model is typically faster than the semi-parametric model, even if no bootstrap samples are taken for the semi-parametric model. This is because the fully-parametric model is of lower dimensional space without constraints.

Suppose we wanted to fit a proportional odds model to the essIncData data with a log-normal distribution. This could be fit by

```
fit_po_ln <- ic_par(cbind(inc_l, inc_u) ~ eduLevel + cntry,
  data = essIncData, model = "po")</pre>
```

We can examine the regression coefficients in the same way as with the semi-parametric model.

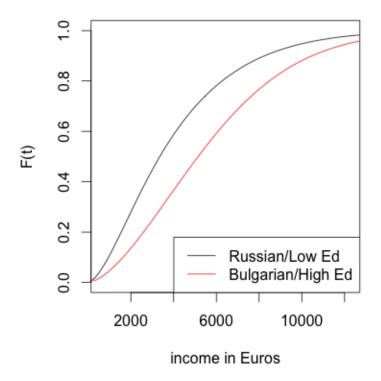
```
> fit_po_ln
```

Model: Proportional Odds

Baseline: weibull

	Estimate	<pre>Exp(Est)</pre>	Std.Error	z-value p
log_shape	0.4725	1.604	0.01038	45.53 0
log_scale	8.6930	5963.000	0.00810	1073.00 0
eduLevel[12,16)	0.8835	2.419	0.04690	18.84 0
eduLevel[16,Inf)	1.7060	5.506	0.06357	26.83 0
cntryPoland	2.4540	11.640	0.06637	36.98 0
cntryRussia	0.8047	2.236	0.05358	15.02 0
cntrvSlovakia	0.6911	1.996	0.06120	11.29 0

final 11k = -14732.97Iterations = 15



We can also examine the survival/cdf plots in the same way.

```
plot(fit_po_ln, newdata, fun = "cdf",
  lgdLocation = "bottomright", xlab = "income in Euros")
```

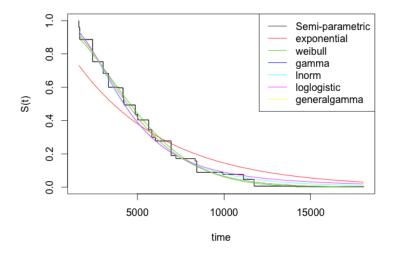
Inspecting model fit

3.1 Examining Baseline Distribution

Althought the semi-parametric model is more flexible, and thus more robust to unusual baseline distributions, there are many reasons one may decide to use a parametric model instead. One reasons is that, as stated earlier, we are not aware of any general distributional theory regarding the baseline distribution, outside of the univariate case with case I interval censored data. Even in this case, the estimator is highly inefficient, observing convergence rates of $n^{1/3}$ instead of the more standard $n^{1/2}$. Because of this, making inference about values that directly require the baseline distribution, such as creating a confindence interval for the median for subjects with a given set of covariates, cannot be done with the semi-parametric model. Secondly, we find later on that when it comes to cross-validation, we often found the semi-parametric estimator to be overly optimistic for some loss functions in comparison with a parametric model.

However, even if a parametric model is used for final inference, the semi-parametric model is still useful for assessing model fit. This is especially important for interval censored data, as we do not have the option of examining typical residuals or histograms as we would if the outcome was uncensored. **icenReg** has the function **diag_baseline** that plots several choices of parametric baseline distributions against the semi-parametric estimate. If the parametric distribution shows no systematic deviations from the semi-parametric fit, this implies the choice of parametric family may do a reason job of describing the underlying distribution. If there are clear deviations, this model should not be trusted.

To use diag_baseline, you must provide either a fitted model, or a



formula, data and model. You then select the parametric families that you would like plotted against the non-parametric estimate (default is to fit all available). As an example, suppose we wanted to examine the different parametric fits for the essIncData dataset. This could be done with

```
diag_baseline(cbind(inc_1, inc_u) ~
  eduLevel + cntry,
  model = "po",
  data = essIncData,
  lgdLocation = "topright")
```

Alternatively, using the fits from earlier, we can just call

```
diag_baseline(fit_po,lgdLocation = "topright")
```

Visual diagnostics are always subjective, but in this case we definitively know that the exponential fit is not appropriate and we believe the lognormal baseline is most appropriate for the proporitional odds model.

3.2 Examining Covariate Effect

Although semi-parametric models do not make assumptions about the parametric family of the baseline distribution, both fully-parametric and semi-parametric models make assumptions about the form of the covariate effect, akin to the link function in generalized linear models.

A rule of thumb for identifying gross violations of proportional hazards is to check if the Kaplan Meier curves cross; if they do, and this cross appears not purely by chance, the proportional hazards assumption seems inappropriate.

This can naturally extend to the case of interval censored data by replacing the Kaplan Meier curves with the NPMLE. Also, this informal test can be generalized to the proportional odds model; the proportional odds assumption also implies that survival curves that differ only by a constant factor of the odds of survival should not cross.

Another method of assessing involves transforming your survival estimates such that if the assumptions are met, the difference in transformed survival will be constant. For the proportional hazards model, this is the complementary log-log transformation (i.e. $\log(-\log(s))$). For the proportional odds model, this is the logit transformation (i.e. $\log(s/(1-s))$).

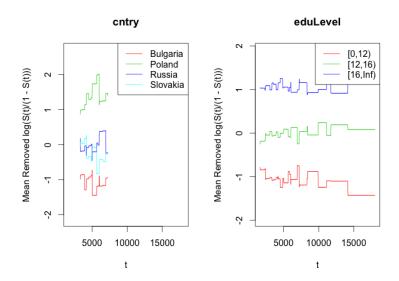
Plotting these functions can be done automatically in **icenReg** using the diag_covar function. The basic flow is that function takes in the fit, divides the data up on a covariate of interest. If it is categorical, it simply breaks up by category, if it is numeric, it attempts to find break point to evenly split up the data. Then, for each subset of the data, it fits the corresponding semi-parametric model and plots the transformation of the baseline distribution.

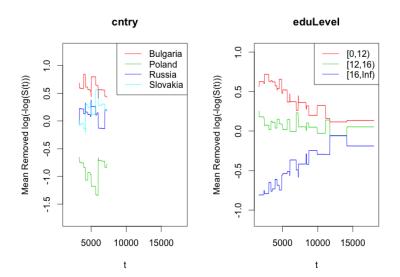
To demonstrate, suppose we wanted to assess whether the Cox-PH or proportional odds model was more appropriate for the essIncData. This could be done by

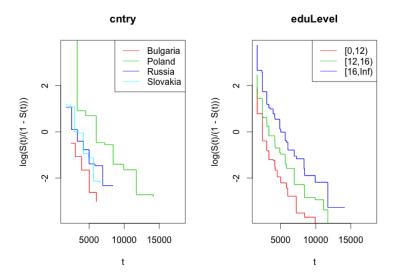
```
diag_covar(fit_po, lgdLocation = "topright")
diag_covar(fit_ph, lgdLocation = "topright")
```

We see that especially for eduLevel, the porportional odds seems much more appropriate (the difference between transformed values seems more constant). This agrees with the fact that the likelihood is almost 100 greater for the proportional odds model than Cox-PH.

Note that the plots for entry are very limited. This is because the mean trend is removed from the plots. However, since the cdf for the semi-







parametric model for the Bulgaria subset of the data is defined as exactly 1 around 10,000, the mean of the transformation is not defined.

We can replot the transformation, without the mean removed, by the following call:

```
diag_covar(fit_po, yType = "transform",
  lgdLocation = "topright")
```

3.3 Imputed Cross Validation

Utility Functions