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stcox PH-assumption tests — Tests of proportional-hazards assumption after stcox

Description Quick start Menu Syntax

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Acknowledgment References Also see

Description

stphplot plots $-\ln\{-\ln(survival)\}$ curves for each category of a nominal or ordinal covariate versus $\ln(analysis time)$ for right-censored data. These are often referred to as "log-log" plots. Optionally, these estimates can be adjusted for covariates. The proportional-hazards assumption is not violated when the curves are parallel.

stcoxkm plots Kaplan-Meier observed survival curves and compares them with the Cox predicted curves for the same variable. The closer the observed values are to the predicted, the less likely it is that the proportional-hazards assumption has been violated.

estat phtest tests the proportional-hazards assumption on the basis of Schoenfeld residuals after fitting a model with stcox.

Quick start

Log-log plot of survival

Check for parallel lines in plot of $-\ln\{-\ln(\text{survival})\}$ versus $\ln(\text{analysis time})$ for each category of covariate a using stset data

stphplot, by(a)

Same as above, but adjust for average values of covariates x1 and x2 stphplot, by(a) adjustfor(x1 x2)

Same as above

stphplot, by(a) adjustfor(x1 x2, atomeans)

Adjust for x1 = 0 and x2 = 0

stphplot, by(a) adjustfor(x1 x2, atzeros)

Kaplan-Meier and predicted survival plot

Compare Kaplan-Meier survival curve with predicted survival from Cox model for each category of covariate a using stset data

```
stcoxkm, by(a)
```

Same as above, but create separate plots for each level of a stcoxkm, by(a) separate

Test using Schoenfeld residuals

Test the proportional-hazards assumption after $\texttt{stcox} \ \texttt{x1} \ \texttt{x2} \ \texttt{x3}$ estat phtest

Same as above, and report separate test for each covariate estat phtest, detail

Menu

stphplot

Statistics > Survival analysis > Regression models > Graphically assess PH assumption

stcoxkm

Statistics > Survival analysis > Regression models > Kaplan-Meier versus predicted survival

estat phtest

Statistics > Survival analysis > Regression models > Test PH assumption

Syntax

Check proportional-hazards assumption:

```
Log-log plot of survival
```

```
stphplot [if], \{by(varname) | \underline{str}ata(varname) \} [stphplot\_options]
```

Kaplan-Meier and predicted survival plot

```
stcoxkm [if], by(varname) [stcoxkm_options]
```

Using Schoenfeld residuals

```
estat phtest [ , phtest_options ]
```

stphplot_options

Description

```
Main
```

* by (varname) fit separate Cox models; the default

* <u>str</u>ata(*varname*) fit stratified Cox model; requires adjustfor() adjustfor(*varlist*[, *suboptions*]) adjust the estimates to specific values of *varlist*;

default is overall means

Options

nonegative plot $ln{-ln(survival)}$

nolntime plot curves against analysis time do not show st setting information

Plot

plot#opts(stphplot_plot_options) affect rendition of the #th connected line and #th plotted points

Add plots

addplot (plot) add other plots to the generated graph

Y axis, X axis, Titles, Legend, Overall

twoway_options any options other than by() documented in [G-3] twoway_options

^{*}Either by(varname) or strata(varname) is required with stphplot.

stphplot_plot_options	Description
cline_options	change look of lines or connecting method
marker_options	change look of markers (color, size, etc.)
stcoxkm_options	Description
- Jain	
by(varname)	report the nominal or ordinal covariate
<u>tie</u> s(<u>bre</u> slow)	use Breslow method to handle tied failures
<u>tie</u> s(<u>efr</u> on)	use Efron method to handle tied failures
<u>tie</u> s(exactm)	use exact marginal-likelihood method to handle tied failures
<u>tie</u> s(exactp)	use exact partial-likelihood method to handle tied failures
<u>sep</u> arate	draw separate plot for predicted and observed curves
<u>nosh</u> ow	do not show st setting information
bserved plot	
obsopts(stcoxkm_plot_options)	affect rendition of the observed curve
<u>obs#</u> opts(stcoxkm_plot_options)	affect rendition of the #th observed curve; not allowed with separate
redicted plot	
<pre>predopts(stcoxkm_plot_options)</pre>	affect rendition of the predicted curve
<pre>pred#opts(stcoxkm_plot_options)</pre>	affect rendition of the #th predicted curve; not allowed with separate
dd plots	
addplot(plot)	add other plots to the generated graph
axis, X axis, Titles, Legend, Overall	
twoway_options	any options other than by() documented in [G-3] <i>twoway_options</i>
byopts(byopts)	how subgraphs are combined, labeled, etc.

stcoxkm_plot_options	Description
connect_options	change look of connecting method
marker_options	change look of markers (color, size, etc.)

