

# Spline terms in a Cox model

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This is a trio of topics that comes up just often enough in my work that I end up re-discovering how to do it correctly about once a year. A note showing how may be useful to others, it is certainly a useful reference for me.

## 1 Plotting smooth terms

Here is a simple example using the MGUS data. I prefer a simpler color palette than the default found in `termplot`.

```
> library(survival)
> mfit <- coxph(Surv(futime, death) ~ sex + pspline(age, df=4), data=mgus)
> mfit
Call:
coxph(formula = Surv(futime, death) ~ sex + pspline(age, df = 4),
      data = mgus)
```

|                           | coef     | se(coef) | se2     |
|---------------------------|----------|----------|---------|
| sexmale                   | 0.22784  | 0.13883  | 0.13820 |
| pspline(age, df = 4), lin | 0.06682  | 0.00703  | 0.00703 |
| pspline(age, df = 4), non |          |          |         |
|                           | Chisq    | DF       | p       |
| sexmale                   | 2.69335  | 1.00     | 0.10    |
| pspline(age, df = 4), lin | 90.22974 | 1.00     | <2e-16  |
| pspline(age, df = 4), non | 3.44005  | 3.05     | 0.34    |

```
Iterations: 5 outer, 16 Newton-Raphson
```

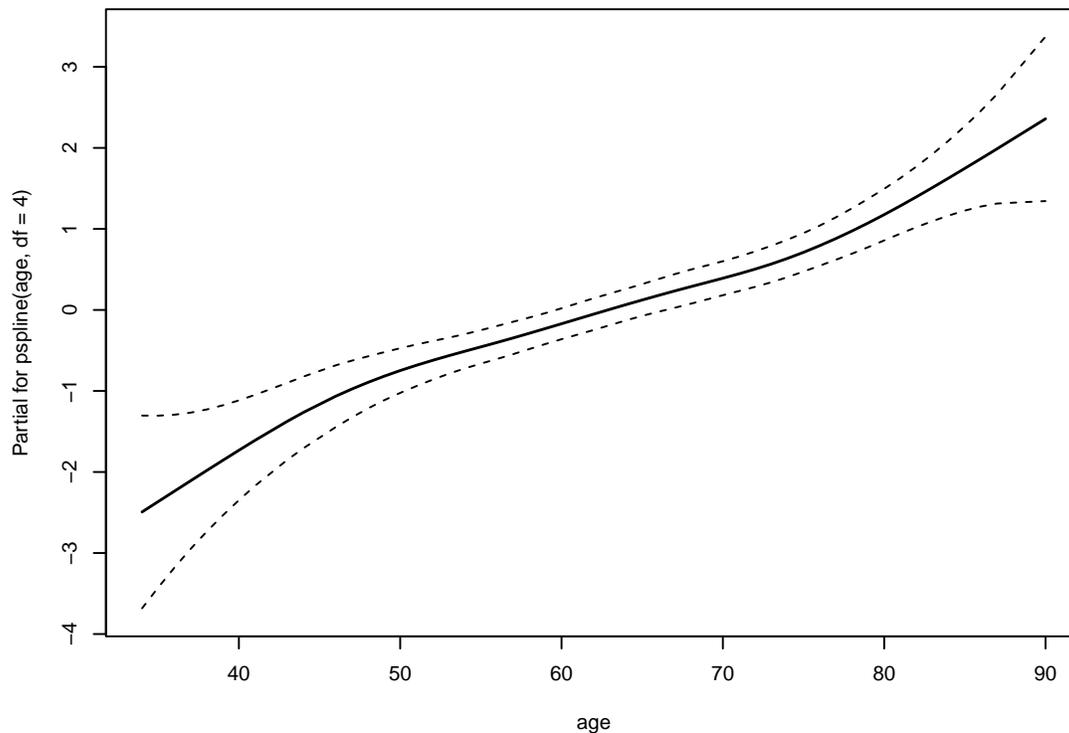
```
Theta= 0.851
```

```
Degrees of freedom for terms= 1.0 4.1
```

```
Likelihood ratio test=108 on 5.04 df, p=<2e-16
```

```
n= 241, number of events= 225
```

```
> termplot(mfit, term=2, se=TRUE, col.term=1, col.se=1)
```



Note that the `term=2` option is passed directly from the `termplot` routine to a `predict(fit, type='terms')` call. For `coxph` models, the `predict` function allows terms to be specified either by position or name. Other routines, e.g. `gam`, respond only to a name. (This can be a bit of a pain since it must exactly match the *printed* call in both spelling and spacing; and the printed spacing may not match what the user typed.)

