Survival Package Functions

Terry Therneau

December 17, 2010

Contents

1 Introduction

Let us change or traditional attitude to the construction of programs. Instead of imagining that our main task is to instruct a *computer* what to do, let us concentrate rather on explaining to *humans* what we want the computer to do. (Donald E. Knuth, 1984).

This is the definition of a coding style called *literate programming*. I first made use of it in the *coxme* library and have become a full convert. For the survival library only selected objects are documented in this way; whenever I find need for a major revision I "convert" the source. An underlying motivation is to leave code that is well enough explained that someone else can take it over.

2 residuals.survreg

The residuals for a survreg model are one of several types

response residual y value on the scale of the original data

deviance an approximate deviance residual. A very bad idea statistically, retained for the sake of backwards compatability.

dfbeta a matrix with one row per observation and one column per parameter showing the approximate influence of each observation on the final parameter value

dfbetas the dfbeta residuals scaled by the standard error of each coefficient

working residuals on the scale of the linear predictor

ldcase likelihood displacement wrt case weights

ldresp likelihood displacement wrt response changes

ldshape likelihood displacement wrt changes in shape

matrix matrix of derivatives of the log-likelihood wrt paramters

The other parameters are

rsigma whether the scale parameters should be included in the result for dfbeta results. I can think of no reason why one would not want them.

collapse optional vector of subject identifiers. This is for the case where a subject has multiple observations in a data set, and one wants to have residuals per subject rather than residuals per observation.

weighted whether the residuals should be multiplied by the case weights. The sum of weighted residuals will be zero.

The routine starts with standard stuff, checking arguments for validity and etc. The two cases of response or working residuals require a lot less computation. and are the most common calls, so they are taken care of first.

```
\langle residuals.survreg \rangle \equiv
  # $Id$
  # Residuals for survreg objects
 residuals.survreg <- function(object, type=c('response', 'deviance',</pre>
                           'dfbeta', 'dfbetas', 'working', 'ldcase',
                           'ldresp', 'ldshape', 'matrix'),
                           rsigma =TRUE, collapse=FALSE, weighted=FALSE, ...) {
      type <-match.arg(type)</pre>
      n <- length(object$linear.predictors)</pre>
      Terms <- object$terms</pre>
      if(!inherits(Terms, "terms"))
                stop("invalid terms component of object")
      # If there was a cluster directive in the model statment then remove
      # it. It does not correspond to a coefficient, and would just confuse
      # things later in the code.
      cluster <- untangle.specials(Terms, "cluster")$terms</pre>
      if (length(cluster) >0 )
           Terms <- Terms[-cluster]</pre>
      strata <- attr(Terms, 'specials')$strata</pre>
      coef <- object$coefficients</pre>
      intercept <- attr(Terms, "intercept")</pre>
      response <- attr(Terms, "response")</pre>
      weights <- object$weights</pre>
      if (is.null(weights)) weighted <- FALSE
      \langle rsr-dist \rangle
      \langle rsr-data \rangle
```

```
\langle rsr\text{-}resid \rangle
\langle rsr\text{-}finish \rangle
```

First retrieve the distribution, which is used multiple times. The common case is a character string pointing to some element of survreg.distributions, but the other is a user supplied list of the form contained there. Some distributions are defined as the transform of another in which case we need to set itrans and dtrans and follow the link, otherwise the transformation and its inverse are the identity.

```
\(\( \text{crr-dist} \) \\
\( \text{if (is.character(object$dist))} \\
\( \text{dd <- survreg.distributions[[object$dist]]} \)
\( \text{else dd <- object$dist} \)
\( \text{if (is.null(dd$itrans)) { \\
\( \text{itrans <- dtrans <-function(x)x } \)
\( \text{} \)
\( \text{else { \\
\( \text{itrans <- dd$itrans } \)
\( \text{dtrans <- dd$dtrans } \)
\( \text{if (!is.null(dd$dist)) dd <- survreg.distributions[[dd$dist]] deviance <- dd$deviance \( \text{deviance dens <- dd$density} \)
\( \text{definition} \)
\( \text{deviance distributions[[dd$dist]] deviance <- dd$density} \)
\( \text{definition} \)
\( \text{deviance distributions[[dd$dist]] deviance <- dd$density} \)
\( \text{definition} \)
\( \text{deviance dens <- dd$density} \)
\( \text{definition} \)
\( \text{deviance distributions[[dd$dist]] deviance } \)
\( \text{definition} \)
\( \text{deviance dens <- dd$density} \)
\( \text{definition} \)
\( \te
```

The next task is to decide what data we need. The response is always needed, but is normally saved as a part of the model. If it is a transformed distribution such as the Weibull (a transform of the extreme value) the saved object y is the transformed data, so we need to replicate that part of the survreg() code. (Why did I even allow for y=F in survreg? Because I was mimicing the lm function — oh the long, long consequences of a design decision.)

The covariate matrix x will be needed for all but response, deviance, and working residuals. If the model included a strata() term then there will be multiple scales, and the strata variable needs to be recovered. The variable sigma is set to a scalar if there are no strata, but otherwise to a vector with n elements containing the appropriate scale for each subject.

The leverage type residuals all need the second derivative matrix. If there was a cluster statement in the model this will be found in naive.var, otherwise in the var component.

```
tranfun <- dd$trans
        exactsurv <- y[,ncol(y)] ==1</pre>
        if (any(exactsurv)) logcorrect <-sum(log(dd$dtrans(y[exactsurv,1])))</pre>
        if (type=='interval') {
             if (any(y[,3]==3))
                      y <- cbind(tranfun(y[,1:2]), y[,3])</pre>
             else y <- cbind(tranfun(y[,1]), y[,3])</pre>
        else if (type=='left')
              y \leftarrow cbind(tranfun(y[,1]), 2-y[,2])
                  y \leftarrow cbind(tranfun(y[,1]), y[,2])
        else
        }
    else {
        if (type=='left') y[,2] \leftarrow 2- y[,2]
        else if (type=='interval' && all(y[,3]<3)) y <- y[,c(1,3)]
        }
    }
if (!is.null(strata)) {
    temp <- untangle.specials(Terms, 'strata', 1)</pre>
    Terms2 <- Terms[-temp$terms]</pre>
    if (length(temp$vars)==1) strata.keep <- mf[[temp$vars]]</pre>
    else strata.keep <- strata(mf[,temp$vars], shortlabel=TRUE)</pre>
    strata <- as.numeric(strata.keep)</pre>
    nstrata <- max(strata)</pre>
    sigma <- object$scale[strata]</pre>
else {
    Terms2 <- Terms
    nstrata <- 1
    sigma <- object$scale
    }
if (need.x) {
   x <- object[['x']] #don't grab xlevels component
   if (is.null(x))
        x <- model.matrix(Terms2, mf, contrasts.arg=object$contrasts)</pre>
```

The most common residual is type response, which requires almost no more work, for the others we need to create the matrix of derivatives before proceeding. We use the **center** component from the deviance function for the distribution, which returns the data point y itself for an exact, left, or right censored observation, and an appropriate midpoint for interval censored ones.

```
\langle rsr\text{-}resid \rangle \equiv
```

```
if (type=='response') {
    yhat0 <- deviance(y, sigma, object$parms)
    rr <- itrans(yhat0$center) - itrans(object$linear.predictor)
    }
else {
    ⟨rtr-deriv⟩
    ⟨rtr-resid2⟩
    }</pre>
```

The matrix of derivatives is used in all of the other cases. The starting point is the density function of the distribution which return a matrix with columns of F(x), 1-F(x), f(x), f'(x)/f(x) and f''(x)/f(x). The matrix type residual contains columns for each of

$$L_i \quad \frac{\partial L_i}{\partial \eta_i} \quad \frac{\partial^2 L_i}{\partial \eta_i^2} \quad \frac{\partial L_i}{\partial \log(\sigma)} \quad \frac{\partial L_i}{\partial \log(\sigma)^2} \quad \frac{\partial^2 L_i}{\partial \eta \partial \log(\sigma)}$$

where L_i is the contribution to the log-likelihood from each individual. Note that if there are multiple scales, i.e. a strata() term in the model, then terms 3–6 are the derivatives for that subject with respect to their *particular* scale factor; derivatives with respect to all the other scales are zero for that subject.

The log-likelihood can be written as

$$L = \sum_{exact} [\log(f(z_i)) - \log(\sigma_i)] + \sum_{censored} \log \left(\int_{z_i^l}^{z_i^u} f(u) du \right)$$

$$\equiv \sum_{exact} [g_1(z_i) - \log(\sigma_i)] + \sum_{censored} \log(g_2(z_i^l, z_i^u))$$

$$z_i = (y_i - \eta_i) / \sigma_i$$

For the interval censored observations we have a z defined at both the lower and upper endpoints. The linear predictor is $\eta = X\beta$.

The derivatives are shown below. Note that $f(-\infty) = f(\infty) = F(-\infty) = 0$, $F(\infty) = 1$, $z^u = \infty$ for a right censored observation and $z^l = -\infty$ for a left censored one.

$$\begin{split} \frac{\partial g_1}{\partial \eta} &= -\frac{1}{\sigma} \left[\frac{f'(z)}{f(z)} \right] \\ \frac{\partial g_2}{\partial \eta} &= -\frac{1}{\sigma} \left[\frac{f(z^u) - f(z^l)}{F(z^u) - F(z^l)} \right] \\ \frac{\partial^2 g_1}{\partial \eta^2} &= \frac{1}{\sigma^2} \left[\frac{f''(z)}{f(z)} \right] - (\partial g_1 / \partial \eta)^2 \\ \frac{\partial^2 g_2}{\partial \eta^2} &= \frac{1}{\sigma^2} \left[\frac{f'(z^u) - f'(z^l)}{F(z^u) - F(z^l)} \right] - (\partial g_2 / \partial \eta)^2 \\ \frac{\partial g_1}{\partial \log \sigma} &= -\left[\frac{zf'(z)}{f(z)} \right] \\ \frac{\partial g_2}{\partial \log \sigma} &= -\left[\frac{z^u f(z^u) - z^l f(z^l)}{F(z^u) - F(z^l)} \right] \end{split}$$

$$\frac{\partial^2 g_1}{\partial (\log \sigma)^2} = \left[\frac{z^2 f''(z) + z f'(z)}{f(z)} \right] - (\partial g_1 / \partial \log \sigma)^2
\frac{\partial^2 g_2}{\partial (\log \sigma)^2} = \left[\frac{(z^u)^2 f'(z^u) - (z^l)^2 f'(z_l)}{F(z^u) - F(z^l)} \right] - \partial g_1 / \partial \log \sigma (1 + \partial g_1 / \partial \log \sigma)
\frac{\partial^2 g_1}{\partial \eta \partial \log \sigma} = \frac{z f''(z)}{\sigma f(z)} - \partial g_1 / \partial \eta (1 + \partial g_1 / \partial \log \sigma)
\frac{\partial^2 g_2}{\partial \eta \partial \log \sigma} = \frac{z^u f'(z^u) - z^l f'(z^l)}{\sigma [F(z^u) - F(z^l)]} - \partial g_2 / \partial \eta (1 + \partial g_2 / \partial \log \sigma)$$