You must stset your data before using stphplot and stcoxkm; see [ST] stset. fweights, iweights, and pweights may be specified using stset; see [ST] stset.

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phtest_options	Description
Main	
log	use natural logarithm time-scaling function
km	use 1 - KM product-limit estimate as the time-scaling function
rank	use rank of analysis time as the time-scaling function
time(varname)	use <i>varname</i> containing a monotone transformation of analysis time as the time-scaling function
plot(varname)	plot smoothed, scaled Schoenfeld residuals versus time
<u>bw</u> idth(#)	use bandwidth of #; default is bwidth(0.8)
$\underline{\mathtt{d}}\mathtt{etail}$	test proportional-hazards assumption separately for each covariate
Scatterplot	
marker_options	change look of markers (color, size, etc.)
marker_label_options	add marker labels; change look or position
Smoothed line	
<pre>lineopts(cline_options)</pre>	affect rendition of the smoothed line
Y axis, X axis, Titles, Legend, Ove	rall
twoway_options	any options other than by() documented in [G-3] <i>twoway_options</i>

estat phtest is not appropriate with svy estimation results.

collect is allowed with estat phtest; see [U] 11.1.10 Prefix commands.

Options

Options are presented under the following headings:

Options for stephplot Options for steoxkm Options for estat phtest

Options for stphplot

Main

by (varname) specifies the nominal or ordinal covariate. Either by() or strata() is required with stphplot.

strata(*varname*) is an alternative to by(). Rather than fitting separate Cox models for each value of *varname*, strata() fits one stratified Cox model. You must also specify adjustfor() with the strata() option; see [ST] sts graph.

adjustfor(varlist , suboptions) adjusts the estimates of the survivor function to specific values of varlist. The default is to adjust to overall mean values of covariates. adjustfor() can be specified with by(); it is required with strata().

suboptions are atomeans (the default), atmeans, atzeros, atbase, and at(); see [ST] adjust-for_option.

```
Options
```

nonegative specifies that $ln\{-ln(survival)\}\$ be plotted instead of $-ln\{-ln(survival)\}\$.

nolntime specifies that curves be plotted against analysis time instead of against ln(analysis time).

noshow prevents stphplot from showing the key st variables. This option is seldom used because most people type stset, show or stset, noshow to set whether they want to see these variables mentioned at the top of the output of every st command; see [ST] stset.

Plot

plot#opts(stphplot_plot_options) affects the rendition of the #th connected line and #th plotted points; see [G-3] cline_options and [G-3] marker_options.

Add plots

addplot(plot) provides a way to add other plots to the generated graph; see [G-3] addplot_option.

∫ Y axis, X axis, Titles, Legend, Overall Ì

twoway_options are any of the options documented in [G-3] twoway_options, excluding by(). These include options for titling the graph (see [G-3] title_options) and for saving the graph to disk (see [G-3] saving_option).

Options for stcoxkm

Main

by (varname) specifies the nominal or ordinal covariate. by () is required.

ties(breslow|efron|exactm|exactp) specifies one of the methods available to stcox for handling tied failures. If none is specified, ties(breslow) is assumed; see [ST] stcox.

separate produces separate plots of predicted and observed values for each value of the variable specified with by().

noshow prevents stcoxkm from showing the key st variables. This option is seldom used because most people type stset, show or stset, noshow to set whether they want to see these variables mentioned at the top of the output of every st command; see [ST] stset.

Observed plot

obsopts (stcoxkm_plot_options) affects the rendition of the observed curve; see [G-3] connect_options and [G-3] marker_options.

obs#opts(stcoxkm_plot_options) affects the rendition of the #th observed curve; see [G-3] connect_options and [G-3] marker_options. This option is not allowed with separate.