Three questions are whether the curve is significantly non-linear, how the curve is centered and whether we can easily plot it on the hazard as opposed to the log hazard scale. The first question is answered by the `printout`, the solution to the others is to use the `plot=FALSE` option of `termplot`, which returns the data points that would be plotted back to the user.

```
> ptemp <- termplot(mfit, se=TRUE, plot=FALSE)
> attributes(ptemp)
$constant
[1] 3.205273

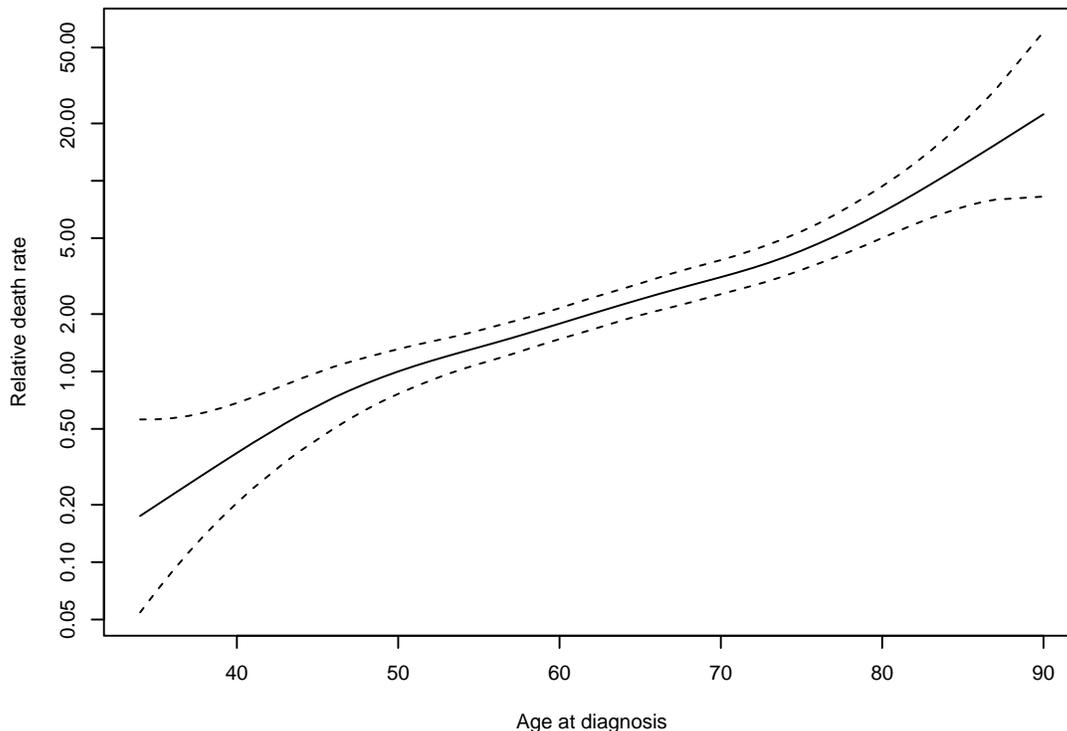
$names
[1] "sex" "age"
> ptemp$age[1:4,]
  x      y      se
1 34 -2.493101 0.5946620
2 35 -2.365152 0.5301362
3 36 -2.237191 0.4720479
```

4 37 -2.109559 0.4208977

The `termplot` function depends on a call to `predict` with `type='terms'`, which returns a centered set of predictions. Like a simple linear model fit, the intercept is a separate term, which is found in the “constant” attribute above, and each column of the result is centered so that the average predicted value is zero. Since any given  $x$  value may appear multiple times in the data and thus in the result of `predict`, and the `termplot` function removes duplicates, the data returned by `termplot` may not be precisely centered at zero.

Now suppose we want to redraw this on log scale with age 50 as the reference, i.e., the risk is 1 for a 50 year old. Since the Cox model is a relative hazards model we can choose whatever center we like. (If there were no one of exactly age 50 in the data set the first line below would need to do an interpolation, e.g. by using the `approx` function.)

```
> ageterm <- ptemp$age # this will be a data frame
> center <- with(ageterm, y[x==50])
> ytemp <- ageterm$y + outer(ageterm$se, c(0, -1.96, 1.96), '*')
> matplot(ageterm$x, exp(ytemp - center), log='y',
          type='l', lty=c(1,2,2), col=1,
          xlab="Age at diagnosis", ylab="Relative death rate")
```



Voila! We now have a plot that is interpretable with respect to a fixed reference.

The approach is appropriate for any term, not just psplines. The above plot uses log scale for the y axis which is appropriate for the question of whether a non-linear age effect was

even necessary for this model (it is not), one could remove the log argument to emphasize the Gompertzian effect of age on mortality.

