

Regression in R

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Contents

1	Understanding Regression	5
1.1	1. Statistical: the matrix formulation of a model	5
1.2	2. Mathematical: the formula for the model	6
1.3	3. Computing: commands and algorithms that fit the model . .	6
1.4	4. Graphical: data space	8

1.5	5. Graphical: beta space	10
1.6	6. Graphical (different definition!): Path diagram of variables . .	13
1.7	7. Geometric: Hilbert space representation of variables or ‘variable space’	13
1.8	8. Most important: Real world interpretation	13
2	Review of the matrix formulation and the general linear hy- pothesis (GLH)	16
2.1	Linear hypotheses	17
2.1.1	Exercise:	18
2.1.2	Example:	18
2.2	Estimation and tests	19
2.2.1	Notes	20
2.2.2	Exercises:	21
3	Interpreting Regression Coefficients: Smoking and Life Ex- pectancy	21
3.1	Functions to test linear hypotheses	44
3.2	Strategies to simplify models	49

3.3	Estimating effects over a grid	50
3.3.1	Exercises	69
3.4	Wald tests vs Likelihood Ratio Tests (LRT)	70
3.4.1	Exercises	76
3.5	Interpreting sequential tests	76
3.6	Working with factors	81
3.6.1	Exercises:	95
3.6.2	Reparametrization to answer different questions	100
3.6.3	Equivalent models	104
3.6.4	Exercise:	110
3.7	Using Lfx with factors	129
3.7.1	Exercises	148
3.8	Using WHO regions as predictors of Life Expectancy	148
3.8.1	Exercise	183
4	Exploring Regression Using R	183
4.1	Interactive 3D	185
4.1.1	Controlling for Health	189
4.2	A more interesting model?	192

4.2.1	Simultaneous tests of groups of coefficients:	200
4.3	Two valid tests:	201
4.4	Explore interactions	206
4.4.1	Some comments on reading a model	209
4.5	Regression diagnostics – quick	220
4.5.1	Visualize fit for diagnostics	233
4.6	Asking questions:	266
4.6.1	Can we simplify the model?	266
4.6.2	Asking specific questions	275
4.7	Understanding coefficients	278
4.7.1	How can we get answers to meaningful questions?	279
4.7.2	Plotting fitted values and bands	297
4.7.3	In the future:	302
5	Appendices	302
5.1	Notes on the Principle of Marginality	302
	References	305

1 Understanding Regression

To really understand regression, you need to be able to approach a problem from many different angles. I can think of at least 8 representations that complement each other. To master regression you need to know how to go from one representation to another and you need to know how to work within the right representation to think about your problem and to solve it.

Here are eight ways of thinking about regression. Some are very powerful for developing the mathematical theory of regression, other are best suited to visualize the interpretation of coefficients for a particular application.

1.1 1. Statistical: the matrix formulation of a model

$$Y = X\beta + \epsilon, \epsilon \sim N(0, \sigma^2)$$

and all the theory that follows, e.g.

$$\text{Var}(\hat{\beta}) = \sigma^2 (X'X)^{-1}$$

$$\hat{Y} = X(X'X)^{-1}X'Y = P_X Y$$

Hat

where P_X is the matrix of the orthogonal projection of \mathbb{R}^n onto $\text{span}(X)$.

1.2 2. Mathematical: the formula for the model

$$y_i = \beta_0 + \beta_1 x_i + \beta_2 x_i^2 + \beta_3 z_i + \beta_4 x_i z_i + \epsilon_i$$

$$\frac{\partial E(y)}{\partial x} = \beta_1 + 2\beta_2 x + \beta_4 z$$

1.3 3. Computing: commands and algorithms that fit the model

```
library(car)
fit <- lm(income ~ education * type, data = Prestige)
summary(fit)
```

|

| Call:

| lm(formula = income ~ education * type, data = Prestige)

Residuals:

Min	1Q	Median	3Q	Max
-6330.8	-1769.2	-356.8	1166.5	17326.2

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	-1865.0	3682.3	-0.506	0.6137
education	866.0	436.4	1.984	0.0502
typeprof	-3068.4	7191.8	-0.427	0.6706
typewc	3646.5	9274.0	0.393	0.6951
education:typeprof	234.0	617.3	0.379	0.7055
education:typewc	-569.2	884.8	-0.643	0.5216

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 3333 on 92 degrees of freedom

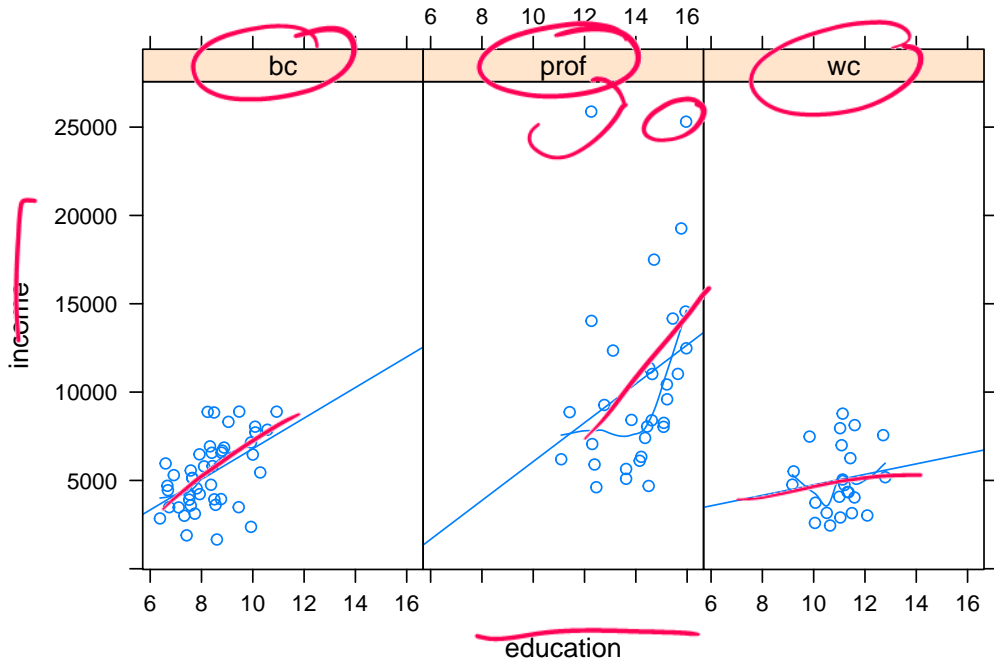
(4 observations deleted due to missingness)

Multiple R-squared: 0.4105, Adjusted R-squared: 0.3785

| F-statistic: 12.81 on 5 and 92 DF, p-value: 1.856e-09

1.4 4. Graphical: data space

```
library(car)
library(lattice)
xyplot(income ~ education | type, Prestige,
       type = c('p', 'r', 'smooth'))
```

1.5 5. Graphical: beta space

```
library(spida2)
library(latticeExtra)
fit <- lm(income ~ education + women, data = Prestige)
summary(fit)
```

```
|
| Call:
| lm(formula = income ~ education + women, data = Prestige)
|
| Residuals:
|      Min       1Q   Median       3Q      Max
| -7257.6 -1160.1  -238.6    681.1 16044.3
|
| Coefficients:
|              Estimate Std. Error t value Pr(>|t|)
| (Intercept) -1491.998    1162.299  -1.284   0.202
| education    944.881     103.731   9.109 9.60e-15 ***
```

$y = \beta_2 - \beta_1$

```
| women ' -64.056  $\beta_2$  8.921 -7.180 1.31e-10 ***  
| ---  
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1  
|  
| Residual standard error: 2839 on 99 degrees of freedom  
| Multiple R-squared:  0.5618,    Adjusted R-squared:  0.5529  
| F-statistic: 63.46 on 2 and 99 DF,  p-value: < 2.2e-16
```

```
plot(rbind(cell(fit),0),type= 'n',  
     xlab = expression(beta[education]),  
     ylab = expression(beta[women]))  
lines(cell(fit,dfn=2), type = 'l', col = 'blue')  
lines(cell(fit,dfn=1), type = 'l', col = 'red')  
abline(h=0)  
abline(v=0)  
points(c(0,0), pch = 18)
```

$H_0: \beta_W = \beta_E = 0$

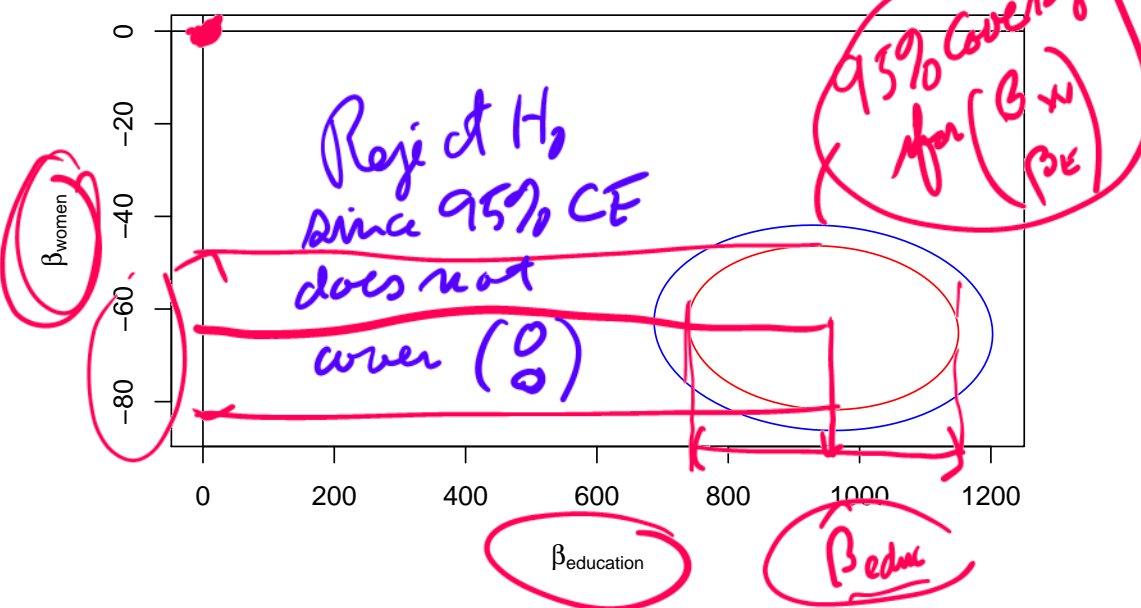


Figure: Confidence ellipse for two parameters jointly. The blue ellipse has 95% coverage in 2 dimensions and its perpendicular shadows onto the vertical and horizontal axes form Scheffe 95% confidence intervals for testing in a space of dimension 2. The similar shadows of the red ellipse provide ordinary 95% confidence intervals.

1.6 6. Graphical (different definition!): Path diagram of variables

1.7 7. Geometric: Hilbert space representation of variables or 'variable space'

1.8 8. Most important: Real world interpretation

The most important representation is the interpretation of the model in the real world. Real world factors, such as the design, the nature of random assignment, the nature of random selection are fundamental in determining the interpretation of the model and on the strategy for model development, selection and interpretation.

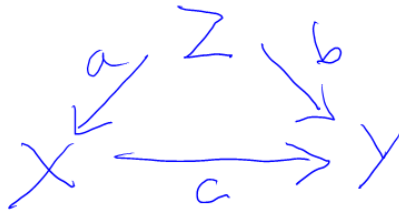
Causal



Mediation



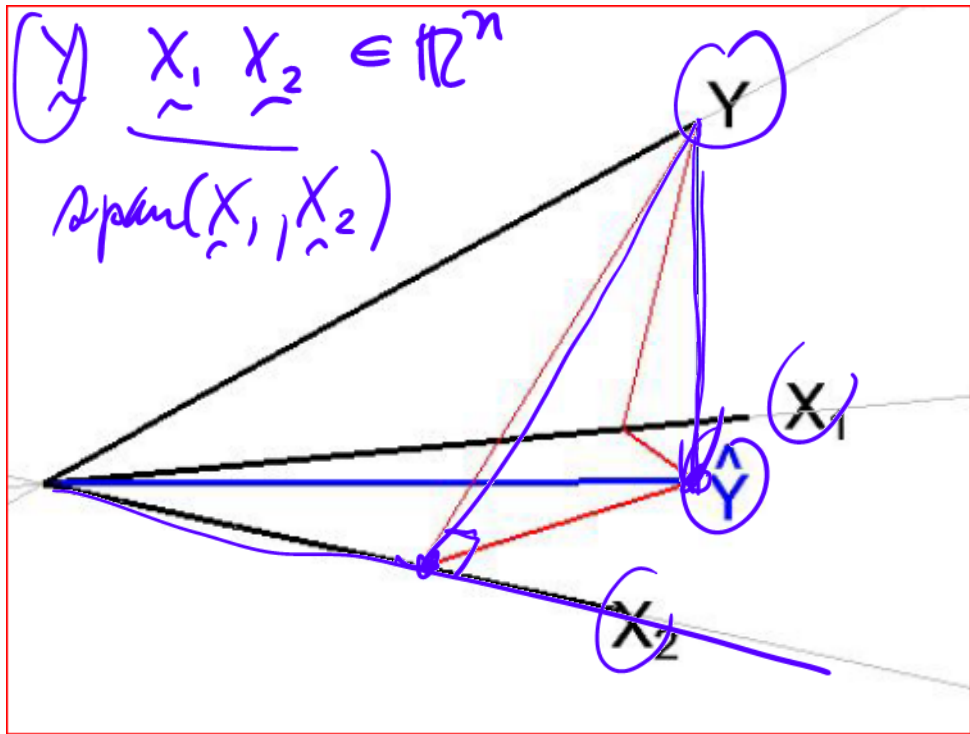
"Total effect"



Confounding

$$\textcircled{Y} \quad \underbrace{\tilde{X}_1, \tilde{X}_2}_{\in \mathbb{R}^n}$$

$$\text{span}(\tilde{X}_1, \tilde{X}_2)$$



This is where you determine the nature of the data: **observational or experimental**, and the nature of the questions: **predictive or causal** or descriptive.