In the code z is the relevant point for exact, left, or right censored data, and z2 the upper endpoint for an interval censored one. The variable tdenom contains the denominator for each subject (which is the same for all derivatives for that subject). For an interval censored observation we try to avoid numeric cancellation by using the appropriate tail of the distribution. For instance with $(z^l, z^u) = (12, 15)$ the value of F(x) will be very near 1 and it is better to subtract two upper tail values (1 - F) than two lower tail ones F.

```
\langle rtr\text{-}deriv \rangle \equiv
  status <- y[,ncol(y)]
  eta <- object$linear.predictors
 z \leftarrow (y[,1] - eta)/sigma
  dmat <- dens(z, object$parms)</pre>
  dtemp < -dmat[,3] * dmat[,4]
                                    #f,
  if (any(status==3)) {
      z2 \leftarrow (y[,2] - eta)/sigma
      dmat2 <- dens(z2, object$parms)</pre>
      }
  else {
      dmat2 <- dmat
                       #dummy values
      z2 <- 0
      }
  tdenom <- ((status==0) * dmat[,2]) + #right censored
             ((status==1) * 1 )
                                      + #exact
             ((status==2) * dmat[,1]) + #left
             ((status==3) * ifelse(z>0, dmat[,2]-dmat2[,2],
                                          dmat2[,1] - dmat[,1])) #interval
  g <- log(ifelse(status==1, dmat[,3]/sigma, tdenom)) #loglik
  tdenom <- 1/tdenom
  dg \leftarrow -(tdenom/sigma) *(((status==0) * (0-dmat[,3])) +
                                                                 #dg/ eta
                            ((status==1) * dmat[,4]) +
                            ((status==2) * dmat[,3]) +
                            ((status==3) * (dmat2[,3]- dmat[,3])))
  ddg <- (tdenom/sigma^2) *(((status==0) * (0- dtemp)) + #ddg/eta^2
                               ((status==1) * dmat[,5]) +
```

Now, we can calcultate the actual residuals case by case. For the dfbetas there will be one column per coefficient, so if there are strata column 4 of the deriv matrix needs to be *uncollapsed* into a matrix with nstrata columns. The same manipulation is needed for the ld residuals.

```
\langle rtr\text{-}resid2\rangle \equiv
  if (type=='deviance') {
      yhat0 <- deviance(y, sigma, object$parms)</pre>
      rr <- (-1)*deriv[,2]/deriv[,3] #working residuals
      rr <- sign(rr)* sqrt(2*(yhat0$loglik - deriv[,1]))</pre>
  else if (type=='working') rr <- (-1)*deriv[,2]/deriv[,3]
  else if (type=='dfbeta' || type== 'dfbetas' || type=='ldcase') {
      score <- deriv[,2] * x # score residuals</pre>
      if (rsigma) {
          if (nstrata > 1) {
               d4 <- matrix(0., nrow=n, ncol=nstrata)</pre>
               d4[cbind(1:n, strata)] <- deriv[,4]
               score <- cbind(score, d4)</pre>
               }
          else score <- cbind(score, deriv[,4])</pre>
          }
      rr <- score %*% vv
      if (type=='dfbetas') rr <- rr %*% diag(1/sqrt(diag(vv)))</pre>
      if (type=='ldcase') rr<- rowSums(rr*score)</pre>
  else if (type=='ldresp') {
      rscore <- deriv[,3] * (x * sigma)
      if (rsigma) {
          if (nstrata >1) {
```

```
d6 <- matrix(0., nrow=n, ncol=nstrata)</pre>
               d6[cbind(1:n, strata)] <- deriv[,6]*sigma
               rscore <- cbind(rscore, d6)</pre>
           else rscore <- cbind(rscore, deriv[,6] * sigma)</pre>
      temp <- rscore %*% vv
      rr <- rowSums(rscore * temp)</pre>
  else if (type=='ldshape') {
      sscore <- deriv[,6] *x</pre>
      if (rsigma) {
           if (nstrata >1) {
               d5 <- matrix(0., nrow=n, ncol=nstrata)</pre>
               d5[cbind(1:n, strata)] <- deriv[,5]
               sscore <- cbind(sscore, d5)</pre>
           else sscore <- cbind(sscore, deriv[,5])</pre>
      temp <- sscore %*% vv
      rr <- rowSums(sscore * temp)</pre>
  else { #type = matrix
      rr <- deriv
   Finally the two optional steps of adding case weights and collapsing over subject id.
\langle rsr\text{-}finish \rangle \equiv
  #case weights
  if (weighted) rr <- rr * weights
  #Expand out the missing values in the result
  if (!is.null(object$na.action)) {
      rr <- naresid(object$na.action, rr)</pre>
      if (is.matrix(rr)) n <- nrow(rr)</pre>
      else
                            n <- length(rr)</pre>
      }
  # Collapse if desired
  if (!missing(collapse)) {
      if (length(collapse) !=n) stop("Wrong length for 'collapse'")
      rr <- drop(rowsum(rr, collapse))</pre>
      }
```

3 Predicted survival from a Cox model

3.1 Individual survival

The survfit method for a Cox model produces individual survival curves. As might be expected these have much in common with ordinary survival curves, and share many of the same methods. The primary differences are first that a predicted curve always refers to a particular set of covariate values. It is often the case that a user wants multiple values at once, in which case the result will be a matrix of survival curves with a row for each time and a column for each covariate set. The second is that the computations are somewhat more difficult.

The input arguments are

formula a fitted object of class 'coxph'. The argument name of 'formula' is historic, from when the survfit function was not a generic and only did Kaplan-Meier type curves.

newdata contains the data values for which curves should be produced, one per row

se.fit TRUE/FALSE, should standard errors be computed.

individual a particular option for time-dependent covariates

type computation type for the survival curve

vartype computation type for the variance

censor if FALSE, remove any times that have no events. This is for backwards compatability with older versions of the code.

id replacement and extension for the individual argument

All the other arguments are common to all the methods, refer to the help pages.

```
 \begin{array}{c} \text{id, y2, strata2)} \\ \langle \mathit{survfit.coxph-finish} \rangle \\ \end{array} \}
```

The third line as.name('survfit') causes the printout to say 'survfit' instead of 'surv-fit.coxph'.

The setup for the routine is fairly pedestrian. If the newdata argument is missing we use object\$means as the default value. This choice has lots of statistical shortcomings, particularly in a stratified model, but is common in other packages and a historic option here. If the type or vartype are missing we use the appropriate one for the method in the Cox model. That is, the coxph computation used for method=''exact'' is the same approximation used in the Kalbfleish-Prentice estimate, that for the Breslow method matches the Aalen survival estimate, and the Efron approximation the Efron survival estimate. The other two rows of labels in temp1 are historical; we include them for backwards compatability but they don't appear in the documentation.

```
\langle survfit.coxph\text{-}setup \rangle \equiv
  if (missing(type)) {
      # Use the appropriate one from the model
      temp1 <- c("exact", "breslow", "efron")</pre>
      survtype <- match(object$method, temp1)</pre>
  else {
      temp1 <- c("kalbfleisch-prentice", "aalen", "efron",</pre>
                   "kaplan-meier", "breslow", "fleming-harrington",
                   "greenwood", "tsiatis", "exact")
      survtype <- match(match.arg(type, temp1), temp1)</pre>
      survtype \leftarrow c(1,2,3,1,2,3,1,2,3)[survtype]
  if (missing(vartype)) {
      vartype <- survtype
      }
  else {
      temp2 <- c("greenwood", "aalen", "efron", "tsiatis")</pre>
      vartype <- match(match.arg(vartype, temp2), temp2)</pre>
      if (vartype==4) vartype<- 2
      }
  if (!se.fit) conf.type <- "none"
 else conf.type <- match.arg(conf.type)</pre>
```

I need to retrieve a copy of the original data. We always need the X matrix and y, both of which may be found in the data object. If the original call included either strata, offset, or weights, or if either x or y are missing from the coxph object, then model.frame is needed. We have to use object['x'] instead of object\$x since the latter will pick off the xlevels component if the x component is missing (which is the default).

```
\langle survfit.coxph-setup \rangle + \equiv
```

```
if (is.null(object$y) || is.null(object[['x']]) ||
  !is.null(object$call$weights) ||
  !is.null(attr(object$terms, 'specials')$strata) ||
  !is.null(attr(object$terms, 'offset'))) {
    mf <- model.frame(object)
    }
else mf <- NULL #useful for if statements later</pre>
```

If a model frame was created, then it is trivial to grab y from the new frame and compare it to object\$y from the original one. This is to avoid nonsense results that arise when someone changes the data set under our feet. For instance

```
fit <- coxph(Surv(time,status) ~ age, data=lung)</pre>
  lung <- lung[1:100,]</pre>
  survfit(fit)
\langle survfit.coxph-setup \rangle + \equiv
  if (is.null(mf)) y <- object[['y']]</pre>
  else {
      y <- model.response(mf)</pre>
      y2 <- object[['y']]</pre>
      if (!is.null(y2) && any(as.matrix(y2) != as.matrix(y)))
           stop("Could not reconstruct the y vector")
      }
  if (is.null(object[['x']])) x <- model.matrix.coxph(object, mf=mf)</pre>
  else x <- object[['x']]</pre>
 n \leftarrow nrow(y)
  if (n != object$n[1] || nrow(x) !=n)
      stop("Failed to reconstruct the original data set")
  if (is.null(mf)) wt \leftarrow rep(1., n)
  else {
      wt <- model.weights(mf)</pre>
      if (is.null(wt)) wt \leftarrow rep(1.0, n)
  type <- attr(y, 'type')</pre>
  if (type != 'right' && type != 'counting')
      stop("Cannot handle \"", type, "\" type survival data")
 missid <- missing(id) # I need this later, and setting id below makes
                           # "missing(id)" alwasy false
  if (!missid) individual <- TRUE
  else if (missid && individual) id <- rep(0,n)
  else id <- NULL
```

```
if (individual && type!= 'counting')
    stop("The individual option is only valid for start-stop data")

if (is.null(mf)) offset <- 0
else {
    offset <- model.offset(mf)
    if (is.null(offset)) offset <- 0
    }

Terms <- object$terms
temp <- untangle.specials(Terms, 'strata')
if (length(temp$terms)==0) strata <- rep(OL,n)
else {
    if (length(temp$vars) ==1) strata <- mf[[temp$vars]]
    else strata <- strata(m[, temp$vars], shortlabel=TRUE)
}</pre>
```

A key variable for the computation is the risk score $\exp(X\beta)$ for each original observation along with the risk score for the target subject(s). There are three choices for the new data

- For predictions with time-dependent covariates the user will have specified the option individual=TRUE, and the new data set contains the covariate trace over time of a single individual; or if id is present we have the covariate trace for multiple subjejects. In either case we need to retrieve the covariates, strata, and repsonse from the new data set.
- For ordinary predictions only the covariates are needed.
- If newdata is not present we assume that this is the ordinary case, and use the value of object\$means as the default covariate set. This is not a good idea statistically since many users view this as an "average" survival curve, which it is not.