## 2 Monotone splines

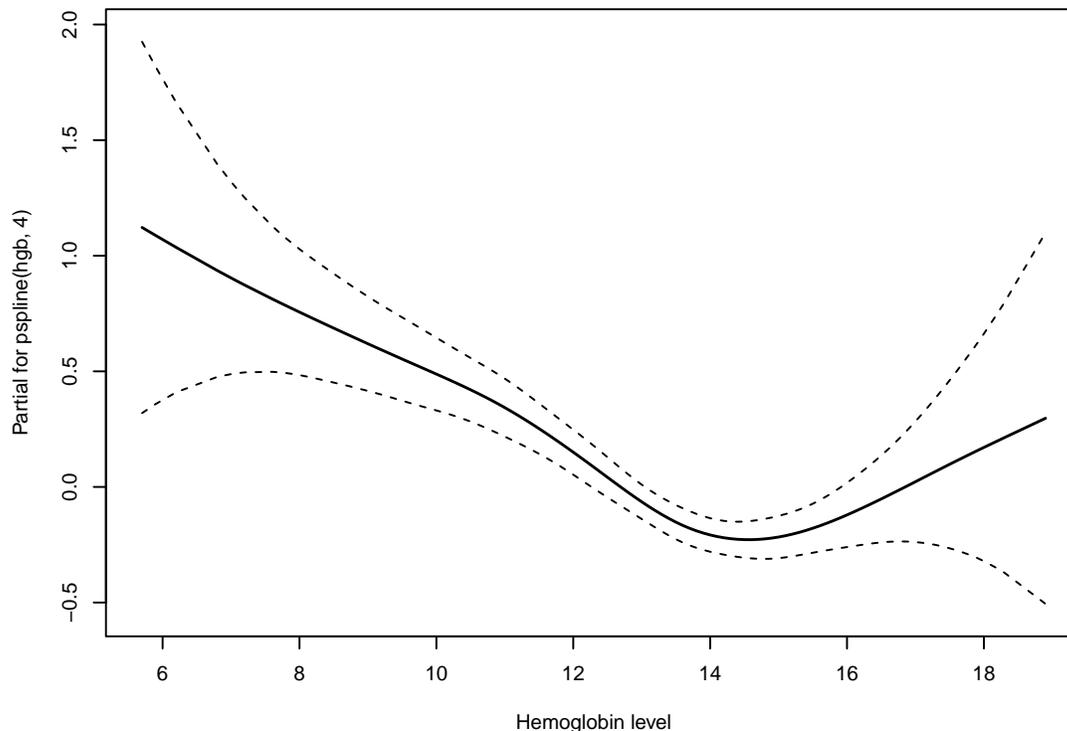
Consider the following model using the `mgus2` data set.

```
> fit <- coxph(Surv(futime, death) ~ age + pspline(hgb, 4), mgus2)
> fit
Call:
coxph(formula = Surv(futime, death) ~ age + pspline(hgb, 4),
      data = mgus2)

              coef  se(coef)      se2
age           0.05394  0.00337  0.00336
pspline(hgb, 4), linear -0.11579  0.01612  0.01612
pspline(hgb, 4), nonlin

              Chisq  DF      p
age           256.31805  1.00 < 2e-16
pspline(hgb, 4), linear  51.56708  1.00 6.9e-13
pspline(hgb, 4), nonlin  18.77963  3.06 0.00033

Iterations: 6 outer, 16 Newton-Raphson
      Theta= 0.942
Degrees of freedom for terms= 1.0 4.1
Likelihood ratio test=420 on 5.06 df, p=<2e-16
n= 1371, number of events= 957
      (13 observations deleted due to missingness)
> termplot(fit, se=TRUE, term=2, col.term=1, col.se=1,
           xlab="Hemoglobin level")
```



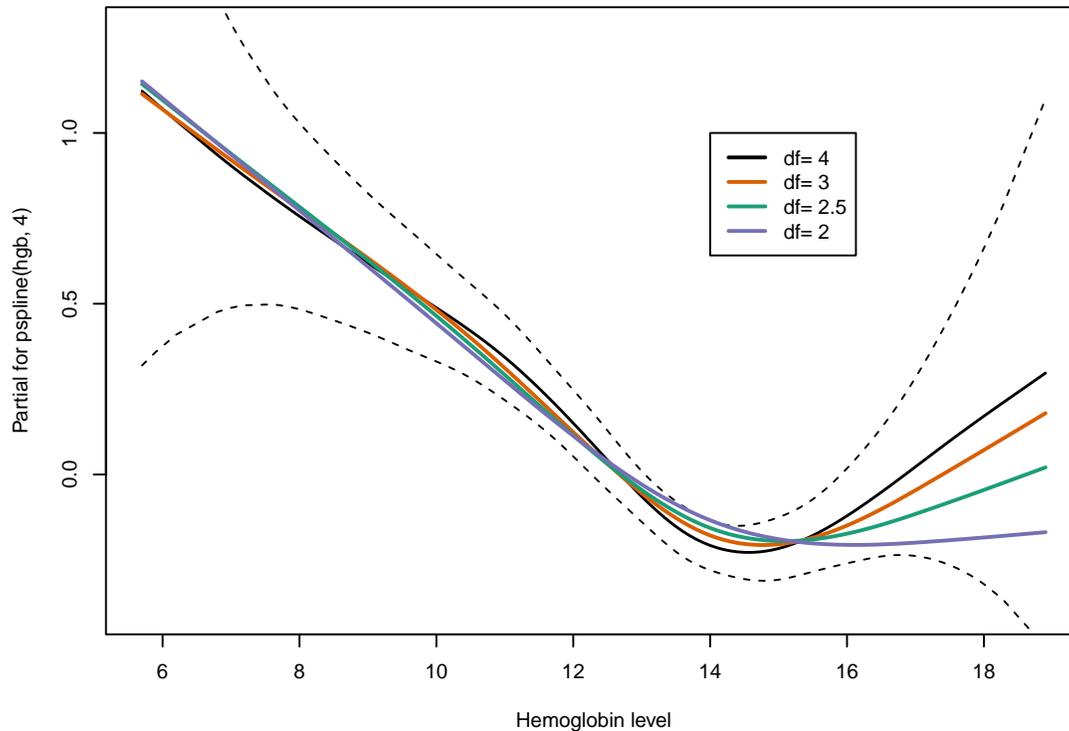
Low hemoglobin or anemia is a recognized marker of frailty in older age, so the rise in risk for low levels is not surprising. The rise on the right hand portion of the curve is less believable — the normal range of HGB is 12-15.5 for women and 13.5 to 17.5 for men, why would we expect a rise there? A monotone fit that forces the curve to be horizontal from 14 onward fits well within the confidence bands, so we might want to force monotonicity.