2 Review of the matrix formulation and the general linear hypothesis (GLH)

$$Y = X\beta + \varepsilon$$

where

1. Y is a vector of length n representing n observations on a 'response' or 'dependent' variable,
2. X is a $n \times p$ matrix representing n observations on each of p 'predictor' or 'independent' variables. The first column frequently consists of 1's.
3. β is a vector of p parameters whose values are unknown and some aspect of which we wish to estimate. If the first column of X consists of 1's it is

customary to number the elements of β starting from 0:

$$\beta = (\beta_0, \beta_1, \dots, \beta_{p-1})'$$

4. ε is a vector of length n representing 'errors' or 'residuals' that are not directly observed.

If X is of full column rank (i.e. $\text{rank}(X) = p$) and if we assume that $\varepsilon \sim N_n(0, \sigma^2 I)$ where I is the $n \times n$ identity matrix, and if $\text{rank}(X) = p$, then the **UMVUE** (Uniformly minimum variance unbiased estimator) of β is

$$\hat{\beta} = (X'X)^{-1}X'Y$$

with $E(\hat{\beta}) = \beta$ and $\text{Var}(\hat{\beta}) = \sigma^2(X'X)^{-1}$

2.1 Linear hypotheses

We can estimate or test hypotheses concerning one or more linear combinations of the β s by forming a $h \times p$ hypothesis matrix L and estimating the function of parameters:

$$\eta = L\beta$$

$$\eta = \beta_2 - \beta_3$$

β_1
 β_2
 β_3

2.1.1 Exercise:

1. Why would we want to estimate a number of linear hypotheses simultaneously? Are the individual estimates of parameters different if we estimate them simultaneously? What difference does it make?

2.1.2 Example:

For a model with three parameters: $\beta = (\beta_0, \beta_1, \beta_2)'$ we can simultaneously estimate the sum and difference of β_1 and β_2 as follows.

Letting

$$L = \begin{bmatrix} 0 & 1 & 1 \\ 0 & 1 & -1 \end{bmatrix}$$

$$\beta_1 + \beta_2$$

$$\beta_1 - \beta_2$$

$$L\beta = \begin{bmatrix} 0 & 1 & 1 \\ 0 & 1 & -1 \end{bmatrix} \begin{bmatrix} \beta_0 \\ \beta_1 \\ \beta_2 \end{bmatrix}$$

we get

$$\eta = \begin{bmatrix} \eta_1 \\ \eta_2 \end{bmatrix} = L\beta$$

$$= \begin{bmatrix} 0 & 1 & 1 \\ 0 & 1 & -1 \end{bmatrix} \begin{bmatrix} \beta_0 \\ \beta_1 \\ \beta_2 \end{bmatrix}$$

$$= \begin{bmatrix} \beta_1 + \beta_2 \\ \beta_1 - \beta_2 \end{bmatrix}$$

$$\text{Var}(Y) = \Sigma$$

2.2 Estimation and tests

$$\text{Var}(AY) = A\Sigma A^T$$

Letting

$$\hat{\eta} = L\hat{\beta}$$

we have

$$E(\hat{\eta}) = E(L\hat{\beta}) = LE(\hat{\beta}) = L\beta$$

and

$$\text{Var}(\hat{\eta}) = \sigma^2 L(X'X)^{-1}L'$$

If L is of full row rank h and X is of full column rank we can test the hypothesis

$$H_0 : \eta = L\beta = 0$$

against the alternative that $\eta \neq 0$ (i.e. that $\eta_i \neq 0$ for at least one i) by using the null distribution:

$$\hat{\eta}' \left(\widehat{Var}(\hat{\eta}) \right)^{-1} \hat{\eta} = \frac{\hat{\beta}' L' (L(X'X)^{-1} L')^{-1} L \hat{\beta}}{s_e^2} \sim h \times F_{h, \nu}$$

where s_e is the 'residual standard error':

$$s_e^2 = \frac{\|Y - X\hat{\beta}\|^2}{\nu}$$

with $\nu = n - p$ the degrees of freedom for the estimate s_e^2 of σ^2 and $F_{h, \nu}$ is the F distribution with h and ν degrees of freedom.

2.2.1 Notes

1. If the rows of L are not linearly independent then it isn't possible to invert $L(X'X)^{-1}L'$ but an equivalent hypothesis can be formed by

replacing L with a matrix whose rows form a basis of the row space of L .

2.2.2 Exercises:

1. Two L matrices with the same row space test equivalent simultaneous hypotheses (**Could you prove this?**).
2. For example, the hypothesis above is equivalent to the hypothesis that $\beta_1 = \beta_2 = 0$. Why?

3 Interpreting Regression Coefficients: Smoking and Life Expectancy

With complex models:

1. most regression coefficients in the standard output are of little interest and
2. most interesting questions are not answered with the standard regression coefficients.

Why do we pay attention to regression output? Because it may make some

$$L_1 \quad L_2$$

$$(a_1 \ a_2 \ \dots \ a_n) \begin{bmatrix} L_2 \end{bmatrix} \quad h \text{ rows}$$

1st row

$$a_1 \tilde{r}_1 + a_2 \tilde{r}_2 + \dots + a_n \tilde{r}_n$$

$$L_1 = AL_2$$

sense for very simple additive models – but even then it is fraught with subtle traps most analysts do not understand

We will illustrate these with the Smoking and Life Expectancy example using country-level data in 2004.

```
dall <- read.csv(paste0("http://",server,"/data/Smoking3.csv"))
dd <- subset( dall, sex == 'BTSX') # subset of a data frame
dd$LifeExp <- dd$lifeexp.Birth # Life expectancy at birth
dd$LE <- dd$LifeExp
dd$smoke <- dd$consumption.cigPC # cigarette consumption
                                     # per adult per year
dd$HE <- dd$HealthExpPC.Tot.ppp # health expenditures per capita
                                     # in US$ PPP
dd$hiv <- dd$hiv_prev15_49 # prevalence of HIV in
                             # population 15 to 49
dd$special <- ifelse(
  dd$country %in% c('Angola','Sierra Leone',
                    'Equatorial Guinea'),
  1,
```

0) # indicator variable for 3 outlying countries

head(dd)

	country	iso3	region	HealthExpPC.Govt.exch	Health
1	Afghanistan	AFG	EMR	8.72	
5	Angola	AGO	AFR	114.61	
7	Albania	ALB	EUR	114.21	
10	Andorra	AND	EUR	2246.75	
15	United Arab Emirates	ARE	EMR	1219.89	
17	Argentina	ARG	AMR	540.82	
	HealthExpPC.Govt.ppp	HealthExpPC.Tot.exch	total	govt	pri
1	7.87	55.93	50.47	7.87	4
5	132.04	186.26	214.58	132.04	8
7	253.49	254.64	565.20	253.49	31
10	2257.23	3058.98	3073.26	2257.23	81
15	1288.52	1639.87	1732.13	1288.52	44
17	869.44	891.80	1433.70	869.44	56
	lifeexp.Birth	lifeexp.At60	smoking.tobacco.current	smoking.to	

	1	60	16	NA		
	5	51	16	NA		
	7	74	19	40		
	10	82	25	35		
	15	76	19	10		
	17	76	21	27		
		smoking.cig.current	smoking.cig.daily	Pop.Total	Pop.MedAge	Pop
	1	NA	NA	29825	16.20	
	5	NA	NA	20821	16.18	
	7	40	36	3162	32.56	
	10	35	31	78	NA	
	15	8	5	9206	29.37	
	17	26	20	41087	30.83	
		Pop.pCntOver60	Pop.pCntAnnGrowth	consumption.cigPC	hiv_prev15	
	1	3.82	-2.4	61		
	5	3.84	-3.1	414		
	7	14.93	-0.3	1116		
	10	22.86	0.0	784		
	15	0.81	-3.1	583		

	17		14.97		-0.9		1042
		smoke	HE	hiv	special		
	1	61	50.47	0.0	0		
	5	414	214.58	2.1	1		
	7	1116	565.20	NA	0		
	10	784	3073.26	NA	0		
	15	583	1732.13	NA	0		
	17	1042	1433.70	0.4	0		

```
fit.hiv2 <- lm( LifeExp ~
  log(HE) * (smoke + I(smoke^2)) + hiv+special,
  data = dd,
  na.action = na.exclude)
summary(fit.hiv2)
```

```
|
| Call:
| lm(formula = LifeExp ~ log(HE) * (smoke + I(smoke^2)) + hiv +
|     special, data = dd, na.action = na.exclude)
```

Residuals:

Min	1Q	Median	3Q	Max
-9.0373	-2.3005	0.2043	2.0760	9.7344

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)	
(Intercept)	3.283e+01	2.674e+00	12.280	< 2e-16	***
log(HE)	6.091e+00	5.024e-01	12.124	< 2e-16	***
smoke	3.642e-02	7.520e-03	4.844	3.31e-06	***
I(smoke^2)	-1.518e-05	3.946e-06	-3.846	0.000181	***
hiv	-7.351e-01	7.593e-02	-9.681	< 2e-16	***
special	-1.822e+01	2.137e+00	-8.526	2.11e-14	***
log(HE):smoke	-4.878e-03	1.155e-03	-4.223	4.30e-05	***
log(HE):I(smoke^2)	2.007e-06	5.726e-07	3.504	0.000614	***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 3.63 on 141 degrees of freedom

```
| (45 observations deleted due to missingness)
| Multiple R-squared: 0.8613, Adjusted R-squared: 0.8544
| F-statistic: 125 on 7 and 141 DF, p-value: < 2.2e-16
```

Do we need curvature in 'smoke'?

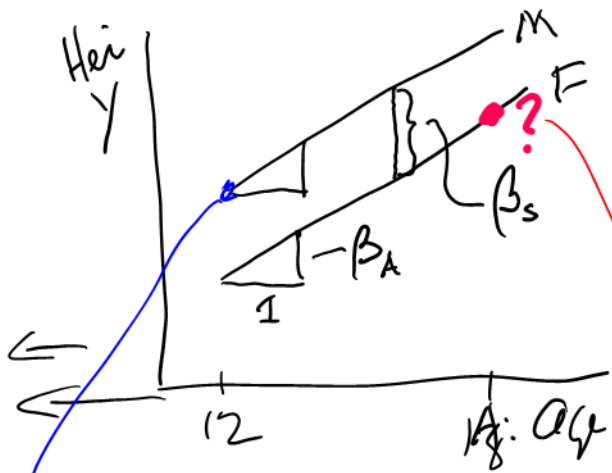
```
wald(fit.hiv2)
```

```
| numDF denDF F.value p.value
|      8  141 6968.992 <.00001
|
| Estimate Std.Error DF t-value p-value Lo
| (Intercept) 32.831342 2.673604 141 12.279805 <.00001 2
| log(HE) 6.091328 0.502427 141 12.123808 <.00001
| smoke 0.036424 0.007520 141 4.843564 <.00001
| I(smoke^2) -0.000015 0.000004 141 -3.846440 0.00018
| hiv -0.735098 0.075932 141 -9.681048 <.00001 -
| special -18.223307 2.137312 141 -8.526276 <.00001 -2
| log(HE):smoke -0.004878 0.001155 141 -4.223209 0.00004 -
| log(HE):I(smoke^2) 0.000002 0.000001 141 3.504458 0.00061
|
| Upper 0.95
```

```
| (Intercept)          38.116875
| log(HE)              7.084592
| smoke                0.051290
| I(smoke^2)           -0.000007
| hiv                  -0.584986
| special              -13.997989
| log(HE):smoke        -0.002595
| log(HE):I(smoke^2)  0.000003
```

```
wald(fit.hiv2, '2') # using '2' as a regular expression
```

```
|      numDF denDF  F.value p.value
|      2      2   141 8.835643 0.00024
|
|              Estimate Std.Error DF  t-value  p-value Lowe
| I(smoke^2)      -1.5e-05 4e-06   141 -3.846440 0.00018 -2.3
| log(HE):I(smoke^2) 2.0e-06 1e-06   141  3.504458 0.00061  1.0
|
|              Upper 0.95
| I(smoke^2)      -7e-06
| log(HE):I(smoke^2) 3e-06
```



$$S: \begin{cases} M \\ F \end{cases}$$

$$Y \sim A + S$$

$$E(Y) = \hat{\beta}_0 + \hat{\beta}_A A + \hat{\beta}_S S$$

$\frac{\partial F}{\partial M}$
 $\frac{\partial F}{\partial M}$

$$\beta_0 + \beta_A 12 + \beta_S 1$$

$$\begin{bmatrix} 1 & 12 & 1 \end{bmatrix} \begin{bmatrix} \beta_0 \\ \beta_A \\ \beta_S \end{bmatrix}$$

$$\beta_0 + \beta_A 18 + \beta_S 0$$

$$= \beta_0 + \beta_A 18$$

$$= \begin{bmatrix} 1 & 18 & 0 \end{bmatrix} \begin{bmatrix} \beta_0 \\ \beta_A \\ \beta_S \end{bmatrix}$$