Since the new data might not have all of the levels present for one of the factors, we need to ensure that levels match the original data. In all cases the computation depends only on the difference between the original covariates and the target set. We subtract the means from each column of the X matrices to avoid any problems with overflow in the exponential function. For a model with a frailty term that was treated as sparse, those coefficients are in the frailty component and do not appear in the means. Since the X variables in this case are all 0/1 no centering is necessary for accuracy, we use 0 as a handy value.

If a variable is deemed redundant the coxph routine will have set its coefficient to NA as a marker. We want to ignore that coefficient: treating it as a zero has the desired effect. Another special case is a null model, having either 1 or only an offset on the right hand side. In that case we create a dummy covariate to allow the rest of the code to work without special if/else. The last special case is a model with a sparse frailty term. We treat the frailty coefficients as 0 variance (in essence as an offset). The frailty is removed from the model variables but kept in the risk score. This isn't statistically very defensible, but it is backwards compatable. A non-sparse frailty is not treated in a special way.

```
\langle survfit.coxph-setup \rangle + \equiv
```

```
if (is.null(x) \mid | ncol(x)==0) { # a model with ~1 on the right hand side
    # Give it a dummy x so the rest of the code goes through
    # (This case is really rare)
    x <- matrix(0., nrow=n)
    coef <- 0.0
    varmat <- matrix(0.0,1,1)</pre>
    risk <- rep(exp(offset- mean(offset)), length=n)
    }
else {
    varmat <- object$var</pre>
    coef <- ifelse(is.na(object$coefficients), 0, object$coefficients)</pre>
    xcenter <- object$means
    if (is.null(object$frail)) {
        x <- scale(x, center=xcenter, scale=FALSE)
        risk <- c(exp(x%*% coef + offset - mean(offset)))
   else {
       x <- x[,!is.na(match(dimnames(x)[[2]], names(coef))), drop=F]
       risk <- exp(object$linear.predictor)</pre>
       x <- scale(x, center=xcenter, scale=FALSE)
    }
```

When grabbing [newdata] we want to keep the model.frame processing, in particular the handling of missing values. For backwards compatability, I allow someone to give an ordinary vector instead of a data frame, when only one curve is required. In this case I also need to verify that the elements have a name. If not we attach the variable names from the model, and assume that it's the right order. Then turn it into a list, like it should have been from the beginning. (Documentation of this ability has been suppressed, however.)

Once we have newdata in the correct form, pull data from it. If individual is false then the repsonse variable and any strata from the original model are not needed. Any cluster terms can be deleted in either case. Last, there is no ability to use sparse frailties and newdata together.

```
temp <- untangle.specials(Terms, 'cluster')
if (length(temp$vars)) Terms2 <- Terms[-temp$terms]
else Terms2 <- Terms
if (!individual) {
   Terms2 <- delete.response(Terms2)
   temp <- untangle.specials(Terms2, 'strata')
   if (length(temp$vars)) Terms2 <- Terms2[-temp$terms]
   }</pre>
```

Now get the newdata model frame using 6 lines of mysterious code. First get an abbreviated form of the original call that has only the calling function, newdata, and id. The calling function is always element 1, the others are found by name. Now manipulate it: change the name of "newdata" to "data", add the formula and xlev components (the second might be NULL), and then change the name. If the original call was survfit(fit1, newdata=mydat, conf.int=.9) the result is model.frame(data=mydat, formula=Terms2, xlev=object\$xlev)

```
\langle survfit.coxph-setup \rangle + \equiv
      tcall <- Call[c(1, match(c('newdata', 'id'), names(Call), nomatch=0))]</pre>
      names(tcall)[2] <- 'data' #rename newdata to data
      tcall$formula <- Terms2
      tcall$xlev <- object$xlevels
      tcall[[1]] <- as.name('model.frame')</pre>
      mf2 <- eval(tcall)
      }
   Extract the data items for individual survival.
\langle survfit.coxph-setup \rangle + \equiv
  if (individual) {
      if (missing(newdata))
          stop("The newdata argument must be present when individual=TRUE")
      if (!missid)) { #grab the id variable
          id <- model.extract("id")</pre>
          if (is.null(id)) stop("id=NULL is an invalid argument")
      temp <- untangle.specials(Terms2, 'strata')</pre>
      if (length(temp$vars) >0) {
          strata2 <- strata(mf2[temp$vars], shortlabel=TRUE)</pre>
          strata2 <- factor(strata2, levels=levels(strata))</pre>
          if (any(is.na(strata2)))
               stop("New data set has strata levels not found in the original")
          Terms2 <- Terms2[-temp$terms]</pre>
      else strata2 <- rep(0, nrow(mf2))</pre>
      x2 <- model.matrix(Terms2, mf2)[,-1, drop=FALSE] #no intercept
      if (length(x2)==0) stop("Individual survival but no variables")
      x2 <- scale(x2, center=xcenter, scale=FALSE)</pre>
```

```
offset2 <- model.offset(mf2)
    if (length(offset2) >0) offset2 <- offset2 - mean(offset)</pre>
    else offset2 <- 0
    y2 <- model.extract(mf2, 'response')
    if (attr(y2,'type') != type)
        stop("Survival type of newdata does not match the fitted model")
    if (attr(y2, "type") != "counting")
        stop("Individual=TRUE is only valid for counting process data")
    y2 <- y2[,1:2] #throw away status, it's never used
else {
    y2 <- strata2 <- NULL #dummy arguments
    if (missing(newdata)) {
        x2 <- matrix(0.0, nrow=1, ncol=ncol(x))</pre>
        offset2 <- 0
        }
    else {
        x2 <- model.matrix(Terms2, mf2)[,-1, drop=FALSE] #no intercept
        x2 <- scale(x2, center=xcenter, scale=FALSE)</pre>
        offset2 <- model.offset(mf2)</pre>
        if (length(offset2) >0) offset2 <- offset2 - mean(offset)</pre>
        else offset2 <- 0
        }
    }
newrisk <- exp(c(x2 %*% coef) + offset2)</pre>
```

Now, we're ready to do the main computation. Before this revision (the one documented here using noweb) there were three C routines used in calculating survival after a Cox model

- 1. agsurv1 creates a single curve, but for the most general case of a *covariate path*. It is used for time dependent covariates.
- 2. agsurv2 creates a set of curves. These curves are for a fixed covariate set, although (start, stop] data is supported. If there were 3 strata in the fit and 4 covariate sets are given, the result will be 12 curves.
- 3. agsurv3 is used to create population survival curves. The result is average survival curve (for 3 different definitions of 'average'). If there were 3 strata and 100 subjects, the first curve returned would be the average for those 100 individual curves in strata 1, the second for strata 2, and the third for strata 3.

In June 2010 the first two were re-written in (mostly) R, in the process of adding functionality and repairing some flaws in the computation of a weighted variance. In effect, the changes are similar to the rewrite of the survfitKM function a few years ago.

Computations are separate for each strata, and each strata will have a different number of time points in the result. Thus we can't preallocate a matrix. Instead we generate an empty list,

one per strata, and then populate it with the survival curves. At the end we unlist the individual components one by one. This is memory efficient, the number of curves is usually small enough that the "for" loop is no great cost, and it's easier to see what's going on than C code.

First, compute the baseline survival curves for each strata. If the strata was a factor we want to leave it in the same order, otherwise sort it. This fitting routine was set out as a separate function for the sake of the rms package. They want to utilize the computation, but have a different setup for recreation of the x and y data.

```
\langle survfit.coxph.fit \rangle \equiv
  survfit.coxph.fit <- function(y, x, wt, x2, risk, newrisk, strata, se.fit,</pre>
                                      survtype, vartype, varmat, id, y2, strata2) {
       if (is.factor(strata)) ustrata <- levels(strata)</pre>
                                  ustrata <- sort(unique(strata))</pre>
       nstrata <- length(ustrata)</pre>
       survlist <- vector('list', nstrata)</pre>
       for (i in 1:nstrata) {
           indx <- which(strata== ustrata[i])</pre>
           survlist[[i]] <- agsurv(y[indx,,drop=F], x[indx,,drop=F],</pre>
                                        wt[indx], risk[indx],
                                        survtype, vartype)
           }
       \langle survfit.coxph-compute \rangle
       if (se.fit) result$std.err <- sqrt(varh)
       }
```

In an ordinary survival curve object with multiple strata, as produced by survfitKM, the time, survival and etc components are each a single vector that contains the results for strata 1, followed by strata 2, The strata compontent is a vector of integers, one per strata, that gives the number of elements belonging to each stratum. The reason is that each strata will have a different number of observations, so that a matrix form was not viable, and the underlying C routines were not capable of handling lists (the code predates the .Call function by a decade). Although a modern redesign would leave the survlist object above as a set of lists we retain the older form.