Predicted plot

predopts(stcoxkm_connect_options) affects the rendition of the predicted curve; see [G-3] connect_options and [G-3] marker_options.

pred#opts(stcoxkm_connect_options) affects the rendition of the #th predicted curve; see [G-3] connect_options and [G-3] marker_options. This option is not allowed with separate.

Add plots

addplot(plot) provides a way to add other plots to the generated graph; see [G-3] addplot_option.

Y axis, X axis, Titles, Legend, Overall

twoway_options are any of the options documented in [G-3] twoway_options, excluding by(). These include options for titling the graph (see [G-3] title_options) and for saving the graph to disk (see [G-3] saving_option).

byopts (*byopts*) affects the appearance of the combined graph when by () and separate are specified, including the overall graph title and the organization of subgraphs. See [G-3] *by_option*.

Options for estat phtest

Main

log, km, rank, and time() are used to specify the time scaling function.

By default, estat phtest performs the tests using the identity function, that is, analysis time itself.

log specifies that the natural log of analysis time be used.

km specifies that 1 minus the Kaplan-Meier product-limit estimate be used.

rank specifies that the rank of analysis time be used.

time(varname) specifies a variable containing an arbitrary monotonic transformation of analysis time. You must ensure that varname is a monotonic transform.

plot(varname) specifies that a scatterplot and smoothed plot of scaled Schoenfeld residuals versus time be produced for the covariate specified by varname. By default, the smoothing is performed using the running-mean method implemented in lowess, mean noweight; see [R] lowess.

bwidth(#) specifies the bandwidth. Centered subsets of bwidth() $\times N$ observations are used for calculating smoothed values for each point in the data except for endpoints, where smaller, uncentered subsets are used. The greater the bwidth(), the greater the smoothing. The default is bwidth(0.8).

detail specifies that a separate test of the proportional-hazards assumption be produced for each covariate in the Cox model. By default, estat phtest produces only the global test.

Scatterplot

marker_options affect the rendition of markers drawn at the plotted points, including their shape, size, color, and outline; see [G-3] marker_options.

marker_label_options specify if and how the markers are to be labeled; see [G-3] marker_label_options.

Smoothed line

lineopts(cline_options) affects the rendition of the smoothed line; see [G-3] cline_options.

Y axis, X axis, Titles, Legend, Overall

twoway_options are any of the options documented in [G-3] twoway_options, excluding by(). These include options for titling the graph (see [G-3] title_options) and for saving the graph to disk (see [G-3] saving_option).

Remarks and examples

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Cox proportional hazards models assume that the hazard ratio is constant over time. Suppose that a group of cancer patients on an experimental treatment is monitored for 10 years. If the hazard of dying for the nontreated group is twice the rate as that of the treated group (HR = 2.0), the proportional-hazards assumption implies that this ratio is the same at 1 year, at 2 years, or at any point on the time scale. Because the Cox model, by definition, is constrained to follow this assumption, it is important to evaluate its validity. If the assumption fails, alternative modeling choices would be more appropriate (for example, a stratified Cox model, time-varying covariates). For examples of testing the proportional-hazards assumption using Stata, see Allison (2014).

stphplot and stcoxkm provide graphical methods for assessing violations of the proportionalhazards assumption. Although using graphs to assess the validity of the assumption is subjective, it can be a helpful tool.

stphplot plots $-\ln\{-\ln(\text{survival})\}$ curves for each category of a nominal or ordinal covariate versus ln(analysis time). These are often referred to as "log-log" plots. Optionally, these estimates can be adjusted for covariates. If the plotted lines are reasonably parallel, the proportional-hazards assumption has not been violated, and it would be appropriate to base the estimate for that variable on one baseline survivor function.

Another graphical method of evaluating the proportional-hazards assumption, though less common, is to plot the Kaplan-Meier observed survival curves and compare them with the Cox predicted curves for the same variable. This plot is produced with stcoxkm. When the predicted and observed curves are close together, the proportional-hazards assumption has not been violated. See Garrett (1997) for more details.