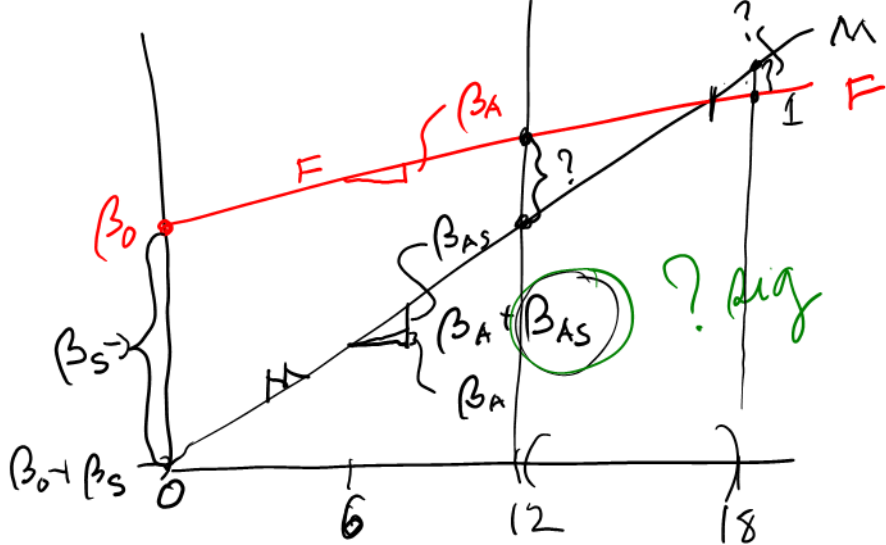
$$L = \begin{bmatrix} 1 & 18 & 0 \\ 1 & 12 & 1 \end{bmatrix} \quad \text{Est } L\beta \text{ with } L\hat{\beta}$$

$$Y \sim A * S \quad E(Y) = \beta_0 + \beta_A A + \beta_S S + \beta_{AS} A \cdot S$$

$$\underline{S = F} \quad E(Y) = \beta_0 + \beta_A A + \underbrace{\beta_S 0 + \beta_{AS} A \cdot 0}_{\text{line}}$$

$$\underline{S = M} \quad E(Y) = \beta_0 + \beta_A A + \beta_S \cdot 1 + \beta_{AS} \cdot A \cdot 1 \\ = \underbrace{(\beta_0 + \beta_S)}_{\text{line}} + \underbrace{(\beta_A + \beta_{AS})}_{\text{line}} \cdot A$$





$$\begin{aligned}
 \text{gap at } A_0 : \frac{\partial E(Y)}{\partial S} \Big|_{A=A_0} &= \frac{\partial}{\partial S} (\beta_0 + \beta_A A + \beta_S S + \beta_{AS} \cdot A \cdot S) \\
 &= (0 + 0 + \beta_S + \beta_{AS} A)
 \end{aligned}$$

$$= \begin{bmatrix} 0 & 0 & 1 & A \end{bmatrix} \begin{bmatrix} \beta_0 \\ \beta_A \\ \beta_S \\ \beta_{KS} \end{bmatrix}$$

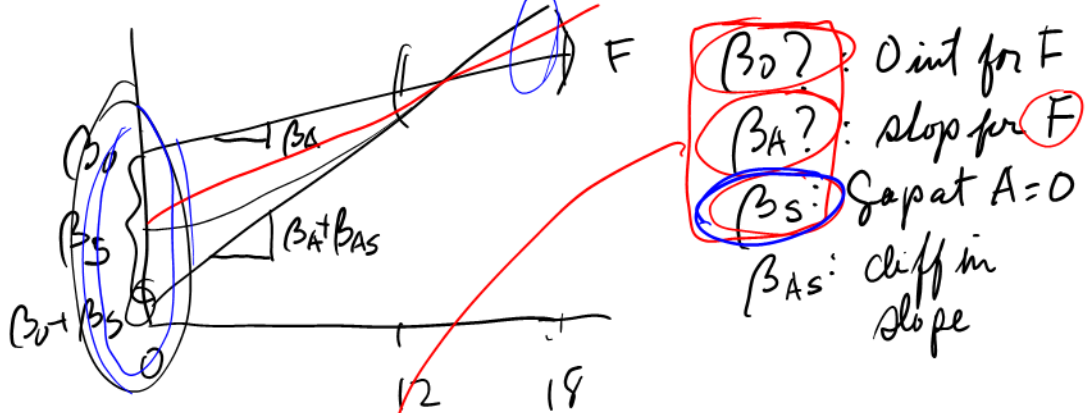
To estimate gap at age 12 & 18

$$L = \begin{bmatrix} 0 & 0 & 1 & 12 \\ 0 & 0 & 1 & 18 \end{bmatrix}$$

L is est of $L\beta$ which is the 2 gaps.

Interpretation of β_S

β_0 β_A β_S β_{KS}



A in AS

S in AS

Int in all terms

Principle of Marginality: Caution interpreting any term that is also in a higher order interaction.

$$Y \sim \underline{A * B * C}$$

$$E(Y) = \beta_0 + \underline{\beta_A} A + \underline{\beta_B} B + \underline{\beta_C} C + \underline{\beta_{AB}} AB \\ + \underline{\beta_{AC}} AC + \underline{\beta_{BC}} BC + \underline{\beta_{ABC}} ABC$$

have conditional interpretations of

Need to ask relevant Q's

How about interaction?

```
wald(fit.hiv2, ':')
```

```
|      numDF denDF F.value p.value
|      :      2    141  9.2372 0.00017
|
|      Estimate Std.Error DF  t-value  p-value Low
| log(HE):smoke -0.004878 0.001155 141 -4.223209 0.00004 -0.
| log(HE):I(smoke^2) 0.000002 0.000001 141 3.504458 0.00061 0.
|
|      Upper 0.95
| log(HE):smoke -0.002595
| log(HE):I(smoke^2) 0.000003
```

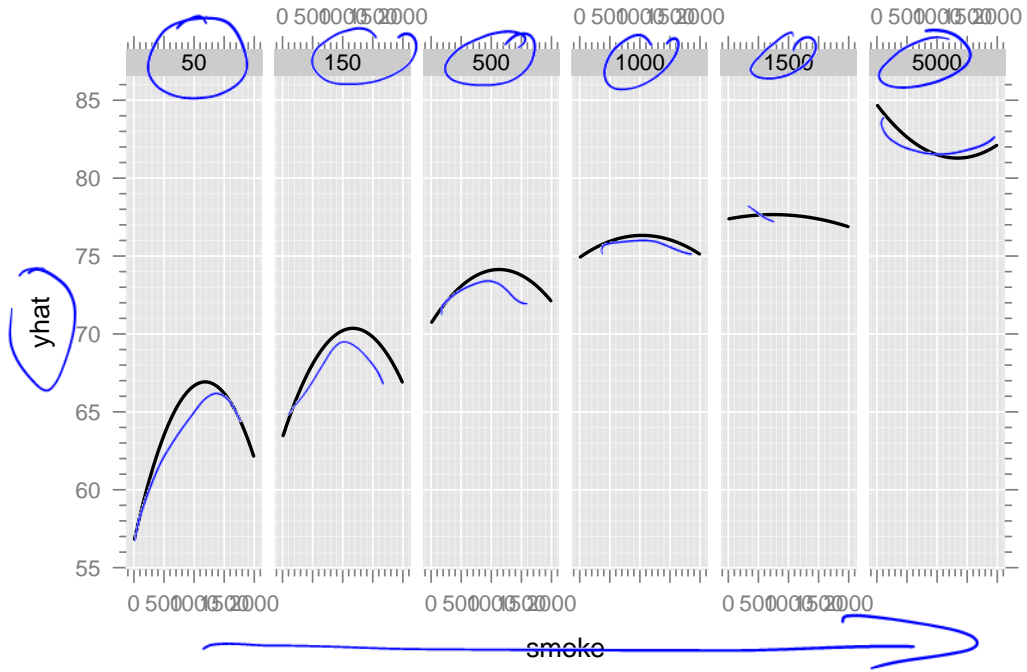
Create a prediction data frame over which to estimate fitted values. We will look at countries with low hiv and exclude outliers.

```
pred <- expand_grid(
  HE = c(50, 150, 500, 1000, 1500, 5000),
  smoke = seq(10, 2000, 20),
  hiv = 0,
```

```
special = 0)
```

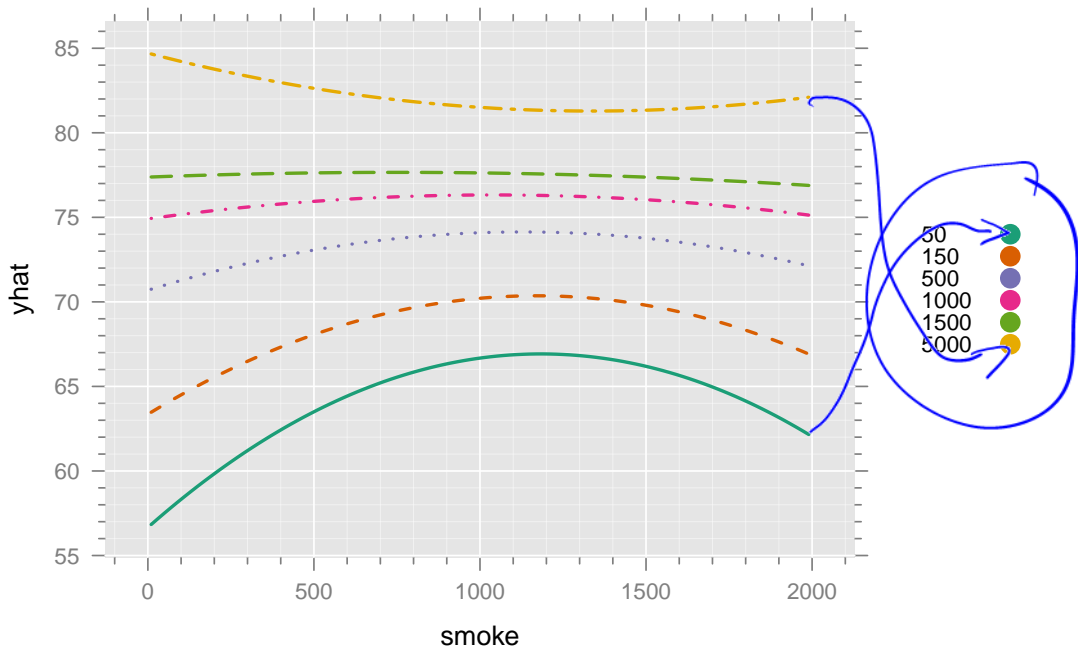
Finding \hat{Y} over a grid of values:

```
pred$yhat <- predict(fit.hiv2, newdata = pred)
gd(lwd = 2) # no groups
gd(lwd = c(2,2)) # groups
xyplot(yhat ~ (smoke | factor(HE)), pred, type = 'l',
       layout = c(6,1))
```



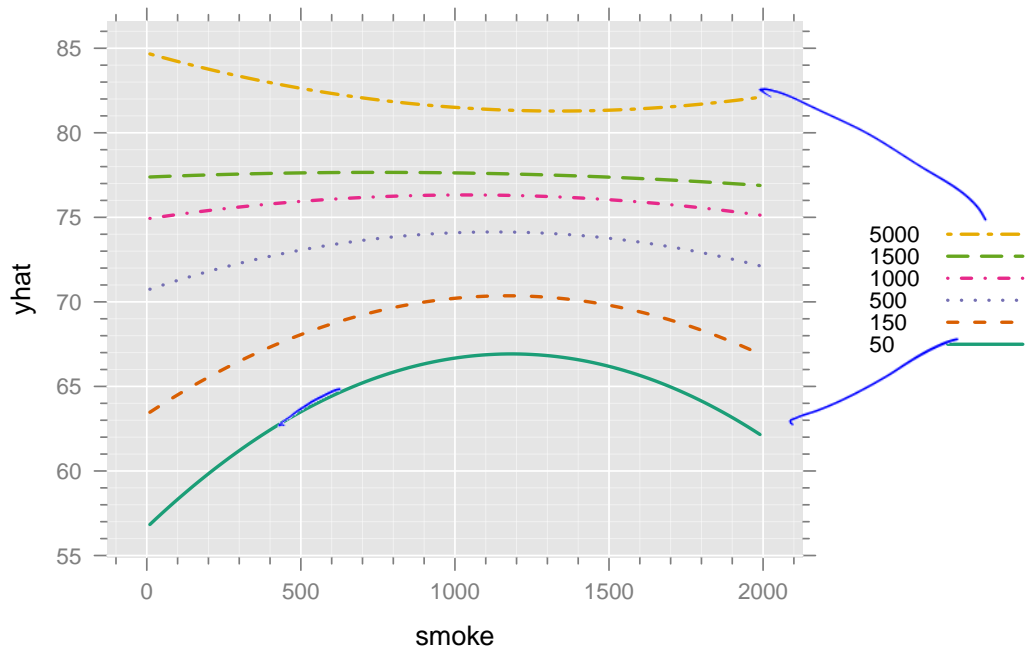
```
xyplot(yhat ~ smoke, groups = factor(HE), pred, type = 'l',  
       auto.key=list(space='right'))
```





It's a good idea to make the order in the legend match the physical location in the graph, as much as feasible.

```
xyplot(yhat ~ smoke , groups = factor(HE), pred, type = 'l',  
       auto.key=list(space='right', lines = T, points = F,  
                     reverse.rows = TRUE))
```



Suppose we want to estimate the slope of these fitted curves: i.e. the 'effect' of an additional cigarette as a function of health expenditures and amount smoked.

We start with the mathematical formula for the model:

Letting $\eta = E(y|HE, Smoke, HIV, Special)$

$$\eta = \beta_0 + \beta_1 \times \ln(HE)$$

$$+ \beta_2 \times Smoke$$

$$+ \beta_3 \times Smoke^2$$

$$+ \beta_4 \times HIV$$

$$+ \beta_5 \times Special$$

$$+ \beta_6 \times \ln(HE) \times Smoke$$

$$+ \beta_7 \times \ln(HE) \times Smoke^2$$

$$\beta_2 \times 1$$

$$\beta_3 \times 2 \times Smoke$$

$$\beta_6 \times \ln(HE)$$

$$\beta_7 \times 2 \times \ln(HE) \times Smoke$$

To understand the interpretation of the coefficients β_i , we differentiate η with respect to each of the independent variables:

$$\frac{\partial \eta}{\partial HE} = \beta_1 \frac{1}{HE} + \beta_6 \frac{Smoke}{HE} + \beta_7 \frac{Smoke^2}{HE}$$

$$\frac{\partial \eta}{\partial Smoke} = \beta_2 + 2\beta_3 Smoke + \beta_6 \ln(HE) + 2\beta_7 Smoke \ln(HE)$$

$$\frac{\partial \eta}{\partial HIV} = \beta_4$$

$$\frac{\partial \eta}{\partial Special} = \beta_5$$

$$\frac{\partial^2 \eta}{\partial HE^2} = \beta_1 \frac{-1}{HE^2} + \beta_6 \frac{-Smoke}{HE^2} + \beta_7 \frac{-Smoke^2}{HE^2}$$

$$\frac{\partial^2 \eta}{\partial Smoke^2} = 2\beta_3$$

$$\frac{\partial^2 \eta}{\partial HE \partial Smoke} = \beta_6 \frac{1}{HE} + 2\beta_7 \frac{Smoke}{HE}$$

Thus β_2 is the **partial derivative** of η with respect to *Smoke* when $\ln(HE) = Smoke = 0$.