There are two tools for this within the pspline function. The first is to decrease the overall degrees of freedom and the second is to use `combine` option to force equality of selected coefficients. Start by decreasing the degrees of freedom. The pspline function automatically picks the number of basis (`nterms`) to be “sufficiently large” for the given degrees of freedom. We fix it at a single value for the rest of this example to better isolate the effects of degrees of freedom and of constraints.

```

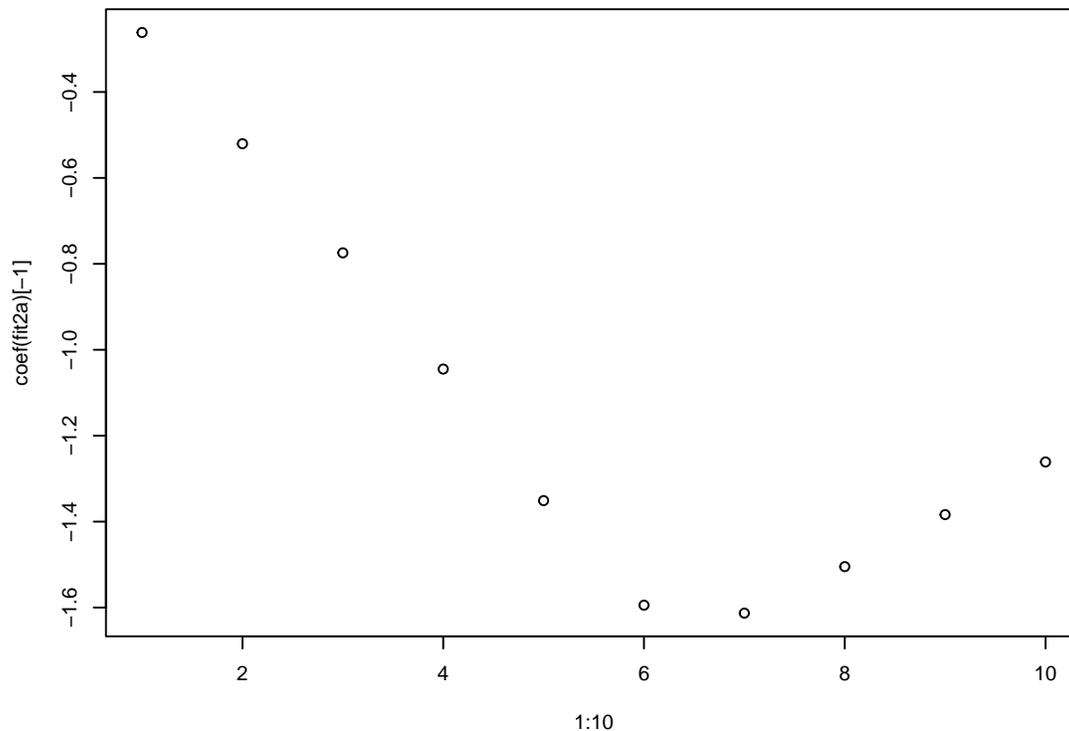
> termplot(fit, se=TRUE, col.term=1, col.se=1, term=2,
           xlab="Hemoglobin level", ylim=c(-.4, 1.3))
> df <- c(3, 2.5, 2)
> for (i in 1:3) {
  tfit <- coxph(Surv(futime, death) ~ age +
               pspline(hgb, df[i], nterm=8), mgus2)
  temp <- termplot(tfit, se=FALSE, plot=FALSE, term=2)
  lines(temp$hgb$x, temp$hgb$y, col=i+1, lwd=2)
}
> legend(14, 1, paste("df=", c(4, df)), lty=1, col=1:4, lwd=2)