For individual=FALSE we have a second dimension, namely each of the target covariate sets (if there are multiples). Each of these generates a unique set of survival and variance(survival) values, but all of the same size since each uses all the strata. The final output structure in this case has single vectors for the time, number of events, number censored, and number at risk values since they are common to all the curves, and a marix of survival and variance estimates, one column for each of the distinct target values. If Λ_0 is the baseline cumulative hazard from the above calculation, then $r_i\Lambda_0$ is the cumulative hazard for the ith new risk score r_i . The variance has two parts, the first of which is $r_i^2H_1$ where H_1 is returned from the agsurv routine, and the second is

$$H_2(t) = d'(t)Vd(t)$$

$$d(t) = \int_0^t [z - \overline{x}(s)] d\Lambda(s)$$

V is the variance matrix for β from the fitted Cox model, and d(t) is the distance between the target covariate z and the mean of the original data, summed up over the interval from 0 to t. Essentially the variance in $\hat{\beta}$ has a larger influence when prediction is far from the mean.

```
\langle survfit.coxph-compute \rangle \equiv
  if (is.null(id)) {
      cumhaz <- unlist(lapply(survlist, function(x) x$cumhaz))</pre>
      varhaz <- unlist(lapply(survlist, function(x) cumsum(x$varhaz)))</pre>
      nevent <- unlist(lapply(survlist, function(x) x$n.event)) #weighted
      ndeath <- unlist(lapply(survlist, function(x) x$ndeath)) #unweighted</pre>
              <- t(matrix(unlist(lapply(survlist, function(x) t(x$xbar))),
                         nrow=ncol(x)))
      hazard <- unlist(lapply(survlist, function(x) x$hazard))</pre>
      if (survtype==1)
          surv <-unlist(lapply(survlist, function(x) cumprod(x$surv)))</pre>
      else surv <- exp(-cumhaz)</pre>
      if (is.matrix(x2) && nrow(x2) >1) { #more than 1 row in newdata
          surv <- outer(surv, newrisk, '^')</pre>
          varh <- matrix(0., nrow=length(varhaz), ncol=nrow(x2))</pre>
          for (i in 1:nrow(x2)) {
               dt <- outer(cumhaz, x2[i,], '*') - xbar</pre>
               varh[,i] <- (varhaz + rowSums((dt %*% varmat)* dt)) *</pre>
                   newrisk[i]^2
          }
      else {
          surv <- surv^newrisk</pre>
          dt <- outer(cumhaz, c(x2)) - xbar
          varh <- (varhaz + rowSums((dt %*% varmat)* dt)) *</pre>
               newrisk^2
          }
```

In the lines just above: I have a matrix dt with one row per death time and one column per variable. For each row d_i separately we want the quadratic form $d_iVd'_i$. The first matrix product can be done for all rows at once: found in the inner parenthesis. Ordinary (not matrix) multiplication followed by rowsums does the rest in one fell swoop.

```
\( \survfit.coxph-compute \rangle +=
\text{result <- list(n=as.vector(table(strata)),}
\time=unlist(lapply(survlist, function(x) x\$time)),
\text{n.risk= unlist(lapply(survlist, function(x) x\$n.risk)),}
\text{n.event=nevent,}
\text{n.censor=unlist(lapply(survlist, function(x) x\$n.censor)),}
\]</pre>
```

```
surv=surv)
```

```
if (nstrata >1) {
    result$strata <- unlist(lapply(survlist, function(x) length(x$n.risk)))
    names(result$strata) <- ustrata
    }
}</pre>
```

For the case with id not missing, we create a single survival curve for each unique id (subject). A subject will spend blocks of time with different covariate sets, sometimes even jumping between strata. Retrieve each one and save it into a list, and then sew them together end to end. The n component is the number of observations in the strata — but this subject might visit several. We report the first one they were in for printout. The time component will be cumulative on this subject's scale. Counting this is a bit trickier than I first thought. Say that the subject's first interval goes from 1 to 10, with observed time points in that interval at 2, 5, and 7, and a second interval from 12 to 20 with observed time points in the data of 15 and 18. On the subject's time scale things happen at days 1, 4, 6, 12 and 15. The deltas saved below are 2-1, 5-2, 7-5, 3+ 14-12, 17-14. Note the 3+ part, kept in the timeforward variable. Why all this "adding up" nuisance? If the subject spent time in two strata, the second one might be on an internal time scale of 'time since entering the strata'. The two intervals in newdata could be 0-10 followed by 0-20. Time for the subject can't go backwards though: the change between internal/external time scales is a bit like following someone who was stepping back and forth over the international date line.

In the code the indx variable points to the set of times that the subject was present, for this row of the new data. Note the > on one end and \le on the other. If someone's interval 1 was 0–10 and interval 2 was 10–20, and there happened to be a jump in the baseline survival curve at exactly time 10 (someone else died), that jump is counted only in the first interval.

```
\langle survfit.coxph-compute \rangle + \equiv
  else {
      onecurve <- function(slist, x2, y2, strata2, ustrata) {
          ntarget <- nrow(x2) #number of different time intervals
          surv <- vector('list', ntarget)</pre>
          n.event <- n.risk <- n.censor <- varh1 <- varh2 <- time <- surv
          stemp <- match(strata2, ustrata)</pre>
          timeforward <- 0
          for (i in 1:ntarget) {
               slist <- survlist[[stemp[i]]]</pre>
               indx <- which(slist$time > y2[i,1] & slist$time <= y2[i,2])</pre>
               if (length(indx)==0) {
                   timeforward <- timeforward + y2[i,2] - y2[i,1]</pre>
                   next # No deaths or censors in user's interval. Possible
                          # user error, but not uncommon at the tail of the curve.
                   }
               time[[i]] <- diff(c(y2[i,1], slist$time[indx])) #time increments</pre>
               time[[i]][1] \leftarrow time[[i]][1] + timeforward
```

```
timeforward <- y2[i,2] - max(slist$time[indx])</pre>
        if (survtype==1) surv[[i]] <- slist$surv[indx]^newrisk[i]</pre>
                          surv[[i]] <- slist$hazard[indx]*newrisk[i]</pre>
        n.event[[i]] <- slist$n.event[indx]</pre>
        n.risk[[i]] <- slist$n.risk[indx]</pre>
        n.censor[[i]]<- slist$n.censor[indx]</pre>
        dt <- outer(slist$cumhaz[indx], x2[i,]) - slist$xbar[indx,,drop=F]</pre>
        varh1[[i]] <- slist$varhaz[indx] *newrisk[i]^2</pre>
        varh2[[i]] <- rowSums((dt %*% varmat)* dt) * newrisk[i]^2</pre>
    varh <- cumsum(unlist(varh1)) + unlist(varh2)</pre>
    if (survtype==1) surv <- cumprod(unlist(surv)) #increments (K-M)
    else surv <- exp(-cumsum(unlist(surv)))</pre>
                                                       #hazards
    list(n=as.vector(table(strata)[stemp[1]]),
               time=cumsum(unlist(time)),
                n.risk = unlist(n.risk),
                n.event= unlist(n.event),
               n.censor= unlist(n.censor),
                surv = surv,
                varh=varh)
    }
if (all(id ==id[1])) {
    result <- onecurve(survlist, x2, y2, strata2, ustrata)
    varh <- result$varh</pre>
    result$varh <-NULL
    }
else {
    onelist <- vector('list', length=length(unique(id)))</pre>
    for (i in unique(id)) {
        indx <- which(id==i)</pre>
        onelist[[i]] <- onecurve(survlist, x2[indx,], y2[indx],</pre>
                                   strata2[indx], ustrata)
    result <- list(n=unlist(lapply(onelist, function(x) x$n)),
                    time=unlist(lapply(onelist, function(x) x$time)),
                    n.risk= unlist(lapply(onelist, function(x) x$n.risk)),
                    n.event=unlist(lapply(onelist, function(x) x$nevent)),
                    n.censor=unlist(lapply(onelist, function(x) x$n.censor)),
                    surv= unlist(lapply(onelist, function(x) x$surv)))
    result$strata <- unlist(lapply(onelist, function(x) length(x$n.risk)))
    names(result$strata) <- unique(id)</pre>
```

```
result$ varh <- unlist(lapply(onelise, function(x) x$varh))
}</pre>
```

Next is the code for the agsurv function, which actually does the work. The estimates of survival are the Kalbfleisch-Prentice (KP), Breslow, and Efron. Each has an increment at each unique death time. First a bit of notation: $Y_i(t)$ is 1 if bservation i is "at risk" at time t and 0 otherwise. For a simple survival (ncol(y)==2) a subject is at risk until the time of censoring or death (first column of y). For (start, stop) data (ncol(y)==3) a subject becomes a part of the risk set at start+0 and stays through stop. $dN_i(t)$ will be 1 if subject i had an event at time t. The risk score for each subject is $r_i = \exp(X_i\beta)$.

The Breslow increment at time t is $\sum w_i dN_i(t) / \sum w_i r_i Y_i(t)$, the number of events at time t over the number at risk at time t. The final survival is exp(-cumsum(increment)).

The Kalbfleish-Prentice increment is a multiplicative term z which is the solution to the equation

$$\sum w_i r_i Y_i(t) = \sum dN_i(t) w_i \frac{r_i}{1 - z(t)^{r_i}}$$

The left hand side is the weighted number at risk at time t, the right hand side is a sum over the tied events at that time. If there is only one event the equation has a closed form solution. If not, and knowing the solution must lie between 0 and 1, we do 35 steps of bisection to get a solution within 1e-8. An alternative is to use the -log of the Breslow estimate as a starting estimate, which is faster but requires a more sophisticated iteration logic. The final curve is $\prod_t z(t)^{r_c}$ where r_c is the risk score for the target subject.

The Efron estimate can be viewed as a modified Breslow estimate under the assumption that tied deaths are not really tied – we just don't know the order. So if there are 3 subjects who die at some time t we will have three psuedo-terms for t, $t + \epsilon$, and $t + 2\epsilon$. All 3 subjects are present for the denominator of the first term, 2/3 of each for the second, and 1/3 for the third terms denominator. All contribute 1/3 of the weight to each numerator (1/3 chance they were the one to die there). The formulas will require $\sum w_i dN_i(t)$, $\sum w_i r_i dN_i(t)$, and $\sum w_i X_i dN_i(t)$, i.e., the sums only over the deaths.

For simple survival data the risk sum $\sum w_i r_i Y_i(t)$ for all the unique death times t is fast to compute as a cumulative sum, starting at the longest followup time an summing towards the shortest. There are two algorithms for (start, stop) data.

- Do a separate sum at each death time. The problem is for very large data sets. For each death time the selection who <- (start<t & stop>=t) is O(n) and can take more time then all the remaining calculations together.
- Use the difference of two cumulative sums, one ordered by start time and one ordered by stop time. This is O(2n) for the intial sums. The problem here is potential round off error if the sums get large, which can happen if the time scale were very, very finely divided. This issue is mostly precluded by subtracting means first.