When $\ln(HE) = 5$ and *Smoke* = 4, the partial derivative of η with respect to

Smoke is

$$\frac{\partial \eta}{\partial \text{Smoke}} = \beta_2 + 8\beta_3 + 5\beta_6 + 40\beta_7$$

whose estimator is

$$\hat{\beta}_2 + 8\hat{\beta}_3 + 5\hat{\beta}_6 + 40\hat{\beta}_7$$

which we can express as a linear transformation of the $\hat{\beta}$ vector. Letting

$$L = \begin{bmatrix} 0 & 0 & 1 & 8 & 0 & 0 & 5 & 40 \end{bmatrix}$$

we have:

$$\hat{\phi} = L\hat{\beta} = \begin{bmatrix} 0 & 0 & 1 & 8 & 0 & 0 & 5 & 40 \end{bmatrix}$$

 β_0 β_1 β_2 β_3 β_4 β_5 β_6 β_7

If we wish to simultaneously estimate the 'effect' of Smoke and the 'effect' of HE given values of HE and Smoke, we can form the L matrix:

$$L = \begin{bmatrix} 0 & 0 & 1 & 2 \text{ Smoke} & 0 & 0 & n(HE) & 2 \text{ Smoke} n(HE) \\ 0 & 1 & 0 & 0 & 0 & 0 & \frac{\text{Smoke}}{HE} & \frac{\text{Smoke}^2}{HE} \end{bmatrix}$$

and

$$\hat{\phi} = L\hat{\beta}$$

In this case $\hat{\phi}$ is a column vector of length 2.

In both cases, inference about ϕ uses the fact that

$$\text{Var}(\hat{\phi}) = L \text{Var}(\hat{\beta}) L'$$

and

$$\text{Var}(\hat{\beta}) = \sigma^2 (X'X)^{-1}$$

With a normal linear model in which

$$Y = X\beta + \epsilon, \epsilon \sim N(0, \sigma^2 I)$$

$L = \begin{bmatrix} 0 & 1 & -1 & 0 \\ 0 & 1 & 0 & -1 \\ 0 & 0 & 1 & -1 \end{bmatrix}$

$\beta_1 = \beta_2$
 $\beta_1 = \beta_3$
 $\beta_2 = \beta_3$

we have that

$$(\hat{\phi} - \phi)' (s^2 L(X'X)^{-1} L')^{-1} (\hat{\phi} - \phi) \sim h \times F_{h,\nu}$$

exists only if L is of full row rank

where h is the number of rows of L (assuming that L is of full row rank) and $\nu = n - p$ where n and p are the number of rows and columns of X respectively, again assuming that X is of full column rank.

rank

We can compute these quantities in R from a fitted model. Note how the 'evalq' function evaluates an expression at the values given in the list provided as the 'envir' argument.

```
L <- evalq(rbind(
  c(0,0,1, 2*smoke, 0, 0, log(HE), 2*smoke*log(HE)),
  c(0,1,0, 0, 0, 0, smoke / HE, smoke^2/HE)),
  envir = list(smoke = 4, HE = exp(5)))
```

```
L
  [,1] [,2] [,3] [,4] [,5] [,6] [,7] [,8]
[1,]  0    0    1    8    0    0 5.00000000 40.0000000
[2,]  0    1    0    0    0    0 0.02695179 0.1078072
```


$\hat{\beta}$:

```
coef(fit.hiv2)
```

```
| (Intercept)          log(HE)          smoke  
| 3.283134e+01        6.091328e+00        3.642387e-02  
|          hiv          special      log(HE):smoke  log(HE)  
| -7.350980e-01      -1.822331e+01      -4.878054e-03        2
```

$\hat{\phi} = L\hat{\beta}$:

```
(phihat <- L %*% coef(fit.hiv2))
```

```
|           [,1]  
| [1,] 0.01199246  
| [2,] 6.09119673
```

$s^2(X'X)^{-1}$:

```
vcov(fit.hiv2)
```

```
|           (Intercept)          log(HE)          smoke
```

	(Intercept)	7.148161e+00	-1.298157e+00	-1.437171e-02	5.
	log(HE)	-1.298157e+00	2.524328e-01	2.379512e-03	-8.
	smoke	-1.437171e-02	2.379512e-03	5.655121e-05	-2.
	I(smoke^2)	5.526193e-06	-8.798735e-07	-2.706223e-08	1.
	hiv	-9.426054e-03	-2.319091e-03	1.603252e-05	-6.
	special	4.335683e-01	-1.122056e-01	-1.123946e-03	4.
	log(HE):smoke	2.438561e-03	-4.363636e-04	-8.441015e-06	3.
	log(HE):I(smoke^2)	-9.000438e-07	1.540858e-07	3.944264e-09	-2.
		hiv	special	log(HE):smoke	log
	(Intercept)	-9.426054e-03	4.335683e-01	2.438561e-03	
	log(HE)	-2.319091e-03	-1.122056e-01	-4.363636e-04	
	smoke	1.603252e-05	-1.123946e-03	-8.441015e-06	
	I(smoke^2)	-6.423962e-09	4.625775e-07	3.949943e-09	
	hiv	5.765616e-03	-1.218220e-03	2.000199e-06	
	special	-1.218220e-03	4.568101e+00	2.166940e-04	
	log(HE):smoke	2.000199e-06	2.166940e-04	1.334160e-06	
	log(HE):I(smoke^2)	-2.425728e-10	-8.002814e-08	-6.010943e-10	

$$\hat{Var}(\hat{\phi}) = L(s^2(X'X)^{-1})L'$$

```
(Vphihat <- L %*% vcov(fit.hiv2) %*% t(L))
```

```
|           [,1]           [,2]  
| [1,] 5.453266e-06 0.0001967712  
| [2,] 1.967712e-04 0.2524093523
```

To test the hypothesis that $\phi = 0$, we have

$$F = \hat{\phi}' (V\hat{ar}(\hat{\phi}))^{-1} \hat{\phi}/h$$

```
(Ftest <- (t(phihat) %*% solve(Vphihat) %*% phihat)/2)
```

```
|           [,1]  
| [1,] 78.44759
```

```
1-pf(Ftest,2, fit.hiv2$df.residual)
```

```
|           [,1]  
| [1,] 0
```

$p\text{val} = 1 - ?$


78.4

```
pf(Ftest,2, fit.hiv2$df.residual, lower.tail = FALSE)
```

```
| [1,]
| [1,] 1.254508e-23
```

Note how rounding error is reduced by using the 'lower.tail' parameter.

3.1 Functions to test linear hypotheses

The functions 'lht' in the 'car' package and 'wald' in the 'spida2' package can be used to test General Linear Hypotheses.

```
require(car)
lht(fit.hiv2,L)
```

```
| Linear hypothesis test
```

```
| Hypothesis:
```

$$L\beta = c$$

```
smoke + 8 I(smoke^2) + 5 log(HE):smoke + 40 log(HE):I(smoke^2)
log(HE) + 0.0269517879963419 log(HE):smoke + 0.107807151985367
```

Model 1: restricted model

Model 2: LifeExp ~ log(HE) * (smoke + I(smoke^2)) + hiv + special

	Res.Df	RSS	Df	Sum of Sq	F	Pr(>F)
1	143	3925.4				
2	141	1858.0	2	2067.4	78.448	< 2.2e-16 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

wald(fit.hiv2, L)

	numDF	denDF	F.value	p.value
1	2	141	78.44759	<.00001

	Estimate	Std.Error	DF	t-value	p-value	Lower 0.95	Upper 0.95
[1,]	0.011992	0.002335	141	5.155465	<.00001	0.007376	0.01660
[2,]	6.091197	0.502404	141	12.124111	<.00001	5.097979	7.08441

$L\beta = 0$

The 'lht' function can take a right-hand side to test hypotheses of the form $H_0 : \phi = \phi_0$. 'wald' can only test $H_0 : \phi = 0$. The 'wald' function can handle L matrices whose rows are not linearly independent. For example, in some uses of 'wald', the 'L' matrix is the whole design matrix.

The second argument of the 'wald' matrix can be a regular expression that is matched against the names of terms in the model. All terms matched by the regular expression are simultaneously tested. Thus one can test the 'overall' significance of an independent variable by testing whether all terms containing that variable are equal to 0. One can also use this approach to test higher-order interactions.

```
wald(fit.hiv2, "smoke") # is there statistical evidence
```

```
|      numDF denDF  F.value p.value  
| smoke      4   141  8.024538  1e-05  
|  
|      Estimate Std.Error DF  t-value p-value Low  
| smoke      0.036424 0.007520  141  4.843564 0.00001 0.  
| f(smoke^2) -0.000015 0.000004  141 -3.846440 0.00018 -0.  
| log(HE):smoke -0.004878 0.001155  141 -4.223209 0.00004 -0.
```

```

| log(HE):I(smoke^2) 0.000002 0.000001 141 3.504458 0.00061 0.
|
| Upper 0.95
| smoke 0.051290
| I(smoke^2) -0.000007
| log(HE):smoke -0.002595
| log(HE):I(smoke^2) 0.000003

```

```

wald(fit.hiv2, "HE") # that 'smoke' improves prediction?
                    # ditto for HE

```

```

| numDF denDF F.value p.value
| HE 3 141 87.10337 <.00001
|
| Estimate Std.Error DF t-value p-value Lower Upper
| log(HE) 6.091328 0.502427 141 12.123808 <.00001 5.086473 7.096183
| log(HE):smoke -0.004878 0.001155 141 -4.223209 0.00004 -0.007191 -0.002565
| log(HE):I(smoke^2) 0.000002 0.000001 141 3.504458 0.00061 0.000000 0.000003
|
| Upper 0.95
| log(HE) 7.084592
| log(HE):smoke -0.002595

```

```
| log(HE):I(smoke^2) 0.000003
```

```
wald(fit.hiv2, ":") # ditto for interactions?
```

```
| numDF denDF F.value p.value
```

```
| : 2 141 9.2372 0.00017
```

```
| Estimate Std.Error DF t-value p-value Low
```

```
| log(HE):smoke -0.004878 0.001155 141 -4.223209 0.00004 -0.
```

```
| log(HE):I(smoke^2) 0.000002 0.000001 141 3.504458 0.00061 0.
```

```
| Upper 0.95
```

```
| log(HE):smoke -0.002595
```

```
| log(HE):I(smoke^2) 0.000003
```

```
wald(fit.hiv2, "2") # ditto for quadratic terms?
```

```
| numDF denDF F.value p.value
```

```
| 2) 2 141 8.835643 0.00024
```

```
| Estimate Std.Error DF t-value p-value Lowe
```

```
| I(smoke^2) -1.5e-05 4e-06 141 -3.846440 0.00018 -2.3
```

```
| log(HE):I(smoke^2) 2.0e-06 1e-06 141 3.504458 0.00061 1.0
```


		Upper 0.95
	I(smoke^2)	-7e-06
	log(HE):I(smoke^2)	3e-06

3.2 Strategies to simplify models

There are many strategies for potentially simplifying large models. The resulting model will depend on the strategy.

One is to attack higher-order interactions and simplify the model by dropping groups of interactions that are not significant but, initially, leaving main effects and lower-order interactions. In many situations there are obvious moderator variables whose interactions should not be dropped as aggressively as those of other variables for which oversimplification to an additive model may be more innocuous. Remember the consequences of dropping an interaction. Main effects become weighted averages of conditional effects, weighted by inverse variance.

Another approach is to drop all terms for selected independent variables if they are not sufficiently significant in an overall test.

Lasso

The two approaches can be combined depending on the role of variables and the goals of the analysis.

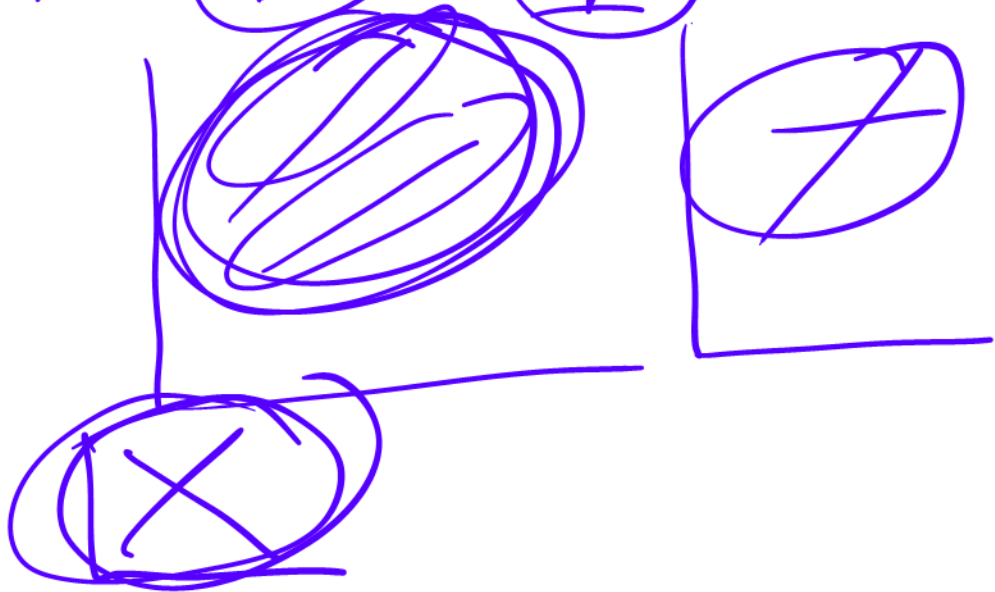
The choice of approach should be guided by many factors: which null hypotheses are likely to be reasonable, the interpretive value of having a simple additive model versus the added validity of estimating conditional effects that are not averaged over levels of variables that may be important, etc. There's a good discussion of these problems in (???)