Many popular tests for proportional hazards are, in fact, tests of nonzero slope in a generalized linear regression of the scaled Schoenfeld residuals on time (see Grambsch and Therneau [1994]). The estat phtest command tests, for individual covariates and globally, the null hypothesis of zero slope, which is equivalent to testing that the log hazard-ratio function is constant over time. Thus rejection of the null hypothesis of a zero slope indicates deviation from the proportional-hazards assumption. The estat phtest command allows three common time-scaling options (log, km, and rank) and also allows you to specify a user-defined function of time through the time() option. When no option is specified, the tests are performed using analysis time without further transformation.

Example 1

These examples use data from a leukemia remission study (Garrett 1997). The data consist of 42 patients who are monitored over time to see how long (weeks) it takes them to go out of remission (relapse: 1 = yes, 0 = no). Half the patients receive a new experimental drug, and the other half receive a standard drug (treatment1: 1 = drug A, 0 = standard). White blood cell count, a strong indicator of the presence of leukemia, is divided into three categories (wbc3cat: 1 = normal, 2 = moderate, 3 = high).

```
. use https://www.stata-press.com/data/r18/leukemia
(Leukemia remission study)
```

. describe

Contains data from https://www.stata-press.com/data/r18/leukemia.dta Observations: 42 Leukemia remission study Variables: 8 23 Mar 2022 10:39

Variable name	Storage type	Display format	Value label	Variable label
weeks	byte	%8.0g		Weeks in remission
relapse	byte	%8.0g	yesno	Relapse
treatment1	byte	%8.0g	trt1lbl	Treatment I
treatment2	byte	%8.0g	trt21b1	Treatment II
wbc3cat	byte	%9.0g	wbclbl	White blood cell count
wbc1	byte	%8.0g		wbc3cat==Normal
wbc2	byte	%8.0g		wbc3cat==Moderate
wbc3	byte	%8.0g		wbc3cat==High

Sorted by: weeks

. stset weeks, failure(relapse)

Survival-time data settings

Failure event: relapse!=0 & relapse<.

Observed time interval: (0, weeks] Exit on or before: failure

	total observations exclusions	
40	abaamwatiana mamainina	manmaganting

42 observations remaining, representing

30 failures in single-record/single-failure data

541 total analysis time at risk and under observation

At risk from t = 0 Earliest observed entry t = 0 Last observed exit t = 35

In this example, we examine whether the proportional-hazards assumption holds for drug A versus the standard drug (treatment1). First, we will use stphplot, followed by stcoxkm.

. stphplot, by(treatment1)
 Failure _d: relapse
Analysis time _t: weeks

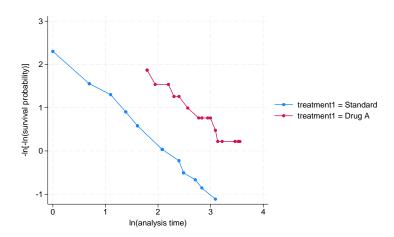


Figure 1.

. stcoxkm, by(treatment1) legend(pos(6) cols(2))

Failure _d: relapse Analysis time _t: weeks

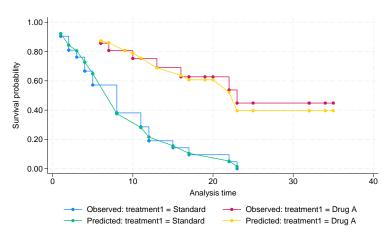


Figure 2.

Figure 1 (stphplot) displays lines that are parallel, implying that the proportional-hazards assumption for treatment1 has not been violated. This is confirmed in figure 2 (stcoxkm), where the observed values and predicted values are close together.

The graph in figure 3 is the same as the one in figure 1, adjusted for white blood cell count. By default, this adjustment sets each level of wbc3cat to its overall mean. In other words, the results are adjusted based on the observed proportions of individuals having normal, moderate, and high white blood cell counts.

·

stphplot, strata(treatment1) adjustfor(i.wbc3cat)

Failure _d: relapse Analysis time _t: weeks

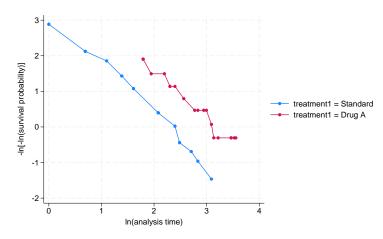


Figure 3.