We compute the extended number still at risk — all whose stop time is \geq each unique death time — in the vector xin. From this we have to subtract all those who haven't actually entered yet found in xout. Remember that (3,20] enters at time 3+. The total at risk at any time is the difference between them. Output is only for the stop times; a call to approx is used to reconcile

the two time sets. The **irisk** vector is for the printout, it is a sum of weighted counts rather than weighted risk scores.

```
\langle agsurv \rangle \equiv
  agsurv <- function(y, x, wt, risk, survtype, vartype) {</pre>
      nvar <- ncol(as.matrix(x))</pre>
      status <- y[,ncol(y)]
      dtime \leftarrow y[,ncol(y)-1]
      death <- (status==1)</pre>
      time <- sort(unique(dtime))</pre>
      nevent <- as.vector(rowsum(wt*death, dtime))</pre>
      ncens <- as.vector(rowsum(wt*(!death), dtime))</pre>
      wrisk <- wt*risk
      rcumsum <- function(x) rev(cumsum(rev(x))) # sum from last to first
      nrisk <- rcumsum(rowsum(wrisk, dtime))</pre>
      irisk <- rcumsum(rowsum(wt, dtime))</pre>
      if (ncol(y) == 2) {
           temp2 <- rowsum(wrisk*x, dtime)</pre>
           xsum
                  <- apply(temp2, 2, rcumsum)</pre>
           }
      else {
           delta <- min(diff(time))/2</pre>
           etime <- c(sort(unique(y[,1])), max(y[,1])+delta) #unique entry times
           indx <- approx(etime, 1:length(etime), time, method='constant',</pre>
                             rule=2, f=1)$y
           esum <- rcumsum(rowsum(wrisk, y[,1])) #not yet entered</pre>
           nrisk <- nrisk - c(esum,0)[indx]</pre>
           irisk <- irisk - c(rcumsum(rowsum(wt, y[,1])),0)[indx]</pre>
                  <- apply(rowsum(wrisk*x, y[,1]), 2, rcumsum) #not yet entered</pre>
           xin <- apply(rowsum(wrisk*x, dtime), 2, rcumsum) # dtime or alive
           xsum <- xin - (rbind(xout,0))[indx,,drop=F]</pre>
           }
      ndeath <- rowsum(status, dtime) #unweighted death count
   The KP estimate requires a short C routine to do the iteration efficiently, and the Efron
estimate a different C routine to efficiently compute the partial sums.
\langle agsurv \rangle + \equiv
      dtimes <- which(nevent >0)
      ntime <- length(time)</pre>
      if (survtype ==1) {
           indx <- (which(status==1))[order(dtime[status==1])] #deaths</pre>
           km <- .C('agsurv4',</pre>
                as.integer(ndeath),
                 as.double(risk[indx]),
                 as.double(wt[indx]),
```

```
as.integer(ntime),
         as.double(nrisk),
         inc = double(ntime))
    }
if (survtype==3 || vartype==3) {
    xsum2 <- rowsum((wrisk*death) *x, dtime)</pre>
    erisk <- rowsum(wrisk*death, dtime) #risk score sums at each death
    tsum <- .C('agsurv5',
                as.integer(length(nevent)),
                as.integer(nvar),
                 as.integer(ndeath),
                 as.double(nrisk),
                as.double(erisk),
                as.double(xsum),
                 as.double(xsum2),
                 sum1 = double(length(nevent)),
                 sum2 = double(length(nevent)),
                xbar = matrix(0., length(nevent), nvar))
haz <- switch(survtype,
                 nevent/nrisk,
                 nevent/nrisk,
                 nevent* tsum$sum1)
varhaz <- switch(vartype,</pre>
                 nevent/(nrisk *
                            ifelse(nevent>=nrisk, nrisk, nrisk-nevent)),
                 nevent/nrisk^2.
                 nevent* tsum$sum2)
xbar <- switch(vartype,</pre>
                (xsum/nrisk)*haz,
                (xsum/nrisk)*haz,
               nevent * tsum$xbar)
result <- list(time=time, n.event=nevent, n.risk=irisk, n.censor=ncens,
               hazard=haz,
               cumhaz=cumsum(haz), varhaz=varhaz, ndeath=ndeath,
               xbar=apply(matrix(xbar, ncol=nvar),2, cumsum))
if (survtype==1) result$surv <- km$inc
result
```

The arguments to this function are the number of unique times n, which is the length of the vectors ndeath (number at each time), denom, and the returned vector km. The risk and wt vectors contain individual values for the subjects with an event. Their length will be equal to sum(ndeath).

```
\langle agsurv4 \rangle \equiv
  #include "survS.h"
  #include "survproto.h"
  void agsurv4(Sint
                        *ndeath,
                                    double *risk,
                                                      double *wt,
                                     double *denom,
                                                       double *km)
                Sint
                        *sn,
  {
      int i, j, k, 1;
      int n; /* number of unique death times */
      double sumt, guess, inc;
      n = *sn;
      j =0;
      for (i=0; i<n; i++) {
           if (ndeath[i] ==0) km[i] =1;
           else if (ndeath[i] ==1) { /* not a tied death */
               km[i] = pow(1- wt[j]*risk[j]/denom[i], 1/risk[j]);
           else { /* biscection solution */
               guess = .5;
               inc = .25;
               for (1=0; 1<35; 1++) { /* bisect it to death */
                   sumt = 0;
                   for (k=j; k<(j+ndeath[i]); k++) {</pre>
                        sumt += wt[k]*risk[k]/(1-pow(guess, risk[k]));
               if (sumt < denom[i]) guess += inc;</pre>
                              guess -= inc;
               else
               inc = inc/2;
               km[i] = guess;
           j += ndeath[i];
      }
  }
   Do a computation which is slow in R, needed for the Efron approximation. Input arguments
n number of observations (unique death times)
d number of deaths at that time
nvar number of covariates
x1 weighted number at risk at the time
x2 sum of weights for the deaths
```

xsum matrix containing the cumulative sum of x values

xsum2 matrix of sums, only for the deaths

On output the values are

- d=0: the outputs are unchanged (they initialize at 0)
- \bullet d=1

```
sum1 1/x1
sum2 1/x1^2
xbar xsum/x1^2
```

 \bullet d=2

```
sum1 (1/2) ( 1/x1 + 1/(x1 - x2/2))
sum2 (1/2) ( same terms, squared)
xbar (1/2) (xsum/x1^2 + (xsum - 1/2 x3)/(x1- x2/2)^2)
```

 \bullet d=3

 \bullet etc

Sum1 will be the increment to the hazard, sum2 the increment to the first term of the variance, and xbar the increment in the hazard times the mean of x at this point.

```
\langle agsurv5 \rangle \equiv
 #include "survS.h"
                                              Sint *dd, double *x1,
 void agsurv5(Sint *n,
                               Sint *nvar,
                               double *xsum, double *xsum2,
                double *x2,
                double *sum1, double *sum2, double *xbar) {
      double temp;
      int i,j, k, kk;
      double d;
      for (i=0; i< *n; i++) {
          d = dd[i];
          if (d==1){
               temp = 1/x1[i];
               sum1[i] = temp;
               sum2[i] = temp*temp;
               for (k=0; k< *nvar; k++)
```

```
xbar[i+ *n*k] = xsum[i + *n*k] * temp*temp;
}
else {
   temp = 1/x1[i];
   for (j=0; j<d; j++) {
        temp = 1/(x1[i] - x2[i]*j/d);
        sum1[i] += temp/d;
        sum2[i] += temp*temp/d;
        for (k=0; k< *nvar; k++){
            kk = i + *n*k;
            xbar[kk] += ((xsum[kk] - xsum2[kk]*j/d) * temp*temp)/d;
        }
    }
}</pre>
```

Finally, the last (somewhat boring) part of the code. First, if given the argument censor=FALSE we need to remove all the time points from the output at which there was only censoring activity. This action is mostly for backwards compatability with older releases that never returned censoring times. Second, add in the variance and the confidence intervals to the result. The code is nearly identical to that in survfitKM.

```
\langle survfit.coxph\text{-}finish \rangle \equiv
  if (!censor) {
      kfun <- function(x, keep){ if (is.matrix(x)) x[keep,,drop=F]</pre>
                                   else if (length(x)==length(keep)) x[keep]
                                   else x}
      keep <- (result$n.event > 0)
      if (!is.null(result$strata)) {
          temp <- rep(names(result$strata), each=result$strata)</pre>
          result$strata <- c(table(temp[keep]))</pre>
      result <- lapply(result, kfun, keep)
  if (se.fit) {
      zval <- qnorm(1- (1-conf.int)/2, 0,1)</pre>
      if (conf.type=='plain') {
          temp1 <- result$surv + zval* result$std.err * result$surv</pre>
          temp2 <- result$surv - zval* result$std.err * result$surv</pre>
          result <- c(result, list(upper=pmin(temp1,1), lower=pmax(temp2,0),</pre>
                            conf.type='plain', conf.int=conf.int))
      if (conf.type=='log') {
          xx <- ifelse(result$surv==0,1,result$surv) #avoid some "log(0)" messages
          temp1 <- ifelse(result$surv==0, 0*result$std.err,</pre>
```

```
exp(log(xx) + zval* result$std.err))
        temp2 <- ifelse(result$surv==0, 0*result$std.err,</pre>
                         exp(log(xx) - zval* result$std.err))
        result <- c(result, list(upper=pmin(temp1,1), lower=temp2,
                         conf.type='log', conf.int=conf.int))
    if (conf.type=='log-log') {
        who <- (result$surv==0 | result$surv==1) #special cases</pre>
        xx <- ifelse(who, .1,result$surv) #avoid some "log(0)" messages
        temp1 <- exp(-exp(log(-log(xx)) + zval*result$std.err/log(xx)))</pre>
        temp1 <- ifelse(who, result$surv + 0*result$std.err, temp1)</pre>
        temp2 <- exp(-exp(log(-log(xx)) - zval*result$std.err/log(xx)))</pre>
        temp2 <- ifelse(who, result$surv + 0*result$std.err, temp2)</pre>
        result <- c(result, list(upper=temp1, lower=temp2,
                         conf.type='log-log', conf.int=conf.int))
result$call <- Call
if (is.R()) class(result) <- c('survfit.cox', 'survfit')</pre>
else
            oldClass(result) <- 'survfit.cox'</pre>
result
```

4 survexp

The arguments for survexp are

formula The model formula. The right hand side consists of grouping variables, identically to survfit and an optional ratetable directive. The "response" varies by method:

- for the Hakulinen method it is a vector of censoring times. This is the actual censoring time for censored subjecs, and is what the censoring time 'would have been' for each subject who died.
- for the conditional method it is the usual Surv(time, status) code
- for the Ederer method no response is given

data, weights, subset, na.action as usual

rmap an optional mapping for rate table variables, see more below.

times An optional vector of time points at which to compute the response. For the Hakulinen and conditional methods the program uses the vector of unique y values if this is missing. For the Ederer the component is not optional.

cohort Should the program produce an overall survival curve (cohort=T) or separate estimates for each subject.

conditional chooses between the Hakulinen and conditional methods.

ratetable the population rate table to use as a reference. This can either be a ratetable object or a previously fitted Cox model

scale Scale the resulting output times, e.g., 365.25 to turn days into years.

npoints This argument is rarely used. If times is not given then output at npoints evenly spaced points spanning the range of y.

se.fit Add standard errors to the output.

```
model, x, y usual
```

The output of survexp contains a subset of the elements in a survfit object, so many of the survfit methods can be applied. In R the result has a class of c('survexp', 'survfit'); in Splus this needs to be a global setting using the setOldClass function.