3.3 Estimating effects over a grid

In a model with interactions and non-linear functions of some independent variables, it is often interesting to characterize how effects (partial derivatives) and inferences about effects vary over a range of predictors.

The 'effects' package (Fox and Hong (2009)) allows easy visualization of predicted values as a variables changes keeping other variables constant. For other variables that interact with the variable whose effect is visualized, the interacting variables are kept constant at a number of selected values. The package is very effective for the easy visualization of moderately complex

$$y \sim \text{Sex} * \text{Age}$$



models with interactions.

The same can be achieved but much more laboriously with the 'Lfx' function in the 'spida2' package (Monette et al. (2018)). With the 'Lfx' function it is possible to estimate derivatives of all orders and features of general parametric splines with the 'gsp' function.

The 'Lfx' function generates an expression which can then be edited to generate large L matrices. The result of the wald test applied to this L matrix can be transformed into a data frame for plotting with error bands.

The following illustrates the use of the 'Lfx' function.

```
Lfx(fit.hiv2)
```

```
| list(1,  
| 1 * M(log(HE)),  
| 1 * smoke,  
| 1 * M(I(smoke^2)),  
| 1 * hiv,  
| 1 * special,
```

$$2LE = k$$

~~2 smoke~~ ~~var int~~

~~2LE~~ vary on other factors
~~2 smoke~~

```
| 1 * M(log(HE)) * smoke,  
| 1 * M(log(HE)) * M(I(smoke^2))  
| )
```

$I = \text{identity}$

The expression generated by 'Lfx' can be edited to generate desired effects. The result is then fed back to 'Lfx' along with a data frame on which to evaluate the edited expression. Note that the 'M' functions preserved the shape of multi-term blocks in the design matrix so that multiplying them by 0 is a way of generating a block of 0s of the right dimension. In the following, we edit the expression to estimate the effect of smoking by differentiating with respect to 'smoke':

```
pred <- expand.grid(  
  HE = c(50, 150, 500, 1000, 1500, 5000),  
  smoke = seq(10, 2000, 20),  
  hiv = 0,  
  special = 0`  
head(pred) # first 6 lines of 'pred'
```

```
| HE smoke hiv special  
| 1 50 10 0 0
```



	2	150	10	0	0
	3	500	10	0	0
	4	1000	10	0	0
	5	1500	10	0	0
	6	5000	10	0	0

Predicted values as a function of HE and smoke

Use the list created above and edit it by differentiating each term with respect to 'smoke' to get the marginal 'effect' of an extra unit of 'smoke':

`Lfx(fit.hiv2)`

```

| list(1,
| 1 * M(log(HE)),
| 1 * smoke,
| 1 * M(I(smoke^2)),
| 1 * hiv,
| 1 * special,
| 1 * M(log(HE)) * smoke,

```

Handwritten annotations in blue ink: A large bracket on the right side of the code block. Circles around the terms '1 * smoke', '1 * M(I(smoke^2))', '1 * M(log(HE)) * smoke', and the '1' in the first line. A circle around 'smoke' in the last line with a diagonal slash through it. A '0' above the first line, a '-0' to the right of the second line, and a '2' to the right of the third line.

```
| 1 * M(log(HE)) * M(I(smoke^2))  
| )
```

Differentiated:

```
L <- Lfx(fit.hiv2,  
list(0,  
0 * M(log(HE)),  
1,  
1 * M(I(2*smoke)),  
0 * hiv,  
0 * special,  
1 * M(log(HE)) * 1,  
1 * M(log(HE)) * M(I(2*smoke))  
), prec)  
dim(L)
```

```
| [1] 600 8
```

~~1111~~

1 + 0 * ~~any~~



head(L) \

	(Intercept)	log(HE)	smoke	I(smoke^2)	hiv	special	log(HE):smoke
1	0	0	1	20	0	0	3.912023
2	0	0	1	20	0	0	5.010635
3	0	0	1	20	0	0	6.214608
4	0	0	1	20	0	0	6.907755
5	0	0	1	20	0	0	7.313220
6	0	0	1	20	0	0	8.517193
	log(HE):I(smoke^2)						
1	78.24046						
2	100.21271						
3	124.29216						
4	138.15511						
5	146.26441						
6	170.34386						


```
ww <- wald(fit.hiv2, L)
```

```
| Warning in wald(fit.hiv2, L): Poorly conditioned L matrix, calcu  
| be incorrect
```

```
ww <- as.data.frame(ww)
```

```
head(ww)
```

		coef	se	U2	L2	HE	smoke
	1	0.0171942856	0.003272967	0.023740220	0.010648352	50	10
	2	0.0118792893	0.002313918	0.016507124	0.007251454	150	10
	3	0.0060545675	0.001764820	0.009584207	0.002524927	500	10
	4	0.0027011781	0.001883655	0.006468487	-0.001066131	1000	10
	5	0.0007395711	0.002094140	0.004927851	-0.003448709	1500	10
	6	-0.0050851507	0.003067648	0.001050145	-0.011220447	5000	10

```
More informative labels:
```

```
ww$HEfac <- factor(ww$HE)
```

```
levels(ww$HEfac) <- paste("Health Exp/Cap:", levels(ww$HEfac))
```

Doing this preserved the order of factor levels

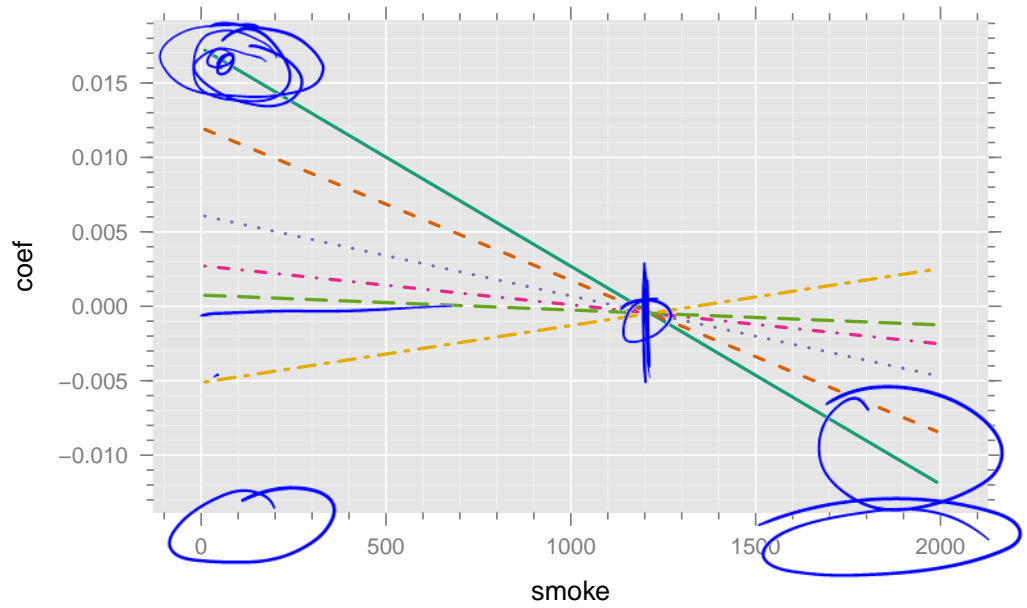
```
tab(ww, ~ HEfac + HE)
```

	HE							
HEfac	50	150	500	1000	1500	5000	Total	
Health Exp/Cap: 50	100	0	0	0	0	0	100	
Health Exp/Cap: 150	0	100	0	0	0	0	100	
Health Exp/Cap: 500	0	0	100	0	0	0	100	
Health Exp/Cap: 1000	0	0	0	100	0	0	100	
Health Exp/Cap: 1500	0	0	0	0	100	0	100	
Health Exp/Cap: 5000	0	0	0	0	0	100	100	
Total	100	100	100	100	100	100	600	

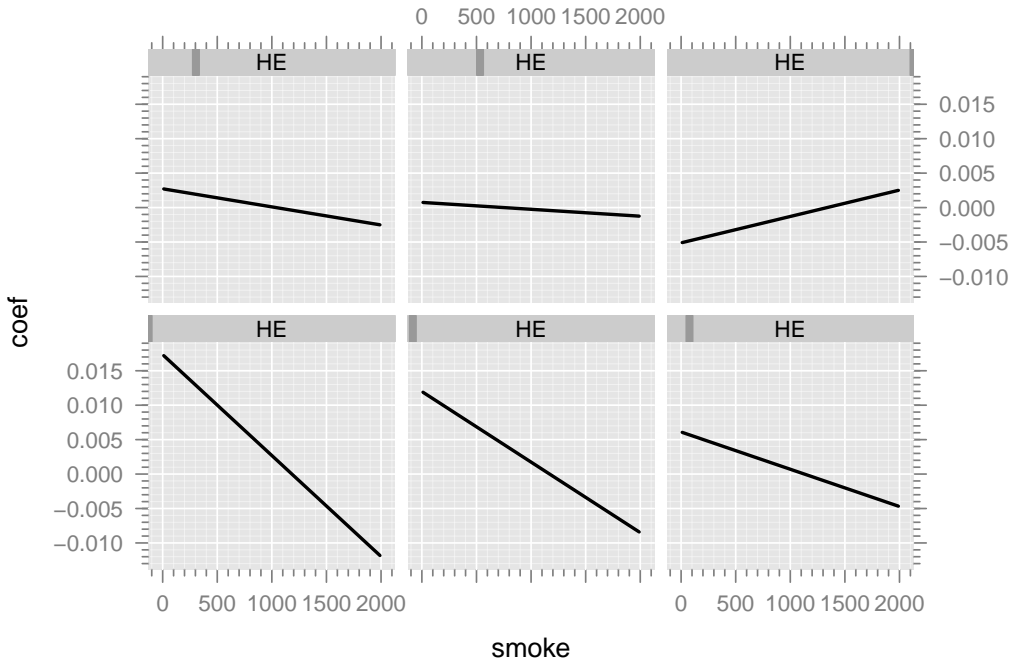
Levels are in numerical order when creating a factor from a numeric object!
This was not always so! We are taking the derivative with respect to 'smoke' of these functions:

```
xyplot(coef ~ smoke, ww, groups = HE,  
       auto.key = list(columns = 3, lines = T,
```

```
points = F),type = 'l')
```



```
xyplot( coef ~ smoke | HE, ww, type = 'l' )
```



```
xyplot( coef ~ smoke | HEfac, ww, type = 'l' )
```

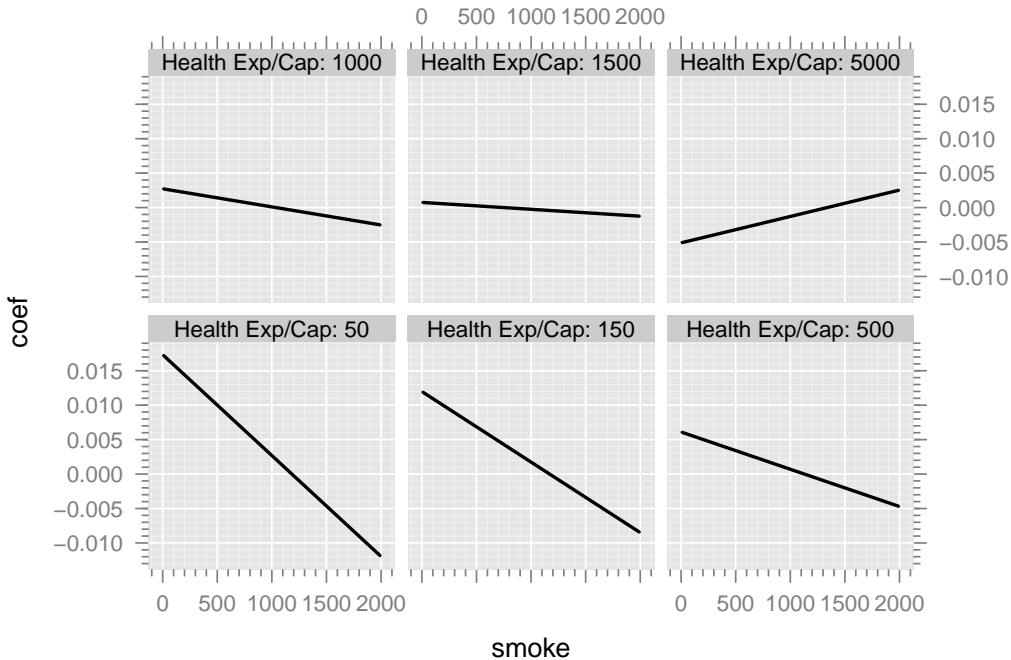
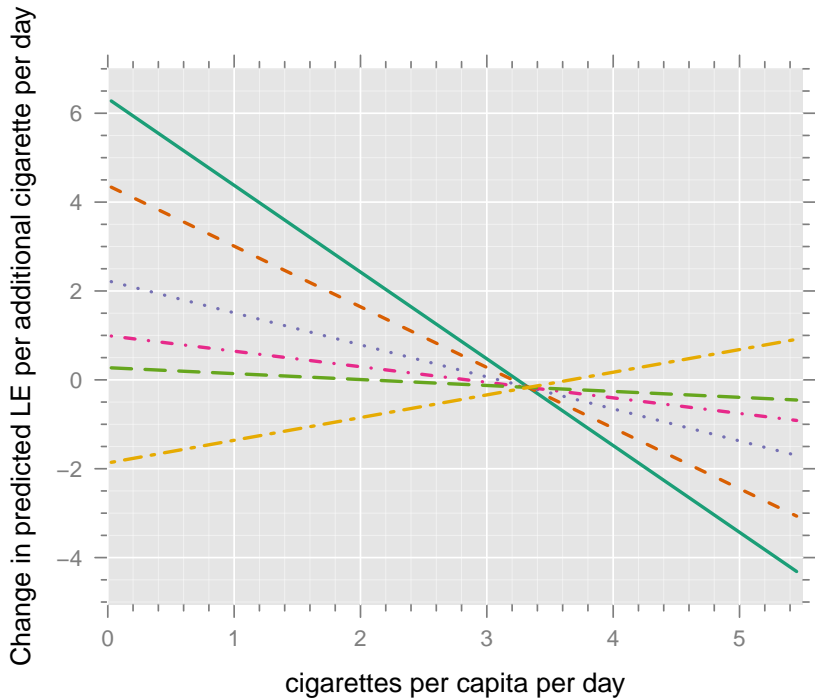


Figure 1: Change in Life Expectancy associated with an increase in cigarette consumption of 1 cigarette per day per capita for different levels of health expenditures per capita per year (US\$).