```



This has reduced, but not eliminated, the right hand rise at the expense of a less sharp transition at the value of 14. The `combine` option makes use of a property of the P-spline basis, which is that the curve will be monotone if and only if the coefficients are monotone. We can then use a pool adjacent violators algorithm to sequentially force equality for those coefficients which go the wrong way. Look at the coefficients for the fit with 2.5 degrees of freedom.

```
> fit2a <- coxph(Surv(futime, death) ~ age + pspline(hgb, 2.5, nterm=8), mgus2)
> coef(fit2a)
      age  ps(hgb)3  ps(hgb)4  ps(hgb)5  ps(hgb)6
0.05399819 -0.26149468 -0.52044496 -0.77438892 -1.04488355
  ps(hgb)7  ps(hgb)8  ps(hgb)9  ps(hgb)10  ps(hgb)11
-1.35106599 -1.59448244 -1.61297463 -1.50478222 -1.38368954
  ps(hgb)12
-1.26113182
> plot(1:10, coef(fit2a)[-1])
```

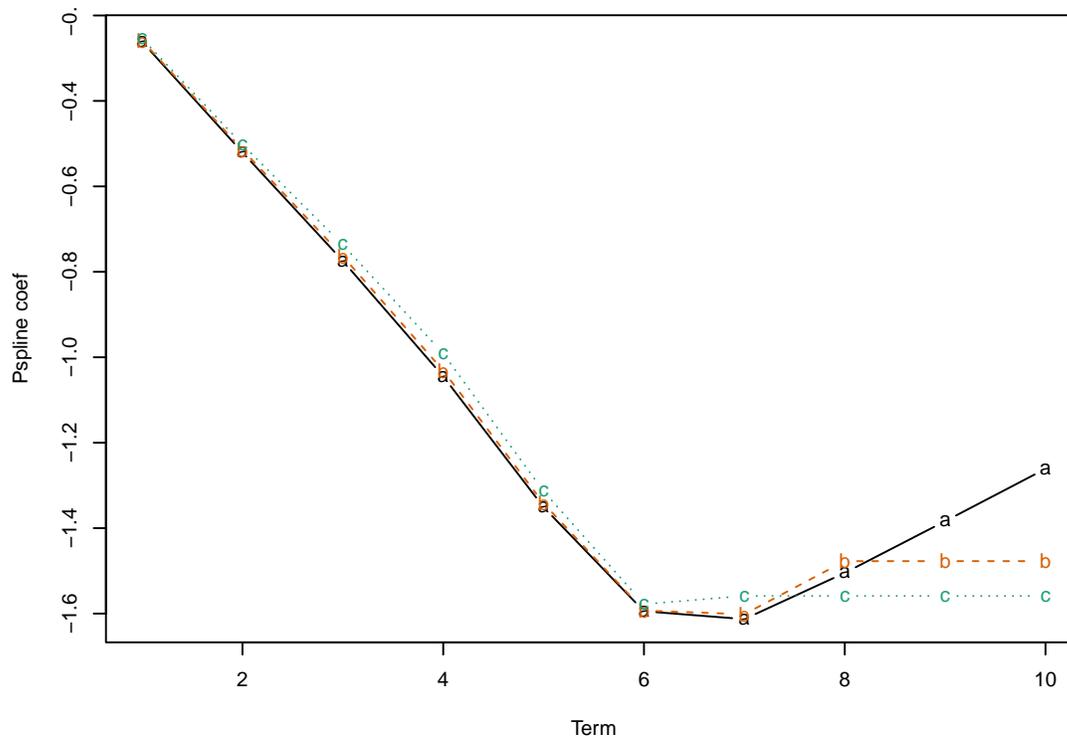


Now force the last 3 to be equal, then the last 4, and see how this changes the fit.

```

> temp <- c(1:7, 8,8,8)
> fit2b <- coxph(Surv(futime, death) ~ age +
  pspline(hgb, 2.5, nterm=8, combine=temp),
  data= mgus2)
> temp2 <- c(1:6, 7,7,7,7)
> fit2c <- coxph(Surv(futime, death) ~ age +
  pspline(hgb, 2.5, nterm=8, combine=temp2),
  data= mgus2)
> matplot(1:10, cbind(coef(fit2a)[-1], coef(fit2b)[temp+1],
  coef(fit2c)[temp2+1]), type='b', pch='abc',
  xlab="Term", ylab="Pspline coef")

```



We see that constraining the last four terms along with a degrees of freedom of is almost enough to force monotonicity; it may be sufficient if our goal is a simple plot for display.

This dance between degrees of freedom, number of terms, and constraints has a component of artistry. When all three values become large the result will begin to approach a step function, reminiscent of non-parametric isotonic regression, whereas small values begin to approach a linear fit. The best compromise of smoothness and constraints will be problem specific.