The lines in figure 3 are still parallel, although they are somewhat closer together. Examining the proportional-hazards assumption on a variable without adjusting for covariates is usually adequate as a diagnostic tool before using the Cox model. However, if you know that adjustment for covariates in a final model is necessary, you may wish to reexamine whether the proportional-hazards assumption still holds.

If we wanted to adjust to the base level of the factor variable wbc3cat instead of the level-specific averages, we could have typed

. stphplot, strata(treatment1) adjustfor(i.wbc3cat, atbase)

Adjusting to a different value, however, would not affect our conclusion about the curves being parallel.

Another variable in this dataset measures a different drug (treatment2: 1 = drug B, 0 = standard). We wish to examine the proportional-hazards assumption for this variable.

. stphplot, by(treatment2) Failure _d: relapse Analysis time _t: weeks

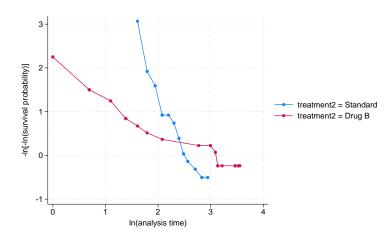


Figure 4.

. stcoxkm, by(treatment2) separate byopts(style(altleg)) Failure _d: relapse

Analysis time _t: weeks

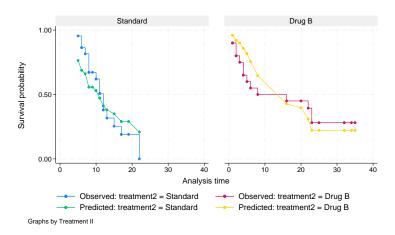


Figure 5.

This variable violates the proportional-hazards assumption. In figure 4, we see that the lines are not only nonparallel but also cross in the data region. In figure 5, we see that there are considerable differences between the observed and predicted values. We have overestimated the positive effect of drug B for the first half of the study and have underestimated it in the later weeks. One hazard ratio describing the effect of this drug would be inappropriate. We definitely would want to stratify on this variable in our Cox model.

Example 2: estat phtest

In this example, we use estat phtest to examine whether the proportional-hazards assumption holds for a model with covariates treatment1 and wbc3cat. After stsetting the data, we first run stcox with these factor variables as regressors. Then we use estat phtest:

. stset weeks, failure(relapse)

Survival-time data settings

Failure event: relapse!=0 & relapse<.

Observed time interval: (0, weeks]
Exit on or before: failure

42 total observations 0 exclusions

42 observations remaining, representing

30 failures in single-record/single-failure data

541 total analysis time at risk and under observation

At risk from t = 0
Earliest observed entry t = 0
Last observed exit t = 35

. stcox i.treatment1 i.wbc3cat, nolog

Failure _d: relapse Analysis time _t: weeks

Cox regression with Breslow method for ties

No. of subjects = 42 No. of failures = 30 Number of obs = 42

Time at risk = 541

Log likelihood = -77.476905

LR chi2(3) = 33.02Prob > chi2 = 0.0000

t	Haz. ratio	Std. err.	z	P> z	[95% conf.	interval]
treatment1 Drug A	. 2834551	.1229874	-2.91	0.004	.1211042	.6634517
wbc3cat Moderate High	3.637825 10.92214	2.201306 7.088783	2.13 3.68	0.033	1.111134 3.06093	11.91015 38.97284

. estat phtest, detail

Test of proportional-hazards assumption

Time function: Analysis time

	rho	chi2	df	Prob>chi2
Ob.treatme~1			1	
1.treatment1	-0.07019	0.15	1	0.6948
1b.wbc3cat			1	
2.wbc3cat	-0.03223	0.03	1	0.8650
3.wbc3cat	0.01682	0.01	1	0.9237
Global test		0.33	3	0.9551

Because we specified the detail option with the estat phtest command, both covariate-specific and global tests were produced. In addition, the rho column reports the correlation between the scaled Schoenfeld residuals and the specified function of time. We can see that there is no evidence that the proportional-hazards assumption has been violated.