With labels that are more informative:



Health Exp. / Cap.

50

150

500

1000

1500

5000

Confidence bands:

lattice Extra

```
xyplot( J(365*coef) ~ I(smoke/365) | HEfac,
```

```
data = ww, type = 'n',
```

```
xlim = c(0,5.5),
```

```
ylab = "Change in predicted LE per additional cigarette",
```

```
xlab = "cigarettes per capita per day",
```

```
fit = 365*(ww$coef),
```

```
lower = 365*(ww$coef - ww$se),
```

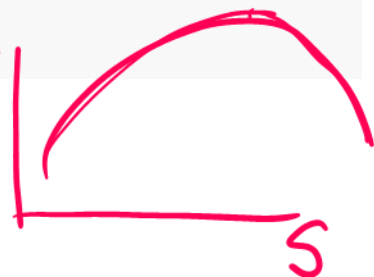
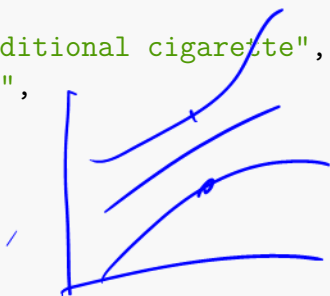
```
upper = 365*(ww$coef + ww$se),
```

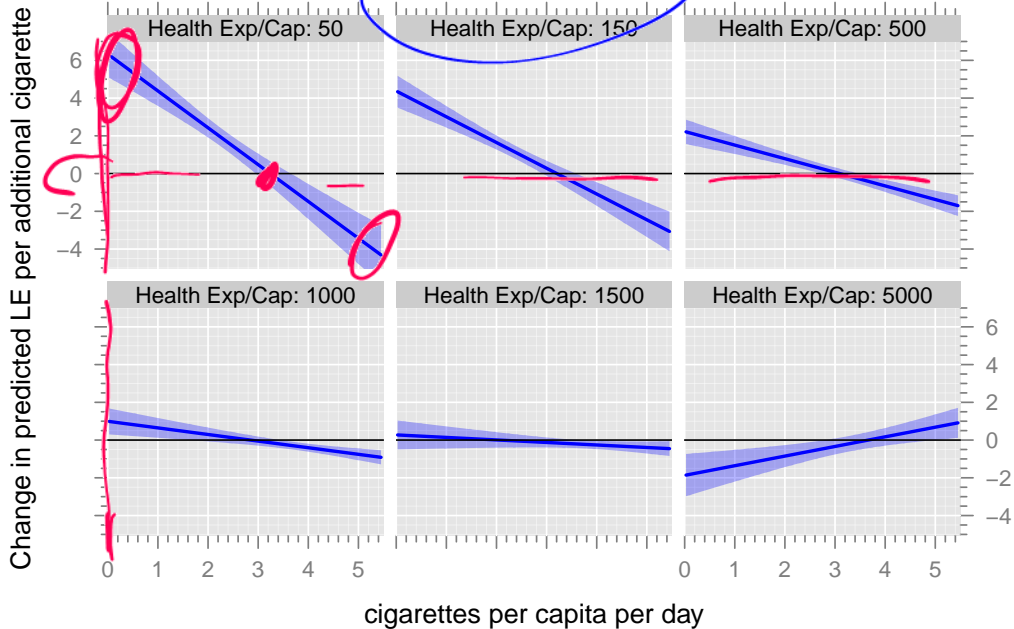
```
subscripts = T,
```

```
as.table = T) +
```

```
layer(panel.fit(...alpha = .3)) +
```

```
layer( panel.abline( h = 0, lwd = 1))
```





Transforming to 'meaningful' coefficients: cigarettes per day

```
## Double checking our previous calculation:
```

```
ptest <- expand.grid(smoke = 4, HE = exp(5), hiv = 0, special = 0)
```

```
(L2 <- Lfx( fit.hiv2,  
  list( 0,  
    0 * M(log(HE)),  
    1 ,  
    1 * M(I(2*smoke)),  
    0 * hiv,  
    0 * special,  
    1 * M(log(HE)) * 1,  
    1 * M(log(HE)) * M(I(2*smoke))  
  ), ptest))
```

```
| (Intercept) log(HE) smoke I(smoke^2) hiv special log(HE):smoke  
| 1           0       0       1           8     0         0           5  
| log(HE):I(smoke^2)
```

```
| 1 40
| attr(,"data")
| smoke HE hiv special
| 1 4 148.4132 0 0
```

```
wald(fit.hiv2, list("At smoke = 4, HE = 148.4"=L2))
```

```
| numDF denDF F.value p.value
| At smoke = 4, HE = 148.4 1 141 26.373 <.00001
| Estimate Std.Error DF t-value p-value Lower 0.95 Upper 0.95
| 1 0.011992 0.002335 141 5.135465 <.00001 0.007376 0.016609
```

3.3.1 Exercises

1. Carry out a similar process to estimate the 'effect' of health expenditures per capita.
2. Study the relative contribution of private versus public health expenditures on life expectancy.
3. Explore the 'effects' package and compare its functionality with 'Lfx'

3.4 Wald tests vs Likelihood Ratio Tests (LRT)

Let's consider a test for the need for a quadratic term in 'smoke'. There are two terms in the model that contain the quadratic term and a test to remove it involves more than one parameter. We need a test of

$$H_0 : \beta_3 = \beta_7 = 0$$

We cannot simply test each hypothesis $H_0 : \beta_3 = 0$ and $H_0 : \beta_7 = 0$ separately. We will see many examples where individual hypotheses are not significant, yet a joint hypothesis is highly significant. This is not a example of this phenomenon since the p-value for each hypothesis is small. Nevertheless, a test of a joint hypothesis needs to be carried out correctly. We consider two ways: a Wald test and a Likelihood Ratio Test executed with the 'anova' function, a clear misnomer.

Wald test using indices of coefficients

```
wald(fit.hiv2, c(4,8))
```

```
|      numDF denDF  F.value p.value
```

	1	2	141	8.835643	0.00024				
	Estimate	Std.Error	DF	t-value	p-value	Lower			
I(smoke^2)	-1.5e-05	4e-06	141	-3.846440	0.00018	-2.3			
log(HE):I(smoke^2)	2.0e-06	1e-06	141	3.504458	0.00061	1.0			
	Upper	0.95							
I(smoke^2)	-7e-06								
log(HE):I(smoke^2)	3e-06								

```
wald(fit.hiv2, list("Quadratic in smoke" = c(4,8)))
```

	numDF	denDF	F.value	p.value				
Quadratic in smoke	2	141	8.835643	0.00024				
	Estimate	Std.Error	DF	t-value	p-value	Lower		
I(smoke^2)	-1.5e-05	4e-06	141	-3.846440	0.00018	-2.3		
log(HE):I(smoke^2)	2.0e-06	1e-06	141	3.504458	0.00061	1.0		
	Upper	0.95						
I(smoke^2)	-7e-06							
log(HE):I(smoke^2)	3e-06							

Wald test using regular expression


```
wald(fit.hiv2, "2")
```

```
|      numDF denDF  F.value p.value  
|      2      2   141 8.835643 0.00024  
|                                     Estimate Std.Error DF  t-value  p-value Lowe  
| I(smoke^2) -1.5e-05 4e-06   141 -3.846440 0.00018 -2.3  
| log(HE):I(smoke^2) 2.0e-06 1e-06   141  3.504458 0.00061  1.0  
|                                     Upper 0.95  
| I(smoke^2) -7e-06  
| log(HE):I(smoke^2) 3e-06
```

Likelihood ratio test

We need to fit the 'null' model

```
fit0 <- update(fit.hiv2, log(HE)*smoke + hiv + special  
summary(fit0)
```

```
|  
| Call:
```

```
lm(formula = LifeExp ~ log(HE) + smoke + hiv + special + log(HE)
    data = dd, na.action = na.exclude)
```

```
Residuals:
```

```
      Min       1Q   Median       3Q      Max
-9.5894 -2.2654  0.0006  2.4404  9.6020
```

```
Coefficients:
```

```
                Estimate Std. Error t value Pr(>|t|)
(Intercept)    3.630e+01  2.082e+00  17.438 < 2e-16 ***
log(HE)         5.728e+00  3.613e-01  15.854 < 2e-16 ***
smoke           1.132e-02  3.149e-03   3.596 0.000444 ***
hiv             -7.617e-01  7.896e-02  -9.647 < 2e-16 ***
special        -1.802e+01  2.243e+00  -8.032 3.23e-13 ***
log(HE):smoke  -1.660e-03  4.628e-04  -3.586 0.000459 ***
```

```
---
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
Residual standard error: 3.824 on 143 degrees of freedom
```

```
| (45 observations deleted due to missingness)
| Multiple R-squared:  0.8439,    Adjusted R-squared:  0.8384
| F-statistic: 154.6 on 5 and 143 DF,  p-value: < 2.2e-16
```

Then compare the null model with the 'full' model:

By default, 'anova' uses an F distribution for the LRT taking advantage of the linear Gaussian model.

```
anova(fit0, fit.hiv2)
```

```
| Analysis of Variance Table
```

```
| Model 1: LifeExp ~ log(HE) + smoke + hiv + special + log(HE):smo
```

```
| Model 2: LifeExp ~ log(HE) * (smoke + I(smoke^2)) + hiv + specia
```

```
|   Res.Df    RSS Df Sum of Sq      F    Pr(>F)
```

```
| 1     143 2090.8
```

```
| 2     141 1858.0  2     232.86 8.8356 0.0002426 ***
```

```
| ---
```

```
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Using the general asymptotic distribution, chi-square, for the LRT gives a slightly different but very close result:

```
anova(fit0, fit.hiv2, test="LRT")
```

```
| Analysis of Variance Table
|
| Model 1: LifeExp ~ log(HE) + smoke + hiv + special + log(HE):smo
| Model 2: LifeExp ~ log(HE) * (smoke + I(smoke^2)) + hiv + specia
|   Res.Df    RSS Df Sum of Sq  Pr(>Chi)
|   1      143 2090.8
|   2      141 1858.0  2    232.86 0.0001455 ***
| ---
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

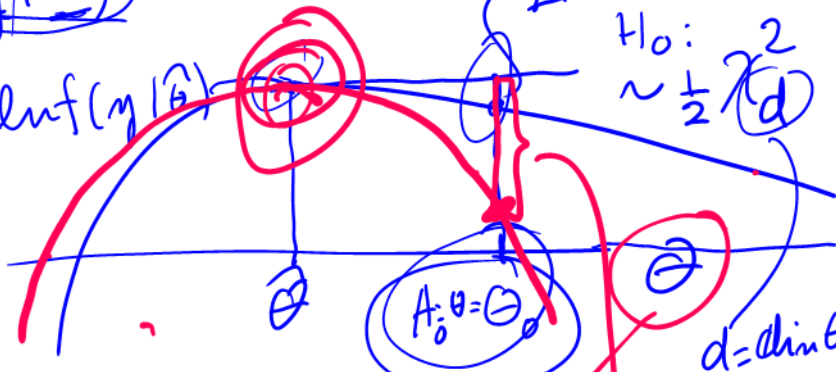
The Wald test and the LRT using the F statistics give identical results. This is the luxury of working with a Gaussian homoskedastic independent linear model.

$\ln f(y|\theta)$

$\ln f(y|\theta)$

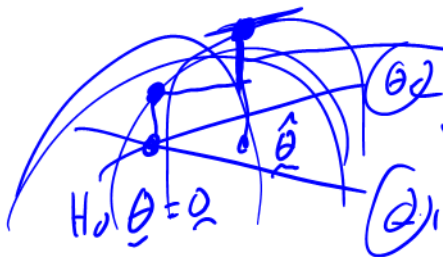
Δ altitude

$H_0: \chi^2$
 $\sim \frac{1}{2} \chi^2(d)$



$A_0: \theta = \theta_0$

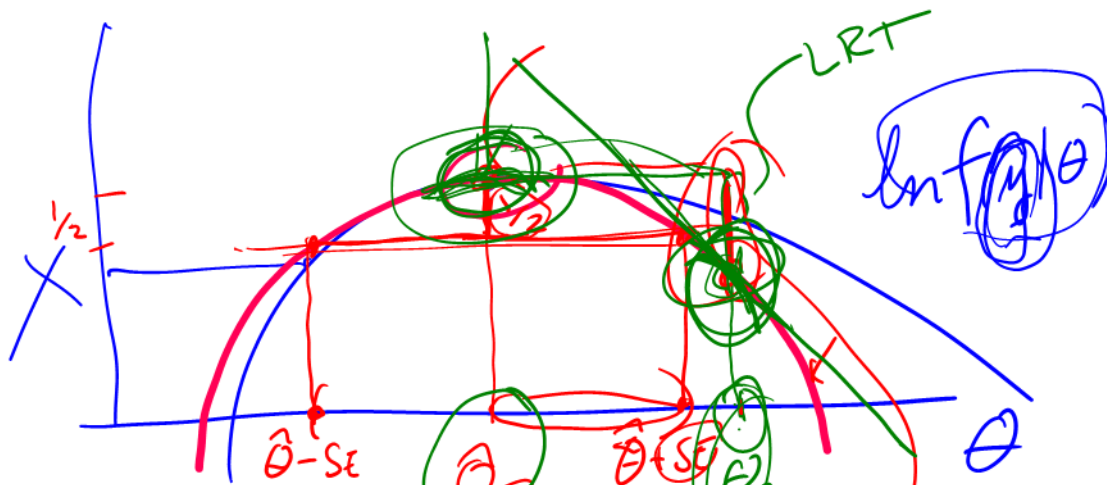
$d = \dim \theta$



$\frac{1}{2} \chi^2_2$

Wald

$\psi = \ln \theta$



$$Z = \frac{\hat{\theta} - \theta_0}{SE}$$

Back at
~ 11 a.m

$$\sigma^2 (S^2 X'X)^{-1}$$

$$H_0: \theta = \theta_0$$

θ_2

θ_1

3.4.1 Exercises

1. Explore the pros and cons of Wald tests versus Likelihood Ratio Tests. Construct an example where they give entirely different results.