### 3 Splines in an interaction

As an example we will use the effect of age on survival in the `flchain` data set, a population based sample of subjects from Olmsted County, Minnesota. If we look at a simple model using age and sex we see that both are very significant.

```
> options(show.signif.stars=FALSE) # display intelligence
> fit1 <- coxph(Surv(futime, death) ~ sex + pspline(age, 3), data=flchain)
> fit1
Call:
coxph(formula = Surv(futime, death) ~ sex + pspline(age, 3),
      data = flchain)

              coef se(coef)      se2   Chisq
sexM              4.09e-01 4.40e-02 4.40e-02 8.64e+01
```

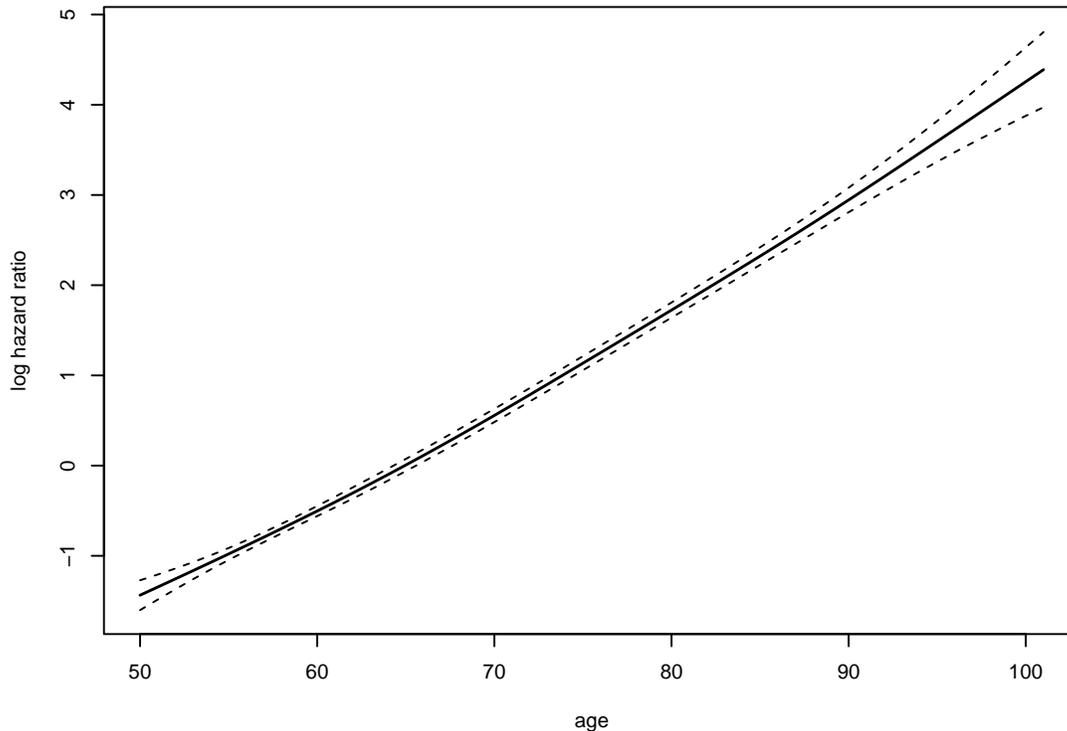
```

pspline(age, 3), linear 1.12e-01 2.20e-03 2.20e-03 2.59e+03
pspline(age, 3), nonlin                                     1.09e+01

sexM              DF      p
pspline(age, 3), linear 1.00 <2e-16
pspline(age, 3), nonlin 2.09 0.0048

Iterations: 7 outer, 20 Newton-Raphson
Theta= 0.989
Degrees of freedom for terms= 1.0 3.1
Likelihood ratio test=2629 on 4.09 df, p=<2e-16
n= 7874, number of events= 2169
> termplot(fit1, term=2, se=TRUE, col.term=1, col.se=1,
           ylab="log hazard ratio")

```



We used a `pspline` term rather than `ns`, say, because the printout for a `pspline` nicely segregates the linear and non-linear age effects. The non-linearity is not very large, as compared to the linear portion, but still may be important.

We would like to go forward and fit separate age curves for the males and the females, since the above fit makes an untested assumption that the male/female ratio of death rates will be the same at all ages. The primary problem is that a formula of `sex * pspline(age)` does not work; the `coxph` routine is not clever enough to do the right thing automatically for `pspline`