Another variable in this dataset measures a different drug (treatment2: 1 = drug B, 0 = standard). We now wish to examine the proportional-hazards assumption for the previous model by substituting treatment2 for treatment1.

We fit a new Cox model and perform the test for proportional hazards:

. stcox i.treatment2 i.wbc3cat, nolog

Failure _d: relapse Analysis time _t: weeks

Cox regression with Breslow method for ties

No. of subjects = 42No. of failures = 30 Time at risk

Number of obs = 42

= 23.93 LR chi2(3) Prob > chi2 = 0.0000

Log likelihood = -82.019053

t	Haz. ratio	Std. err.	z	P> z	[95% conf.	interval]
treatment2 Drug B	.8483777	.3469054	-0.40	0.688	.3806529	1.890816
wbc3cat Moderate High	3.409628 14.0562	2.050784 8.873693	2.04 4.19	0.041	1.048905 4.078529	11.08353 48.44314

. estat phtest, detail

Test of proportional-hazards assumption

Time function: Analysis time

	rho	chi2	df	Prob>chi2
0b.treatme~2			1	
1.treatment2	-0.51672	10.19	1	0.0014
1b.wbc3cat			1	•
2.wbc3cat	-0.09860	0.29	1	0.5903
3.wbc3cat	-0.03559	0.04	1	0.8448
Global test		10.24	3	0.0166

treatment2 violates the proportional-hazards assumption. A single hazard ratio describing the effect of this drug is inappropriate.

The test of the proportional-hazards assumption is based on the principle that, for a given regressor, the assumption restricts $\beta(t_j) = \beta$ for all t_j . This implies that a plot of $\beta(t_j)$ versus time will have a slope of zero. Grambsch and Therneau (1994) showed that $E(s_i^*) + \widehat{\beta} \approx \beta(t_j)$, where s_i^* is the scaled Schoenfeld residual at failure time t_j and $\widehat{\beta}$ is the estimated coefficient from the Cox model. Thus a plot of $s_i^* + \widehat{\beta}$ versus some function of time provides a graphical assessment of the assumption.

Continuing from above, if you type

. predict sch*, scaledsch

you obtain five variables—sch1, sch2, sch3, sch4, and sch5—corresponding to the regressors. Ignoring the base categories, sch2 corresponds to 1.treatment2, sch4 corresponds to 2.wbc3cat, and sch5 corresponds to 3.wbc3cat. Given the utility of $s_i^* + \widehat{\beta}$, what is stored in variable sch2 is actually $s_{j2}^*+\widehat{eta}_2$ and not just the scaled Schoenfeld residual for the 1.treatment2, s_{j2}^* , itself. The estimated coefficient, $\hat{\beta}_2$, is added automatically. The same holds true for the variable representing the next regressor, $sch4 = s_{j4}^* + \hat{\beta}_4$, and so on.

As such, a graphical assessment of the proportional-hazards assumption for the first regressor is as simple as

```
. scatter sch2 _t || lfit sch2 _t
```

which plots a scatter of $s_{j2}^* + \widehat{\beta}_2$ versus analysis time, $_{-}$ t, and overlays a linear fit. Is the slope zero? The answer is no for 1.treatment2, and that agrees with our results from estat phtest.

4

□ Technical note

The tests of the proportional-hazards assumption assume homogeneity of variance across risk sets. This allows the use of the estimated overall (pooled) variance—covariance matrix in the equations. Although these tests have been shown by Grambsch and Therneau (1994) to be fairly robust to departures from this assumption, exercise care where this assumption may not hold, particularly when performing a stratified Cox analysis. In such cases, we recommend that you check the proportional-hazards assumption separately for each stratum.

Video example

How to fit a Cox proportional hazards model and check proportional-hazards assumption

Stored results

estat phtest stores the following in r():

```
Scalars r(df) global test degrees of freedom r(chi2) global test \chi^2 r(p) global test p-value Matrices r(phtest) separate tests for each covariate
```

Methods and formulas

For one covariate, x, the Cox proportional hazards model reduces to

$$h(t;x) = h_0(t) \exp(x\beta)$$

where $h_0(t)$ is the baseline hazard function from the Cox model. Let $S_0(t)$ and $H_0(t)$ be the corresponding Cox baseline survivor and baseline cumulative hazard functions, respectively.