3.5 Interpreting sequential tests

Type 1: sequential tests

```
anova(fit.hiv2) # sequential - Type 1 tests
```

```
| Analysis of Variance Table
|
| Response: LifeExp
|
|      Df Sum Sq Mean Sq  F value    Pr(>F)
| log(HE)      1  8631.5   8631.5  655.0400 < 2.2e-16 ***
| smoke        1   152.3    152.3   11.5572 0.0008779 ***
| I(smoke^2)    1   399.7    399.7   30.3307 1.675e-07 ***
| hiv          1  1228.0   1228.0   93.1880 < 2.2e-16 ***
| special      1   878.4    878.4   66.6576 1.637e-13 ***
```



```
| log(HE):smoke          1   81.6    81.6    6.1932 0.0139881 *
| log(HE):I(smoke^2)    1  161.8   161.8   12.2812 0.0006136 ***
| Residuals             141 1858.0    13.2
| ---
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Type 2: Each term added last except for higher-order interactions

```
require(car)
Anova(fit.hiv2)  # Type 2 is default
```

```
| Anova Table (Type II tests)
|
| Response: LifeExp
|
|          Sum Sq  Df  F value    Pr(>F)
| log(HE)      3199.9   1 242.8357 < 2.2e-16 ***
| smoke         129.5   1   9.8297 0.0020900 **
| I(smoke^2)     71.0   1   5.3901 0.0216845 *
| hiv          1235.0   1  93.7227 < 2.2e-16 ***
| special       957.9   1  72.6974 2.109e-14 ***
```

```
| log(HE):smoke          235.0    1  17.8355 4.297e-05 ***
| log(HE):I(smoke^2)    161.8    1  12.2812 0.0006136 ***
| Residuals             1858.0  141
| ---
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Type 3: Each term added last

Note that ‘Type 1’ and ‘Type 2’ sums of squares are interpreted consistently (as far as I’ve seen) in different packages. ‘Type 3’, however, has quite different interpretations. Each variable, or group of variables in the case of a factor with 3 or more levels, is added last but different packages set the other variables at different values. Some set them at their 0 values and others set them at their mean values. What does `car::Anova` do?

```
require(car)
Anova(fit.hiv2, type = 3)
```

```
| Anova Table (Type III tests)
|
```

```

| Response: LifeExp
|
|           Sum Sq  Df F value    Pr(>F)
| (Intercept) 1987.03  1 150.794 < 2.2e-16 ***
| log(HE)     1936.86  1 146.987 < 2.2e-16 ***
| smoke       309.14  1  23.460 3.315e-06 ***
| I(smoke^2)  194.96  1  14.795 0.0001809 ***
| hiv        1235.00  1  93.723 < 2.2e-16 ***
| special     957.94  1  72.697 2.109e-14 ***
| log(HE):smoke 235.02  1  17.835 4.297e-05 ***
| log(HE):I(smoke^2) 161.83  1  12.281 0.0006136 ***
| Residuals   1857.97 141
|
| ---
|
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

Type 3 is identical to regression output except that it uses equivalent F tests and a single test for terms with multiple degrees of freedom

```
summary(fit.hiv2)
```

```
|
```

Call:

```
lm(formula = LifeExp ~ log(HE) * (smoke + I(smoke^2)) + hiv +  
    special, data = dd, na.action = na.exclude)
```

Residuals:

Min	1Q	Median	3Q	Max
-9.0373	-2.3005	0.2043	2.0760	9.7344

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)	
(Intercept)	3.283e+01	2.674e+00	12.280	< 2e-16	***
log(HE)	6.091e+00	5.024e-01	12.124	< 2e-16	***
smoke	3.642e-02	7.520e-03	4.844	3.31e-06	***
I(smoke^2)	-1.518e-05	3.946e-06	-3.846	0.000181	***
hiv	-7.351e-01	7.593e-02	-9.681	< 2e-16	***
special	-1.822e+01	2.137e+00	-8.526	2.11e-14	***
log(HE):smoke	-4.878e-03	1.155e-03	-4.223	4.30e-05	***
log(HE):I(smoke^2)	2.007e-06	5.726e-07	3.504	0.000614	***

```
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1  
|  
| Residual standard error: 3.63 on 141 degrees of freedom  
|   (45 observations deleted due to missingness)  
| Multiple R-squared:  0.8613,    Adjusted R-squared:  0.8544  
| F-statistic:    125 on 7 and 141 DF,  p-value: < 2.2e-16
```

3.6 Working with factors

Controlling for WHO regions provides a non-trivial example of the use of factors in regression.

When you create a data frame in R, non-numeric variables are automatically turned into **factors**. Factors are both a strength of R and a frequent source of annoyance and confusion. See traps and pitfalls with factors.

Let's create a small data frame to illustrate how factors work:

```
set.seed(147)  
sdf <- data.frame( x = c(1:7,6:10),
```

```
g = rep(c('a','b','c'),c(2,5,5))
```

```
sdf
```

```
|      x g
|     1 1 a
|     2 2 a
|     3 3 b
|     4 4 b
|     5 5 b
|     6 6 b
|     7 7 b
|     8 6 c
|     9 7 c
|    10 8 c
|    11 9 c
|    12 10 c
```

```
sdf$y <- with(sdf, x + c(1,0,2)[g] + .5* rnorm(12))
```

```
sdf
```

```
|      x g      y
|  1  1 a  2.120108
|  2  2 a  2.853852
|  3  3 b  2.722852
|  4  4 b  4.825593
|  5  5 b  4.623128
|  6  6 b  5.732132
|  7  7 b  7.221543
|  8  6 c  7.060536
|  9  7 c  9.465820
| 10  8 c  9.748093
| 11  9 c 11.580972
| 12 10 c 11.661232
```

```
sdf$g
```

```
| [1] a a b b b b b c c c c c
| Levels: a b c
```

```
unclass(sdf$g)      # the innards of g
```

```
|      [1] 1 1 2 2 2 2 2 3 3 3 3 3  
|      attr(,"levels")  
|      [1] "a" "b" "c"
```

g is actually a numeric variable consisting of indices into a vector of 'levels'.

```
sfit <- lm( y ~ x + g, sdf, na.action = na.exclude)  
summary(sfit)
```

```
|  
| Call:  
| lm(formula = y ~ x + g, data = sdf, na.action = na.exclude)  
|  
| Residuals:  
|      Min       1Q   Median       3Q      Max  
| -0.7367 -0.3465 -0.1574  0.2736  0.8536  
|  
| Coefficients:
```


	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	0.9074	0.4421	2.052	0.0742 .
x	1.0530	0.1250	8.421	3.01e-05 ***
gb	-1.1476	0.6449	-1.779	0.1131
gc	0.5716	0.9408	0.608	0.5603

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.5662 on 8 degrees of freedom
Multiple R-squared: 0.9797, Adjusted R-squared: 0.972
F-statistic: 128.4 on 3 and 8 DF, p-value: 4.178e-07

Note:

- There is no term called 'ga' although there are 3 levels: 'a', 'b' and 'c'
- The 'missing' level, 'a', is called the **reference level**
- Each term shows a comparison with the reference level

To work out what the coefficients for 'gb' and 'gc' mean, you need to look at the X matrix:

```
model.matrix(sfit, na.action = na.exclude)
```

```
|      (Intercept)  x gb gc
|  1             1  1  0  0
|  2             1  2  0  0
|  3             1  3  1  0
|  4             1  4  1  0
|  5             1  5  1  0
|  6             1  6  1  0
|  7             1  7  1  0
|  8             1  6  0  1
|  9             1  7  0  1
| 10             1  8  0  1
| 11             1  9  0  1
| 12             1 10  0  1
| attr(,"assign")
| [1] 0 1 2 2
| attr(,"contrasts")
```

```
| attr("contrasts")$g  
| [1] "contr.treatment"
```

```
model.matrix(~ x + g, sdf)
```

```
|      (Intercept)  x gb gc  
|  1              1  1  0  0  
|  2              1  2  0  0  
|  3              1  3  1  0  
|  4              1  4  1  0  
|  5              1  5  1  0  
|  6              1  6  1  0  
|  7              1  7  1  0  
|  8              1  6  0  1  
|  9              1  7  0  1  
| 10              1  8  0  1  
| 11              1  9  0  1  
| 12              1 10  0  1  
| attr("assign")
```

```
| [1] 0 1 2 2
| attr(,"contrasts")
| attr(,"contrasts")$g
| [1] "contr.treatment"
```

```
model.matrix(~ g, sdf)
```

```
|      (Intercept) gb gc
| 1             1  0  0
| 2             1  0  0
| 3             1  1  0
| 4             1  1  0
| 5             1  1  0
| 6             1  1  0
| 7             1  1  0
| 8             1  0  1
| 9             1  0  1
| 10            1  0  1
| 11            1  0  1
```

```
| 12          1 0 1
| attr(,"assign")
| [1] 0 1 1
| attr(,"contrasts")
| attr(,"contrasts")$g
| [1] "contr.treatment"
```

If you work through the model:

$$E(y|x, g) = \beta_0 + \beta_x x + \beta_{gb} gb + \beta_{gc} gc$$

where $gb = 1$ if $g = b$ and 0 otherwise, and $gc = 1$ if $g = c$ and 0 otherwise, you see that

1. β_{gb} is the difference between the expected level for group 'b' versus the reference group 'a' keeping x constant and
2. β_{gc} is the same comparison for group 'c' compared with the reference group 'a'.

Plotting fits within groups and panels using `latticeExtra`:

See more elegant but perhaps less flexible approaches in the 'car' and in the 'effects' package by John Fox

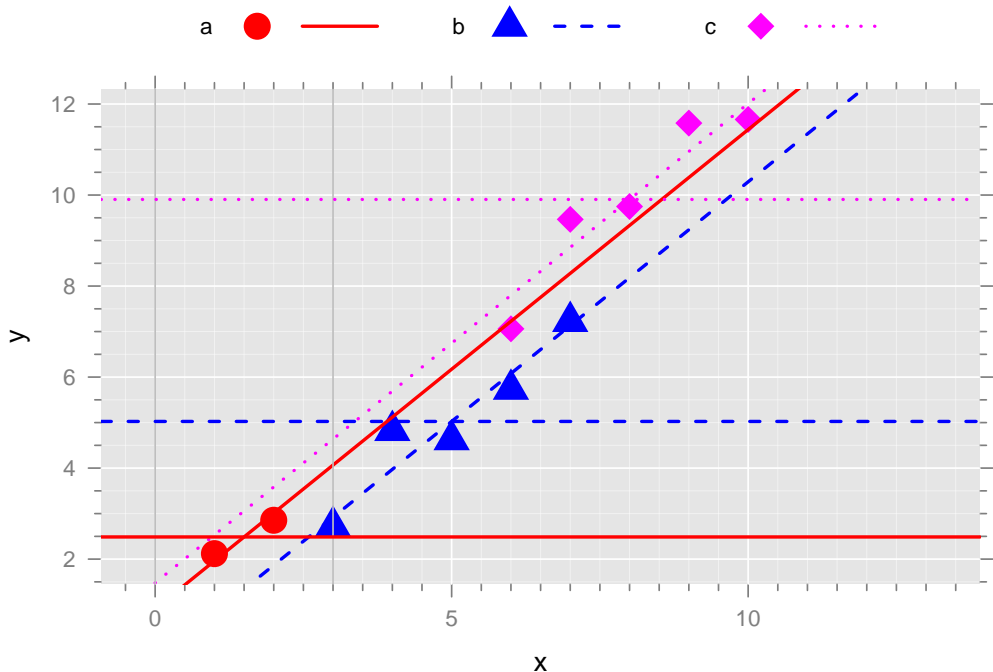
```
pred <- expand.grid( x = 0:13, g = levels( sdf$g))
```

The values over which we want to see predicted lines every combination of x and g

```
pred <- merge( sdf, pred, all = T) # merge with data
pred$y1 <- predict(sfit, newdata = pred) # the predicted value
pred <- sortdf(pred, ~ x) # order so lines won't be interrupted
require(latticeExtra)
require(spida2)
gd(cex=2, lty=1:3, # this gives control over line styles,
  # colour, etc.

  pch = 16:18,
  col = c('red','blue','magenta'),
  lwd =2) # from spida2
xyplot( y ~ x , pred, groups = g ,
  auto.key = list(columns = 3,lines = T),
```

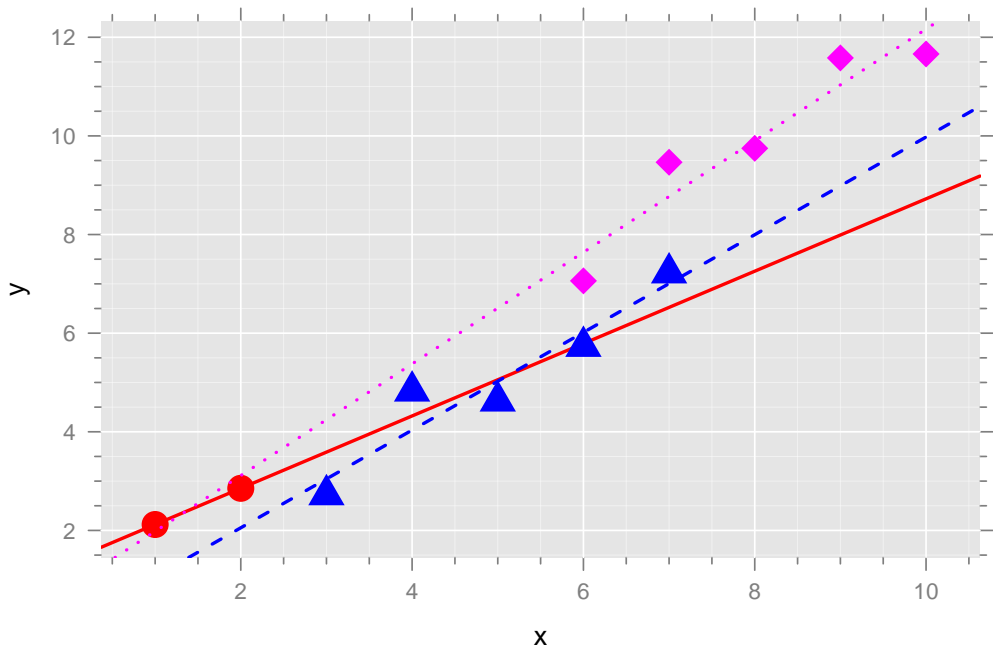
```
y1 = pred$y1, subscripts = T,  
sub =  
"compare adjusted and unadjusted differences between groups") +  
glayer( panel.lines( x, y1[subscripts],...,type = 'l')) +  
layer( panel.abline( v = c(0,3), col = 'grey')) +  
glayer( panel.abline( h = mean(y,na.rm=T),...))
```



compare adjusted and unadjusted differences between groups

an alternative ... but

```
xyplot( y ~ x, sdf, groups = g, type = c('p', 'r'))
```



3.6.1 Exercises:

1. Draw by hand the values of estimated coefficients in the plot.
2. How would you estimate the differences between horizontal lines?
3. Does 'g' matter?