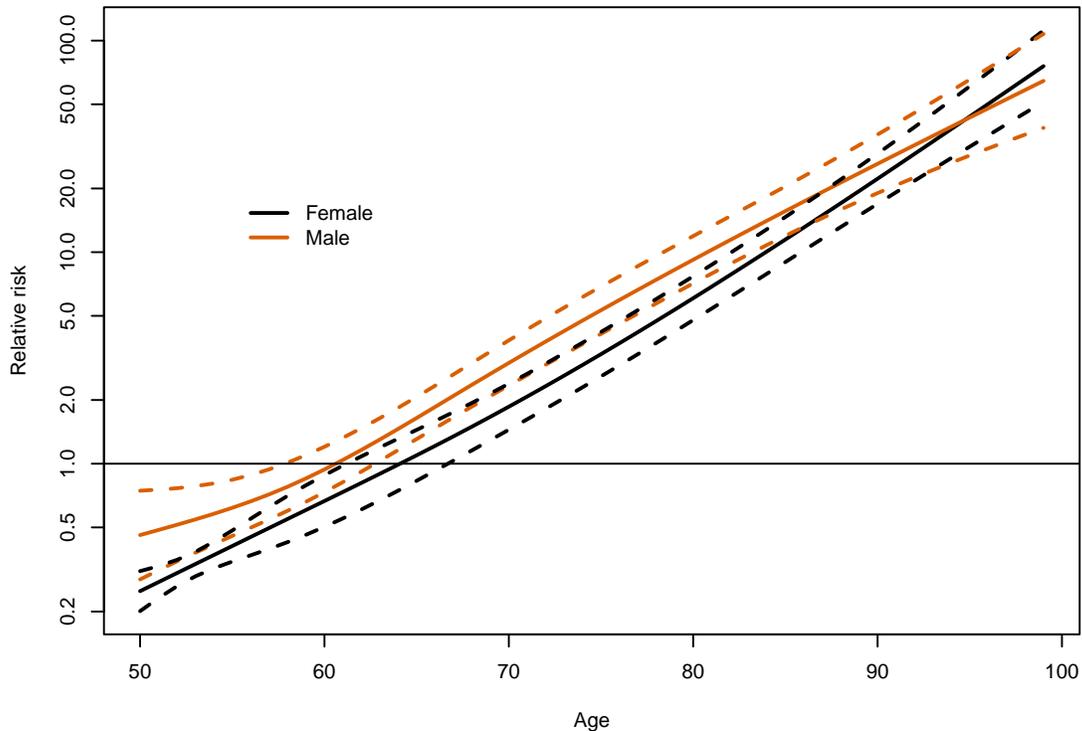
interactions. (Perhaps some future version will be sufficiently intelligent, but don't hold your breath).

As a first solution we will use regression splines, i.e., splines that can be represented using a basis matrix.

```
> options(show.signif.stars=FALSE) # display statistical intelligence
> require(splines, quietly=TRUE)
> nfit1 <- coxph(Surv(futime, death) ~ sex + age, flchain)
> nfit2 <- coxph(Surv(futime, death) ~ sex + ns(age, df=3), flchain)
> nfit3 <- coxph(Surv(futime, death) ~ sex * ns(age, df=3), flchain)
> anova(nfit1, nfit2, nfit3)
Analysis of Deviance Table
Cox model: response is Surv(futime, death)
Model 1: ~ sex + age
Model 2: ~ sex + ns(age, df = 3)
Model 3: ~ sex * ns(age, df = 3)
  loglik  Chisq Df Pr(>|Chi|)
1 -17558
2 -17554 8.8627 2    0.0119
3 -17552 4.4947 3    0.2128
```

The nonlinear term is significant but the interaction is not. Nevertheless we would like to plot the two estimated curves for `nfit3`, expecting that they will be approximately parallel. The `termplot` routine is not able to deal with models that have an interaction and will bow out with a warning message; we use explicit prediction instead, which is nearly as easy.

```
> pdata <- expand.grid(age= 50:99, sex=c("F", "M"))
> pdata[1:5,]
  age sex
1  50  F
2  51  F
3  52  F
4  53  F
5  54  F
> ypred <- predict(nfit3, newdata=pdata, se=TRUE)
> yy <- ypred$fit + outer(ypred$se, c(0, -1.96, 1.96), '*')
> matplot(50:99, exp(matrix(yy, ncol=6)), type='l', lty=c(1,1,2,2,2,2),
  lwd=2, col=1:2, log='y',
  xlab="Age", ylab="Relative risk")
> legend(55, 20, c("Female", "Male"), lty=1, lwd=2, col=1:2, bty='n')
> abline(h=1)
```



The `ns` function generates a basis of dummy variables to represent the spline, which will work automatically in interactions. The coefficients that result are not very interpretable, but the result of `predict` is invariant to this. The issues as compared to using `termplot` are

1. We need to provide our own set of predictor values for the plot, whereas `termplot` would automatically use the set of unique age and sex values.
2. Keeping track of the indexing. The `predict` function produces two vectors each of length `nrow(pdata)` with a predicted value and its standard error, one value for each row of `pdata`. The data set is in order of females, then males. We fold it into an appropriate matrix for use with `matplot`.
3. The vertical centering of the curves corresponds to an average population predictor of  $\eta = X\beta = 0$ ; i.e., the average over the subjects in the data set. This is a consequence of the variable centering that `coxph` does for numerical stability. To center at some particular (age, sex) pair obtain the predicted value for that fictional subject and subtract the resulting value from `yy`.