The proportional-hazards assumption implies that

$$H(t) = H_0(t) \exp(x\beta)$$

or

$$\ln H(t) = \ln H_0(t) + x\beta$$

where H(t) is the cumulative hazard function. Thus, under the proportional-hazards assumption, the logs of the cumulative hazard functions at each level of the covariate have equal slope. This is the basis for the method implemented in stphplot.

The proportional-hazards assumption also implies that

$$S(t) = S_0(t) \exp(x\beta)$$

Let $\widehat{S}(t)$ be the estimated survivor function based on the Cox model. This function is a step function like the Kaplan-Meier estimate and, in fact, reduces to the Kaplan-Meier estimate when x=0. Thus for each level of the covariate of interest, we can assess violations of the proportional-hazards assumption by comparing these survival estimates with estimates calculated independently of the model. See Kalbfleisch and Prentice (2002) or Hess (1995).

stcoxkm plots Kaplan-Meier estimated curves for each level of the covariate together with the Cox model predicted baseline survival curve. The closer the observed values are to the predicted values, the less likely it is that the proportional-hazards assumption has been violated.

Grambsch and Therneau (1994) presented a scaled adjustment for the Schoenfeld residuals that permits the interpretation of the smoothed residuals as a nonparametric estimate of the log hazard-ratio function. These scaled Schoenfeld residuals, $\mathbf{r}_{S_i}^*$, can be obtained directly with predict's scaledsch option; see [ST] stcox postestimation.

Scaled Schoenfeld residuals are centered at $\widehat{\beta}$ for each covariate and, when there is no violation of proportional hazards, should have slope zero when plotted against functions of time. The estat phtest command uses these residuals, tests the null hypothesis that the slope is equal to zero for each covariate in the model, and performs the global test proposed by Grambsch and Therneau (1994). The test of zero slope is equivalent to testing that the log hazard-ratio function is constant over time. With the detail option, estat phtest also reports the correlation between the scaled Schoenfeld residuals and the specified function of time.

For a specified function of time, q(t), the statistic for testing the pth individual covariate is, for $\overline{g}(t) = d^{-1} \sum_{i=1}^{N} \delta_i g(t_i),$

$$\chi_c^2 = \frac{\left[\sum_{i=1}^N \left\{\delta_i g(t_i) - \overline{g}(t)\right\} r_{S_{pi}}^*\right]^2}{d \operatorname{Var}(\widehat{\beta}_p) \sum_{i=1}^N \left\{\delta_i g(t_i) - \overline{g}(t)\right\}^2}$$

which is asymptotically distributed as χ^2 with 1 degree of freedom. $r_{S_{ni}}^*$ is the scaled Schoenfeld residual for observation i, and δ_i indicates failure for observation i, with $d = \sum \delta_i$.

The statistic for the global test is calculated as

$$\chi_g^2 = \left[\sum_{i=1}^N \left\{ \delta_i g(t_i) - \overline{g}(t) \right\} \mathbf{r}_{S_i} \right]' \left[\frac{d \operatorname{Var}(\widehat{\boldsymbol{\beta}})}{\sum_{i=1}^N \left\{ \delta_i g(t_i) - \overline{g}(t) \right\}^2} \right] \left[\sum_{i=1}^N \left\{ \delta_i g(t_i) - \overline{g}(t) \right\} \mathbf{r}_{S_i} \right]$$

for \mathbf{r}_{S_i} , a vector of the m (unscaled) Schoenfeld residuals for the ith observation; see [ST] **stcox postestimation**. The global test statistic is asymptotically distributed as χ^2 with m degrees of freedom.

The equations for the scaled Schoenfeld residuals and the two test statistics just described assume homogeneity of variance across risk sets. Although these tests are fairly robust to deviations from this assumption, care must be exercised, particularly when dealing with a stratified Cox model.

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Also see

[ST] **stcox** — Cox proportional hazards model

[ST] sts — Generate, graph, list, and test the survivor and related functions

[ST] stset — Declare data to be survival-time data

[ST] adjustfor_option — Adjust survivor and related functions for covariates at specific values

[U] 20 Estimation and postestimation commands

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