The first few lines are standard. Keep a copy of the call, then manufacture a call to model.frame that contains only the arguments relevant to that function.

The function works with two data sets, the user's data on an actual set of subjects and the reference ratetable. This leads to a particular nuisance, that the variable names in the data set may not match those in the ratetable. For instance the United States overall death rate table survexp.us expects 3 variables, as shown by summary(survexp.us)

• age = age in days for each subject at the start of follow-up

- sex = sex of the subject, "male" or "female" (the routine accepts any unique abbreviation and is case insensitive)
- year = date of the start of follow-up

Up until the most recent revision, the formula contained any necessary mapping between the variables in the data set and the ratetable. For instance

In this case the user's data set has a variable 'age' containing age in years, along with sex and an entry date. This had to be changed for two reasons. The primary one is that the data in a ratetable call had to be converted into a matrix in order to "pass through" the model.frame logic. With the recent updates to coxph so that it remembers factor codings correctly in new data sets, it is advantageous to keep factors as factors. The second is that a coxph model with a large number of covariates induces a very long ratetable clause; at about 40 variable it causes one of the R internal routines to fail due to a long expression. A third reason, perhaps the most pressing in reality, is that I've always felt that the prior code was confusing since it used the same term 'ratetable' for two different tasks.

The new process adds the rmap argument, an example would be rmap=list(age =age*365.25, year=entry.dt). Any variables in the ratetable that are not found in rmap are assumed to not need a mapping, this would be sex in the above example. For backwards compatability we allow the old style argument, converting it into the new style.

The rmap argument needs to be examined without evaluating it; we then add the appropriate extra variables into a temporary formula so that the model frame has all that is required. The ratetable variables then can be retrieved from the model frame.

```
\langle survexp\text{-}setup \rangle + \equiv
 rate <- attr(Terms, "specials")$ratetable</pre>
  if(length(rate) > 1)
           stop("Can have only 1 ratetable() call in a formula")
  if(length(rate) == 1) {
      if (!missing(rmap))
           stop("The ratetable() call in the formula is depreciated")
      stemp <- untangle.specials(Terms, 'ratetable')</pre>
      rcall <- as.call(parse(text=stemp$var)[[1]])</pre>
                                                           # as a call object
      rcall[[1]] <- as.name('list')</pre>
                                                           # make it a call to list(...
      Terms <- Terms[-stemp$terms]</pre>
                                                           # remove from the formula
      }
  else if (!missing(rmap)) {
      rcall <- substitute(rmap)</pre>
      if (!is.call(rcall) || rcall[[1]] != as.name('list'))
           stop ("Invalid rcall argument")
      }
```

```
else rcall <- NULL
# Check that there are no illegal names in rcall, then expand it
# to include all the names in the ratetable
if(is.ratetable(ratetable))
                              varlist <- attr(ratetable, "dimid")</pre>
else if(inherits(ratetable, "coxph")) {
    ## Remove "log" and such things, to get just the list of
        variable names
    varlist <- all.vars(delete.response(ratetable$terms))</pre>
    }
else stop("Invalid rate table")
temp <- match(names(rcall)[-1], varlist) # 2,3,... are the argument names
if (any(is.na(temp)))
    stop("Variable not found in the ratetable:", (names(rcall))[is.na(temp)])
if (any(!(varlist %in% names(rcall)))) {
    to.add <- varlist[!(varlist %in% names(rcall))]</pre>
    temp1 <- paste(text=paste(to.add, to.add, sep='='), collapse=',')</pre>
    if (is.null(rcall)) rcall <- parse(text=paste("list(", temp1, ")"))[[1]]</pre>
    else {
        temp2 <- deparse(rcall)</pre>
        rcall <- parse(text=paste("c(", temp2, ",list(", temp1, "))"))[[1]]
```

The formula below is used only in the call to model.frame to ensure that the frame has both the formula and the ratetable variables. We don't want to modify the original formula, since we use it to create the X matrix and the response variable. The non-obvious bit of code is the addition of an environment to the formula. The model.matrix routine has a non-standard evaluation - it uses the frame of the formula, rather than the parent.frame() argument below, along with the data to look up variables.

```
⟨survexp-setup⟩+≡

# Create a temporary formula, used only in the call to model.frame
newvar <- all.vars(rcall)
if (length(newvar) > 0) {
    tform <- paste(deparse(Terms), paste(newvar, collapse='+'), sep='+')
    if (is.R()) m$formula <- as.formula(tform, environment(Terms))
    else m$formula <- as.formula(tform)
    }

if (is.R()) m <- eval(m, parent.frame())
else m <- eval(m, sys.parent())
</pre>
```

If the user data has 0 rows, e.g. from a mistaken **subset** statement that eliminated all subjects, we need to stop early. Otherwise the .C code fails in a nasty way.

```
\langle survexp\text{-}setup \rangle + \equiv
n <- nrow(m)
```

}

```
if (n==0) stop("Data set has 0 rows")
weights <- model.extract(m, 'weights')
if (!is.null(weights)) warning("Weights ignored")

if (any(attr(Terms, 'order') >1))
            stop("Survexp cannot have interaction terms")
if (!missing(times)) {
    if (any(times<0)) stop("Invalid time point requested")
    if (length(times) >1 )
        if (any(diff(times)<0)) stop("Times must be in increasing order")
}</pre>
```

If a response variable was given, we only need the times and not the status. To be correct, computations need to be done for each of the times given in the times argument as well as for each of the unique y values. This ends up as the vector newtime. If a times argument was given we will subset down to only those values at the end.

The next step is to check out the ratetable. For a population rate table a set of consistency checks is done by the match.ratetable function, giving a set of sanitized indices R. For a Cox model R will be a model matix whose covariates are coded in exactly the same way that variables were coded in the original Cox model. We call the model.matrix.coxph function to avoid the steps found there (remove cluster statements, etc). We also need to use the mf argument of the function, otherwise it will call model.frame internally and fail when it can't find the response variable (which we don't need).

Note that for a population rate table the standard error of the expected is by definition 0 (the population rate table is based on a huge sample). For a Cox model rate table, an se formula is currently only available for the Ederer method.

```
\langle survexp-compute \rangle \square ovars <- attr(Terms, 'term.labels')
rdata <- data.frame(eval(rcall, m))  # the variables matching the ratetable
if (is.ratetable(ratetable)) {
   israte <- TRUE</pre>
```

```
if (no.Y) {
          if (missing(times))
              stop("There is no times argument, and no follow-up times are given in the formula")
          else newtime <- sort(unique(times))</pre>
          Y <- rep(max(times), n)
          }
      se.fit <- FALSE
      rtemp <- match.ratetable(rdata, ratetable)</pre>
      R <- rtemp$R
      }
  else if (inherits(ratetable, 'coxph')) {
      israte <- FALSE
      Terms <- ratetable$terms</pre>
      if (!is.null(attr(Terms, 'offset')))
          stop("Cannot deal with models that contain an offset")
      strats <- attr(Terms, "specials")$strata</pre>
      if (length(strats))
          stop("survexp cannot handle stratified Cox models")
      if (any(names(m[,rate]) != attr(ratetable$terms, 'term.labels')))
            stop("Unable to match new data to old formula")
      R <- model.matrix.coxph(ratetable, data=rdata)</pre>
      if (no.Y) {
          if (missing(se.fit)) se.fit <- TRUE</pre>
      else se.fit <- FALSE
  else stop("Invalid ratetable")
   Now for some calculation. If cohort is false, then any covariates on the right hand side (other
than the rate table) are irrelevant, the function returns a vector of expected values rather than
survival curves.
\langle survexp\text{-}compute \rangle + \equiv
  if (!cohort) { #individual survival
      if (no.Y) stop("For non-cohort, an observation time must be given")
      if (israte)
            temp <- survexp.fit (cbind(1:n,R), Y, max(Y), TRUE, ratetable)</pre>
      else temp<- survexp.cfit(cbind(1:n,R), Y, FALSE, TRUE, ratetable, FALSE)
      xx <- temp$surv
      names(xx) <- row.names(m)</pre>
      na.action <- attr(m, "na.action")</pre>
      if (length(na.action)) return(naresid(na.action, xx))
      else return(xx)
```

}

Now for the more commonly used case: returning a survival curve. First see if there are any grouping variables. The results of the tcut function are often used in person-years analysis, which is somewhat related to expected survival. However tcut results aren't relevant here and we put in a check for the confused user. The strata command creates a single factor incorporating all the variables.

```
\langle survexp\text{-}compute \rangle + \equiv
  if (length(ovars)==0) X <- rep(1,n) #no categories
  else {
      odim <- length(ovars)</pre>
      for (i in 1:odim) {
           temp <- m[[ovars[i]]]</pre>
           ctemp <- class(temp)</pre>
           if (!is.null(ctemp) && ctemp=='tcut')
                stop("Can't use tcut variables in expected survival")
      X <- strata(m[ovars])</pre>
  #do the work
  if (israte)
      temp <- survexp.fit(cbind(as.numeric(X),R), Y, newtime,</pre>
                            conditional, ratetable)
  else {
      temp <- survexp.cfit(cbind(as.numeric(X),R), Y, conditional, FALSE,</pre>
                            ratetable, se.fit=se.fit)
      newtime <- temp$times
      }
```