To create the same figure with `pspline` curves it is necessary to code the males and females as two separate terms. To do this create our own dummy variables to handle the interaction.

```
> agem <- with(flchain, ifelse(sex=="M", age, 60))
> agef <- with(flchain, ifelse(sex=="F", age, 60))
> fit2 <- coxph(Surv(futime, death) ~ sex + pspline(agef, df=3))
```

```

+ pspline(agem, df=3), data=flchain)
> anova(fit2, fit1)
Analysis of Deviance Table
Cox model: response is Surv(futime, death)
Model 1: ~ sex + pspline(agef, df = 3) + pspline(agem, df = 3)
Model 2: ~ sex + pspline(age, 3)
  loglik Chisq    Df Pr(>|Chi|)
1 -17551
2 -17554 5.8211 2.8583    0.1096

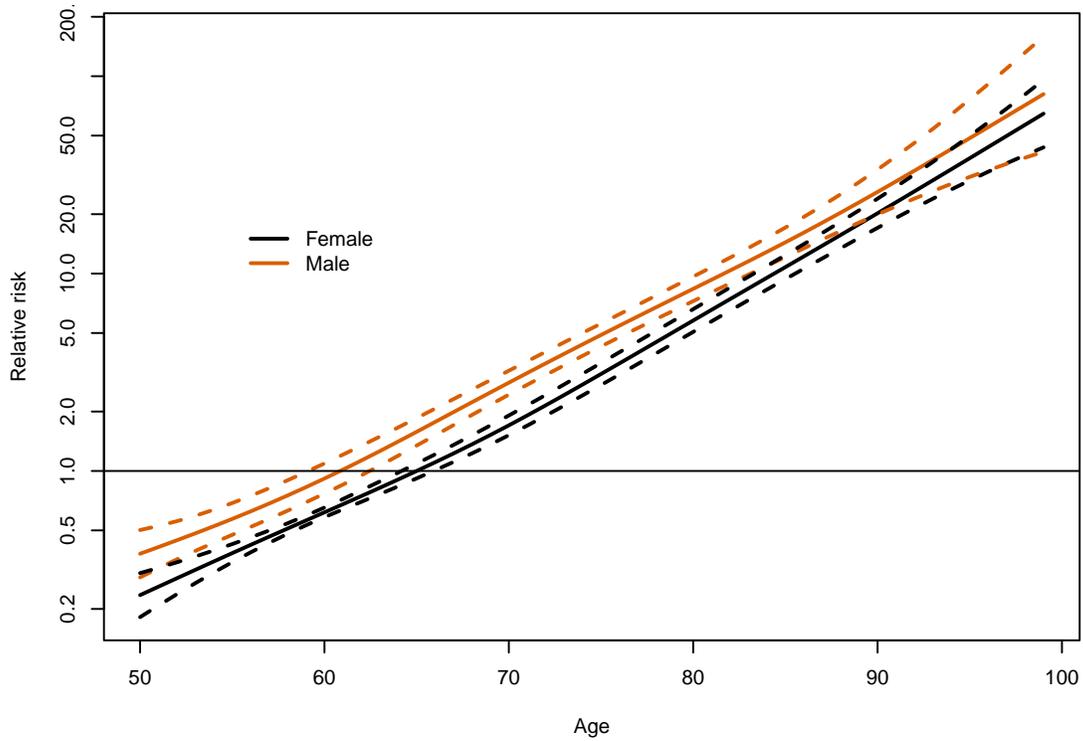
```

You might well ask why we used 60 as a dummy value of `agem` for the females instead of 0? If a value of 0 is used it forces the `pspline` function to create a basis set that includes all the empty space between 0 and 50, and do predictions at 0; these last can become numerically unstable leading to errors or incorrect values. Best is to pick a value close to the mean, though any value within the range will do. For this plot we will use a 65 year old female as the reference.

```

> # predictions
> pdata2 <- pdata
> pdata2$agem <- with(pdata2, ifelse(sex=="M", age, 60))
> pdata2$agef <- with(pdata2, ifelse(sex=="F", age, 60))
> ypred2 <- predict(fit2, pdata2, se=TRUE)
> yy <- ypred2$fit + outer(ypred2$se, c(0, -1.96, 1.96), '*')
> # reference
> refdata <- data.frame(sex='F', agef=65, agem=60)
> ref <- predict(fit2, newdata=refdata, type="lp")
> # plot
> matplot(50:99, exp(matrix(yy-ref, ncol=6)), type='l', lty=c(1,1,2,2,2,2),
  lwd=2, col=1:2, log='y',
  xlab="Age", ylab="Relative risk")
> legend(55, 20, c("Female", "Male"), lty=1, lwd=2, col=1:2, bty='n')
> abline(h=1)

```



The final curves for males and female are not quite parallel, most of the difference is at the highest ages, however, where there are very few subjects. One thing the plot does not display is that the spacing between the male and female points also has a standard error. This moves the entire bundle of three red curves up and down. It is not clear how best to add this information into the plot. For questions of parallelism and shape, as here, it seemed best to ignore it, which is what the termplot function also does. If someone were reading individual male/female differences off the plot a different choice would be appropriate.