Now we need to package up the curves properly. Until such time as the program accepts Cox models with strata (on the some-day list), all the results can be returned as a single matrix of survivals with a common vector of times. If there was a times argument we need to subset to selected rows of the computation.

```
if (missing(times)) {
    n.risk <- temp$n
    surv <- temp$surv
    if (se.fit) err <- temp$se
    }
else {
    if (israte) keep <- match(times, newtime)
    else {
        # The result is from a Cox model, and it's list of
        # times won't match the list requested in the user's call
        # Interpolate the step function, giving survival of 1 and
        # se of 0 for requested points that precede the Cox fit's
        # first downward step. The code is like summary.survfit.</pre>
```

```
n <- length(newtime)</pre>
         keep <- approx(c(0, newtime), 0:n, xout=times,</pre>
                          method='constant', f=0, rule=2)$y
         }
    if (is.matrix(temp$surv)) {
         surv <- (rbind(1,temp$surv))[keep+1,,drop=FALSE]</pre>
         n.risk <- temp$n[pmax(1,keep),,drop=FALSE]</pre>
         if (se.fit) err <- (rbind(0,temp$se))[keep+1,,drop=FALSE]</pre>
         }
    else {
         surv <- (c(1,temp$surv))[keep+1]</pre>
         n.risk <- temp$n[pmax(1,keep)]</pre>
         if (se.fit) err \leftarrow (c(0,temp$se))[keep+1]
         }
    newtime <- times
    }
newtime <- newtime/scale</pre>
if (is.matrix(surv)) {
    dimnames(surv) <- list(NULL, levels(X))</pre>
    out <- list(call=call, surv=surv, n.risk=c(n.risk[,1]),</pre>
                       time=newtime)
    if (se.fit) out$std.err <- err</pre>
    }
else {
     out <- list(call=call, surv=c(surv), n.risk=c(n.risk),</pre>
                     time=newtime)
     if (se.fit) out$std.err <- c(err)</pre>
     }
```

Last do the standard things: add the model, x, or y components to the output if the user asked for them. (For this particular routine I can't think of a reason they every would.) Copy across summary information from the rate table computation if present, and add the method and class to the output.

```
else oldClass(out) <- c('survexp')
out</pre>
```

5 predict.coxph

This function produces various types of predicted values from a Cox model. The arguments are **obejct** The result of a call to coxph.

newdata Optionally, a new data set for which prediction is desired. If this is absent predictions are for the observations used fit the model.

type The type of prediction

- lp = the linear predictor for each observation
- risk = the risk score exp(lp) for each observation
- expected = the expected number of events
- terms = a matrix with one row per subject and one column for each term in the model.

se.fit Whether or not to return standard errors of the predictions.

na.action What to do with missing values if there is new data.

terms The terms that are desired. This option is almost never used, so rarely in fact that it's hard to justify keeping it.

collapse An optional vector of subject identifiers, over which to sum or 'collapse' the results

... All predict methods need to have a ... argument; we make no use of it however.

5.1 Setup

The first task of the routine is to reconsruct necessary data elements that were not saved as a part of the coxph fit. We will need the following components:

- for type='expected' residuals we need the original survival y. This is saved in coxph objects by default so will only need to be fetched in the highly unusual case that a user specified y=FALSE in the original call.
- for any call with either newdata, standard errors, or type='terms' the original X matrix, weights, strata, and offset. When checking for the existence of a saved X matrix we can't use object\$x since that will also match the xlevels component.
- the new data matrix, if any

```
 \langle predict.coxph \rangle \equiv \\ predict.coxph \leftarrow function(object, newdata, \\ type=c("lp", "risk", "expected", "terms"), \\ se.fit=FALSE, na.action=na.pass, \\ terms=names(object\$assign), collapse, ...) \{ \\ \langle pcoxph-init \rangle \\ \langle pcoxph-getdata \rangle \\ \langle pcoxph-expected \rangle \\ \langle pcoxph-simple \rangle \\ \langle pcoxph-terms \rangle \\ \langle pcoxph-finish \rangle \\ \}
```

We start of course with basic argument checking. Then retrieve the model parameters: does it have a strata statement, offset, etc. The $\mathtt{Terms2}$ object is a model statement without the strata or cluster terms, appropriate for recreating the matrix of covariates X. For type=expected the response variable needs to be kept, if not we remove it as well since the user's newdata might not contain one.

```
\langle pcoxph-init \rangle \equiv
  if (!inherits(object, 'coxph'))
      stop("Primary argument much be a coxph object")
 type <-match.arg(type)</pre>
 n <- object$n
 Terms <- object$terms
  if (!missing(terms)) {
      if (is.numeric(terms)) {
          if (any(terms != floor(terms) |
                   terms > length(object$assign) |
                   terms <1)) stop("Invalid terms argument")</pre>
          }
      else if (any(is.na(match(terms, names(object$assign)))))
         stop("a name given in the terms argument not found in the model")
 #Do I have strata or a cluster statment? Don't conflict names with the strata()
  # function
 strat <- attr(Terms, 'specials')$strata</pre>
 dropx <- NULL
  if (length(strat)) {
      stemp <- untangle.specials(Terms, 'strata', 1)</pre>
      dropx <- stemp$terms</pre>
  if (length(attr(Terms, 'specials')$cluster)) {
      temp <- untangle.specials(Terms, 'cluster', 1)</pre>
```

```
dropx <- c(dropx, temp$terms)
}
if (length(dropx)) Terms2 <- Terms[-dropx]
else Terms2 <- Terms
if (type != 'expected') Terms2 <- delete.response(Terms2)
na.action.used <- object$na.action
has.offset <- !is.null(attr(Terms, 'offset'))</pre>
```

The next task of the routine is to reconsruct necessary data elements that were not saved as a part of the coxph fit. We will need the following components:

- for type='expected' residuals we need the original survival y. This is saved in coxph objects by default so will only need to be fetched in the highly unusual case that a user specified y=FALSE in the original call.
- for any call with either newdata, standard errors, strata, or type='terms' the original X matrix, weights, strata, and offset. When checking for the existence of a saved X matrix we can't use object\$x since that will also match the xlevels component.
- the new data matrix, if any

For the case that none of the above are needed, we can use the linear.predictors component of the fit. The variable need.x signals this case, which takes up almost none of the code but is common in usage.

```
\langle pcoxph\text{-}getdata \rangle \equiv
  if (type == 'expected') {
      y <- object[['y']]</pre>
      if (is.null(y)) { # very rare case
           mf <- model.frame(object)</pre>
           y <- model.extract(mf, 'response')</pre>
           }
      }
  if (se.fit || !missing(newdata) || type=='terms' || length(strat)) {
      need.x <- TRUE</pre>
      if (is.null(object[['x']])) {
           mf <- model.frame(object)</pre>
           x <- model.matrix(delete.response(Terms2), mf,
                           contr=object$contrasts)[,-1,drop=FALSE]
           if (length(strat)) {
                if (length(stemp$vars)==1) oldstrat <- mf[[stemp$vars]]</pre>
                else oldstrat <- strata(mf[,stemp$vars], shortlabel=TRUE)</pre>
           else oldstrat <- rep(OL, nrow(mf))</pre>
           weights <- model.weights(mf)</pre>
           offset <- model.offset(mf)</pre>
           }
```

```
else {
        x <- object[['x']]
        oldstrat <- object$strata
        weights <- object$weights</pre>
        offset <- object$offset
    if (is.null(weights)) weights <- rep(1.0, nrow(x))</pre>
    if (is.null(offset)) offset <- rep(0.0, nrow(x))</pre>
    if (length(oldstrat)==0) oldstrat <- rep(OL, nrow(x))</pre>
    }
else {
    oldstrat <- rep(OL, n)
    offset <- 0
    need.x <- FALSE</pre>
    }
if (!missing(newdata)) {
    mf <- model.frame(Terms2, data=newdata, xlev=object$xlevels,</pre>
                        na.action=na.action)
    newx <- model.matrix(Terms2, mf,</pre>
                           contr=object$contrasts)[,-1,drop=FALSE]
    if (length(strat)) {
        if (length(stemp$vars)==1) newstrat <- mf[[stemp$vars]]</pre>
        else newstrat <- strata(mf[,stemp$vars], shortlabel=TRUE)</pre>
        if (any(is.na(match(newstrat, oldstrat))))
             stop("New data has a strata not found in the original model")
        }
    else newstrat <- rep(OL, nrow(mf))</pre>
    newoffset <- model.offset(mf)</pre>
    if (is.null(newoffset)) newoffset <- rep(0.0, nrow(newx))</pre>
    na.action.used <- attr(mf, 'na.action')</pre>
    if (type== 'expected') {
        newy <- model.response(mf)</pre>
        if (attr(newy, 'type') != attr(y, 'type'))
             stop("New data has a different survival type than the model")
        }
    }
```

5.2 Expected hazard

When we do not need standard errors the computation of expected hazard is very simple since the martingale residual is defined as status - expected. The 0/1 status is saved as the last column of y.

```
\langle pcoxph\text{-}expected \rangle \equiv
```

The more general case makes use of the [agsurv] routine to calculate a survival curve for each strata. The routine is defined in the section on individual Cox survival curves. The code here closely matches that. The routine only returns values at the death times, so we need approx to get a complete index.

One non-obvious, but careful choice is to use the residuals for the predicted value instead of the computation below, whenever operating on the original data set. This is a consequence of the Efron approx. When someone in a new data set has exactly the same time as one of the death times in the original data set, the code below implicitly makes them the "last" death in the set of tied times. The Efron approx puts a tie somewhere in the middle of the pack. This is way too hard to work out in the code below, but thankfully the original Cox model already did it. However, it does mean that a different answer will arise if you set newdata = the original coxph data set. Standard errors have the same issue, but 1. they are hardly used and 2. the original coxph doesn't do that calculation. So we do what's easiest.

```
\langle pcoxph\text{-}expected2\rangle \equiv
 ustrata <- unique(oldstrat)</pre>
 risk <- exp(object$linear.predictors)</pre>
 x <- scale(x, center=object$means, scale=FALSE)
  if (se.fit) {
      se <- double(nrow(mf))</pre>
      }
  if (missing(newdata))
      se <- double(nrow(mf))</pre>
  if (!missing(newdata)) {
      se <- double(nrow(mf))</pre>
      pred <- se
      newx <- scale(newx, center=object$means, scale=FALSE)</pre>
      newrisk <- c(exp(newx %*% object$coef))</pre>
 survtype= ifelse(fit$method=='efron', 3,2)
  for (i in ustrata) {
      indx <- which(oldstrat == i)</pre>
      afit <- agsurv(y[indx,,drop=F], x[indx,,drop=F],</pre>
                                         weights[indx], risk[indx],
                                         survtype, survtype)
      afit.n <- length(afit$time)
      if (missing(newdata)) {
           # In this case we need se.fit, nothing else
           j1 <- approx(afit$time, 1:afit.n, y[indx,1], method='constant',</pre>
```

```
f=0, yleft=0, yright=afit.n)$y
    chaz <- c(0, afit$cumhaz)[j1 +1]</pre>
    varh <- c(0, cumsum(afit$varhaz))[j1 +1]</pre>
    xbar <- rbind(0, afit$xbar)[j1+1,,drop=F]</pre>
    if (ncol(y)==2) {
        dt <- (chaz * x[indx,]) - xbar</pre>
        se[indx] <- sqrt(varh + rowSums((dt %*% object$var) *dt)) *</pre>
             risk[indx]
    else {
        j2 <- approx(afit$time, 1:afit.n, y[indx,2], method='constant',</pre>
                  f=0, yleft=0, yright=afit.n)$y
        chaz2 \leftarrow c(0, afit\$cumhaz)[j2 +1]
        varh2 <- c(0, cumsum(afit$varhaz))[j2 +1]</pre>
        xbar2 <- rbind(0, afit$xbar)[j2+1,,drop=F]</pre>
        dt <- (chaz * x[indx,]) - xbar</pre>
        v1 <- varh + rowSums((dt %*% object$var) *dt)
        dt2 \leftarrow (chaz2 * x[indx,]) - xbar2
        v2 <- varh2 + rowSums((dt2 %*% object$var) *dt2)</pre>
        se[indx] <- sqrt(v2-v1)* risk[indx]</pre>
    }
else {
    #there is new data
    indx2 <- which(newstrat == i)</pre>
    j1 <- approx(afit$time, 1:afit.n, newy[indx2,1],</pre>
                  method='constant', f=0, yleft=0, yright=afit.n)$y
    chaz <-c(0, afit$cumhaz)[j1+1]</pre>
    pred[indx2] <- chaz * newrisk[indx2]</pre>
    if (se.fit) {
        varh <- c(0, cumsum(afit$varhaz))[j1+1]</pre>
        xbar <- rbind(0, afit$xbar)[j1+1,,drop=F]</pre>
    if (ncol(y)==2) {
        if (se.fit) {
             dt <- (chaz * newx[indx2,]) - xbar[indx2,]</pre>
             se[indx2] <- sqrt(varh + rowSums((dt %*% object$var) *dt)) *</pre>
                 newrisk[indx2]
             }
        }
    else {
        j2 <- approx(afit$time, 1:afit.n, newy[indx2,2],</pre>
                  method='constant', f=0, yleft=0, yright=afit.n)$y
                      chaz2 <- approx(-afit$time, afit$cumhaz, -newy[indx2,2],</pre>
                     method="constant", rule=2, f=0)$y
```

```
chaz2 <-c(0, afit$cumhaz)[j2+1]
pred[indx2] <- (chaz2 - chaz) * newrisk[indx2]

if (se.fit) {
    varh2 <- c(0, cumsum(afit$varhaz))[j1+1]
    xbar2 <- rbind(0, afit$xbar)[j1+1,,drop=F]
    dt <- (chaz * newx[indx2,]) - xbar[indx2,]
    dt2 <- (chaz2 * newx[indx2,]) - xbar2[indx2,]

    v2 <- varh2 + rowSums((dt2 %*% object$var) *dt2)
    v1 <- varh + rowSums((dt %*% object$var) *dt)
    se[indx2] <- sqrt(v2-v1)* risk[indx2]
    }
}
</pre>
```

5.3 Linear predictor, risk, and terms

For these three options what is returned is a relative prediction which compares each observation to the average for the data set. Partly this is practical. Say for instance that a treatment covariate was coded as 0=control and 1=treatment. If the model were refit using a new coding of 3=control 4=treatment, the results of the Cox model would be exactly the same with respect to coefficients, variance, tests, etc. The raw linear predictor $X\beta$ however would change, increasing by a value of 3β . The relative predictor

$$\eta_i = X_i \beta - (1/n) \sum_j X_j \beta \tag{1}$$

will stay the same. The second reason for doing this is that the Cox model is a relative risks model rather than an absolute risks model, and thus relative predictions are almost certainly what the user was thinking of.

When the fit was for a stratified Cox model more care is needed. For instance assume that we had a fit that was stratified by sex with covaritate x, and a second data set were created where for the females x is replaced by x+3. The Cox model results will be unchanged for the two models, but the 'normalized' linear predictors $(x-\overline{x})'\beta$ will not be the same. This reflects a more fundamental issue that the for a stratified Cox model relative risks are well defined only within a stratum, i.e. for subject pairs that share a common baseline hazard. The example above is artificial, but the problem arises naturally whenever the model includes a strata by covariate interaction. So for a stratified Cox model the predictions should be forced to sum to zero within each stratum, or equivalently be made relative to the weighted mean of the stratum. Unfortunately, this important issue was not realized until late in 2009 when a puzzling query was sent to the author involving the results from such an interaction. Note that this issue did not arise with type='expected', which has a natural scaling.

An offset variable, if specified, is treated like any other covariate with respect to centering. The logic for this choice is not as compelling, but it seemed the best that I could do. Note that offsets play no role whatever in predicted terms, only in the lp and risk.

```
Start with the simple ones
\langle pcoxph\text{-}simple \rangle \equiv
  if (is.null(object$coefficients))
      coef<-numeric(0)</pre>
  else {
      # Replace any NA coefs with 0, to stop NA in the linear predictor
      coef <- ifelse(is.na(object$coefficients), 0, object$coefficients)</pre>
      }
  if (missing(newdata)) {
      offset <- offset - mean(offset)
      if (length(strat)) {
          for (i in unique(oldstrat)) {
               j <- which(oldstrat==i)</pre>
               if (length(j)==1) x[j,] \leftarrow 0 # only 1 subject in the strata
               else if (length(j) >1) {
                   xmean <- colSums(x[j,] * weights[j]) / sum(weights[j])</pre>
                   x[j,] <- scale(x[j,], center=xmean, scale=FALSE)</pre>
               }
          newx <- x
      else if (need.x) newx <- scale(x, center=object$means, scale=FALSE)
      }
  else {
      offset <- newoffset - mean(offset)
      if (length(strat)) {
          for (i in unique(oldstrat)) {
               j <- which(newstrat==i)</pre>
                                            #no matches is a possibility
               if (length(j) ==1) newx[j,] <- 0
               else if (length(j) >1) {
                   xmean <- colSums(x[j,,drop=FALSE] * weights[j]) / sum(weights[j])</pre>
                   newx[j,] <- scale(newx[j,], center=xmean, scale=FALSE)</pre>
                   }
          }
      else newx <- scale(newx, center=object$means, scale=FALSE)</pre>
  if (type=='lp' || type=='risk') {
      if (need.x) pred <- newx %*% coef + offset
      else pred <- object$linear.predictors</pre>
      if (se.fit) se <- sqrt(rowSums((newx %*% object$var) *newx))</pre>
      if (type=='risk') {
```

```
pred <- exp(pred)
if (se.fit) se <- se * sqrt(pred) # standard Taylor series approx
}
</pre>
```

The type=terms residuals are a bit more work. In Splus this code used the Build.terms function, which was essentially the code from predict.lm extracted out as a separate function. As of March 2010 (today) a check of the Splus function and the R code for predict.lm revealed no important differences. A lot of the bookkeeping in both is to work around any possible NA coefficients resulting from a singularity. The basic formula is to

- 1. If the model has an intercept, then sweep the column means out of the X matrix. We've already done this.
- 2. For each term separately, get the list of coefficients that belong to that term; call this list
- 3. Restrict X, β and V (the variance matrix) to that subset, then the linear predictor is $X\beta$ with variance matrix XVX'. The standard errors are the square root of the diagonal of this latter matrix. This can be computed, as colSums((X

Note that the assign component of a coxph object is the same as that found in Splus models (a list), most R models retain a numeric vector which contains the same information but it is not as easily used. The first first part of predict.lm in R rebuilds the list form as its asgn variable. I can skip this part since it is already done.

```
\langle pcoxph\text{-}terms \rangle \equiv
  else if (type=='terms') {
      asgn <- object$assign
      nterms<-length(asgn)
      pred<-matrix(ncol=nterms,nrow=NROW(newx))</pre>
      dimnames(pred) <- list(rownames(newx), names(asgn))</pre>
      if (se.fit) se <- pred
      for (i in 1:nterms) {
           tt <- asgn[[i]]
           tt <- tt[!is.na(object$coefficients[tt])]</pre>
           xtt <- newx[,tt, drop=F]</pre>
           pred[,i] <- xtt %*% object$coefficient[tt]</pre>
           if (se.fit)
                se[,i] <- sqrt(rowSums((xtt %*% object$var[tt,tt]) *xtt))</pre>
      pred <- pred[,terms, drop=F]</pre>
      if (se.fit) se <- se[,terms, drop=F]</pre>
      attr(pred, 'constant') <- sum(object$coefficients*object$means, na.rm=T)
```

To finish up we need to first expand out any missings in the result based on the na action, and optionally collapse the results within a subject. What should we do about the standard errors when collapse is specified? We assume that the individual pieces are independent and thus var(sum) = sum(variances). The statistical justification of this is quite solid for the linear predictor, risk and terms type of prediction due to independent increments in a martingale. For expecteds the individual terms are positively correlated so the se will be too small. One solution would be to refuse to return an se in this case, but the bias should usually be small, and besides it would be unkind to the user.

Prediction of type='terms' is expected to always return a matrix, or the R termplot() function gets unhappy.

```
\langle \mathit{pcoxph-finish} \rangle {\equiv}
  if (type != 'terms') {
      pred <- drop(pred)</pre>
      if (se.fit) se <- drop(se)
      }
  if (!is.null(na.action.used)) {
      pred <- naresid(na.action.used, pred)</pre>
      if (is.matrix(pred)) n <- nrow(pred)</pre>
                            n <- length(pred)</pre>
      if(se.fit) se <- naresid(na.action.used, se)</pre>
  if (!missing(collapse)) {
      if (length(collapse) != n) stop("Collapse vector is the wrong length")
      pred <- rowsum(pred, collapse) # in R, rowsum is a matrix, always</pre>
      if (se.fit) se <- sqrt(rowsum(se^2, collapse))</pre>
      if (type != 'terms') {
           pred <- drop(pred)</pre>
           if (se.fit) se <- drop(se)
      }
  if (se.fit) list(fit=pred, se.fit=se)
  else pred
```