```
summary(sfit) # p-values not significant
```

```
|  
| Call:  
| lm(formula = y ~ x + g, data = sdf, na.action = na.exclude)  
|  
| Residuals:  
|      Min       1Q   Median       3Q      Max  
| -0.7367 -0.3465 -0.1574  0.2736  0.8536  
|  
| Coefficients:  
|              Estimate Std. Error t value Pr(>|t|)  
| (Intercept)   0.9074     0.4421   2.052  0.0742 .  
| x              1.0530     0.1250   8.421 3.01e-05 ***
```

```

|   gb          -1.1476      0.6449  -1.779   0.1131
|   gc           0.5716      0.9408   0.608   0.5603
|   ---
|   Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
|
|   Residual standard error: 0.5662 on 8 degrees of freedom
|   Multiple R-squared:  0.9797,    Adjusted R-squared:  0.972
|   F-statistic: 128.4 on 3 and 8 DF,  p-value: 4.178e-07

```

But

```
wald(sfit, "g") # simultaneous test that both are 0: different ans
```

```

|   numDF denDF  F.value p.value
|   g      2      8 9.711573 0.00724
|   Estimate Std.Error DF t-value  p-value Lower 0.95 Upper 0.9
|   gb -1.147573 0.644944  8  -1.779337 0.11306 -2.634817  0.339671
|   gc  0.571585 0.940783  8   0.607563 0.56032 -1.597865  2.741035

```

Using the GLH (General Linear Hypothesis)

```
Lmu.6 <-list( "at x = 6" = rbind(  
  'g = a' = c(1,6,0,0),  
  'g = b' = c(1,6,1,0),  
  'g = c' = c(1,6,0,1)))  
Lmu.6
```

```
| $`at x = 6`  
|      [,1] [,2] [,3] [,4]  
| g = a    1    6    0    0  
| g = b    1    6    1    0  
| g = c    1    6    0    1
```

```
wald(sfit, Lmu.6)
```

```
|          numDF denDF  F.value p.value  
| at x = 6      3      8 613.0644 <.00001  
|          Estimate Std.Error DF t-value  p-value Lower 0.95 Upper 0.  
| g = a  7.225664  0.690607  8  10.46277 1e-05  5.633120  8.818207  
| g = b  6.078091  0.282401  8  21.52290 <.00001  5.426872  6.729309
```

```
| g = c 7.797249 0.355897 8 21.90875 <.00001 6.976550 8.617948
```

```
Ldiff <- rbind(  
  'b - a' = c(0,0,1,0),  
  'c - a' = c(0,0,0,1),  
  'c - b' = c(0,0,-1,1))
```

```
Ldiff
```

```
|          [,1] [,2] [,3] [,4]  
| b - a      0    0    1    0  
| c - a      0    0    0    1  
| c - b      0    0   -1    1
```

```
wald(sfit,Ldiff)
```

```
|      numDF denDF  F.value p.value  
|      1      2      8 9.711573 0.00724  
|      Estimate Std.Error DF t-value  p-value Lower 0.95 Upper  
| b - a -1.147573 0.644944  8  -1.779337 0.11306 -2.634817  0.3396  
| c - a  0.571585 0.940783  8   0.607563 0.56032 -1.597865  2.7410
```

```
| c - b 1.719159 0.518616 8 3.314900 0.01062 0.523229 2.9150
wald(sfit, 'g')
```

```
| numDF denDF F.value p.value
| g      2      8 9.711573 0.00724
| Estimate Std.Error DF t-value p-value Lower 0.95 Upper 0.9
| gb -1.147573 0.644944 8 -1.779337 0.11306 -2.634817 0.339671
| gc 0.571585 0.940783 8 0.607563 0.56032 -1.597865 2.741035
```

This illustrated the crucial point that separate tests of

$$H_0 : \beta_1 = 0$$

and

$$H_0 : \beta_2 = 0$$

can yield very different ‘conclusions’ that a test of the joint hypothesis:

$$H_0 : \beta_1 = \beta_2 = 0$$

Later, we will see how the relationship between confidence ellipses(oids) and tests makes this clear.

3.6.2 Reparametrization to answer different questions

```
sdf$g2 <- relevel(sdf$g, 'b') # makes 'b' the reference level
fitr <- lm( y ~ x + g2, sdf)
summary(fitr)
```

```
|
| Call:
| lm(formula = y ~ x + g2, data = sdf)
|
| Residuals:
|      Min       1Q   Median       3Q      Max
| -0.7367 -0.3465 -0.1574  0.2736  0.8536
|
| Coefficients:
|              Estimate Std. Error t value Pr(>|t|)
| (Intercept)  -0.2402     0.6746  -0.356   0.7310
| x             1.0530     0.1250   8.421 3.01e-05 ***
| g2a          1.1476     0.6449   1.779   0.1131
```



```
| g2c          1.7192      0.5186      3.315      0.0106 *  
| ---  
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1  
|  
| Residual standard error: 0.5662 on 8 degrees of freedom  
| Multiple R-squared:  0.9797,    Adjusted R-squared:  0.972  
| F-statistic: 128.4 on 3 and 8 DF,  p-value: 4.178e-07
```

```
fitr2 <- lm( y ~ x + g2 -1 , sdf)  # dropping the intercept  
summary(fitr2)
```

```
|  
| Call:  
| lm(formula = y ~ x + g2 - 1, data = sdf)  
|  
| Residuals:  
|      Min       1Q   Median       3Q      Max  
| -0.7367 -0.3465 -0.1574  0.2736  0.8536
```

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)	
x	1.0530	0.1250	8.421	3.01e-05	***
g2b	-0.2402	0.6746	-0.356	0.7310	
g2a	0.9074	0.4421	2.052	0.0742	.
g2c	1.4790	1.0319	1.433	0.1897	

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.5662 on 8 degrees of freedom

Multiple R-squared: 0.9961, Adjusted R-squared: 0.9941

F-statistic: 508.3 on 4 and 8 DF, p-value: 1.176e-09

```
fitr3 <- lm( y ~ I(x-6) + g2 -1 , sdf) # recentering x
summary(fitr3) # compare with earlier
```

Call:

```
lm(formula = y ~ I(x - 6) + g2 - 1, data = sdf)
```

Residuals:

Min	1Q	Median	3Q	Max
-0.7367	-0.3465	-0.1574	0.2736	0.8536

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)	
I(x - 6)	1.0530	0.1250	8.421	3.01e-05	***
g2b	6.0781	0.2824	21.523	2.29e-08	***
g2a	7.2257	0.6906	10.463	6.05e-06	***
g2c	7.7972	0.3559	21.909	1.99e-08	***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.5662 on 8 degrees of freedom

Multiple R-squared: 0.9961, Adjusted R-squared: 0.9941

F-statistic: 508.3 on 4 and 8 DF, p-value: 1.176e-09

```
wald(sfit, Lmu.6)
```

```
|          numDF denDF  F.value p.value
|  at x = 6      3      8 613.0644 <.00001
|          Estimate Std.Error DF t-value  p-value Lower 0.95 Upper 0.
|  g = a 7.225664 0.690607  8  10.46277 1e-05   5.633120   8.818207
|  g = b 6.078091 0.282401  8  21.52290 <.00001 5.426872   6.729309
|  g = c 7.797249 0.355897  8  21.90875 <.00001 6.976550   8.617948
```

3.6.3 Equivalent models

What makes the last three models equivalent?

```
summary(lm( model.matrix(fitr) ~ model.matrix(sfit)-1))
```

```
| Warning in summary.lm(object, ...): essentially perfect fit: sum
| unreliable
```

```
| Warning in summary.lm(object, ...): essentially perfect fit: sum
| unreliable
```

```
| Warning in summary.lm(object, ...): essentially perfect fit: sum
| unreliable
```

```
| Warning in summary.lm(object, ...): essentially perfect fit: sum
| unreliable
```

```
| Response (Intercept) :
```

```
| Call:
```

```
| lm(formula = `(Intercept)` ~ model.matrix(sfit) - 1)
```

```
| Residuals:
```

```
|           Min           1Q           Median           3Q           Max
| -7.504e-16 -4.690e-17  0.000e+00  4.690e-17  7.504e-16
```

```
| Coefficients:
```

```
|                                     Estimate Std. Error  t value P
| model.matrix(sfit)(Intercept)  1.000e+00  2.966e-16  3.371e+15
```

```
| model.matrix(sfit)x          -3.752e-17  8.390e-17 -4.470e-01
| model.matrix(sfit)gb       -6.379e-16  4.327e-16 -1.474e+00
| model.matrix(sfit)gc       -5.253e-16  6.312e-16 -8.320e-01
```

```
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
| Residual standard error: 3.799e-16 on 8 degrees of freedom
```

```
| Multiple R-squared:      1, Adjusted R-squared:      1
```

```
| F-statistic: 2.079e+31 on 4 and 8 DF,  p-value: < 2.2e-16
```

```
| Response x :
```

```
| Call:
```

```
| lm(formula = x ~ model.matrix(sfit) - 1)
```

```
| Residuals:
```

```
|           Min           1Q           Median           3Q           Max
| -8.742e-16 -8.917e-17 -2.706e-17  1.417e-16  8.742e-16
```

Coefficients:

	Estimate	Std. Error	t value	P
model.matrix(sfit)(Intercept)	-1.026e-15	3.623e-16	-2.831e+00	
model.matrix(sfit)x	1.000e+00	1.025e-16	9.759e+15	
model.matrix(sfit)gb	-8.052e-16	5.285e-16	-1.523e+00	
model.matrix(sfit)gc	-9.918e-16	7.709e-16	-1.287e+00	

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 4.64e-16 on 8 degrees of freedom

Multiple R-squared: 1, Adjusted R-squared: 1

F-statistic: 5.459e+32 on 4 and 8 DF, p-value: < 2.2e-16

Response g2a :

Call:

lm(formula = g2a ~ model.matrix(sfit) - 1)

Residuals:

Min	1Q	Median	3Q	Max
-5.639e-17	-2.701e-17	-8.327e-18	3.385e-17	6.024e-17

Coefficients:

	Estimate	Std. Error	t value	P
model.matrix(sfit)(Intercept)	1.000e+00	3.675e-17	2.721e+16	
model.matrix(sfit)x	-4.223e-17	1.039e-17	-4.063e+00	
model.matrix(sfit)gb	-1.000e+00	5.361e-17	-1.865e+16	
model.matrix(sfit)gc	-1.000e+00	7.820e-17	-1.279e+16	

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 4.706e-17 on 8 degrees of freedom

Multiple R-squared: 1, Adjusted R-squared: 1

F-statistic: 2.258e+32 on 4 and 8 DF, p-value: < 2.2e-16

Response g2c :

Call:

```
lm(formula = g2c ~ model.matrix(sfit) - 1)
```

Residuals:

Min	1Q	Median	3Q	Max
-1.054e-17	-6.564e-18	-8.950e-20	4.320e-18	1.981e-17

Coefficients:

	Estimate	Std. Error	t value	Pr(>
model.matrix(sfit)(Intercept)	0.000e+00	7.899e-18	0.00e+00	
model.matrix(sfit)x	0.000e+00	2.234e-18	0.00e+00	
model.matrix(sfit)gb	0.000e+00	1.152e-17	0.00e+00	
model.matrix(sfit)gc	1.000e+00	1.681e-17	5.95e+16	<2

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 1.012e-17 on 8 degrees of freedom

```
| Multiple R-squared:      1, Adjusted R-squared:      1  
| F-statistic: 1.222e+34 on 4 and 8 DF,  p-value: < 2.2e-16
```

```
# note the Resid. SE
```

Each model matrix spans exactly the same linear space. Thus their columns are just different bases for the same space and the β s for one model are just a linear transformation of the β s for the other model.

3.6.4 Exercise:

What do the coefficients estimate in each of the following models?. Indicate how each coefficient is related to the first graph shown below.

Factor alone: g - 1

```
summary( fit1 <- lm( y ~ g - 1, sdf))
```

```
|  
| Call:  
| lm(formula = y ~ g - 1, data = sdf)
```

Residuals:

Min	1Q	Median	3Q	Max
-2.8428	-0.4108	-0.1774	0.9497	2.1965

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)	
ga	2.4870	1.1855	2.098	0.0653	.
gb	5.0250	0.7498	6.702	8.83e-05	***
gc	9.9033	0.7498	13.209	3.39e-07	***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 1.677 on 9 degrees of freedom

Multiple R-squared: 0.9613, Adjusted R-squared: 0.9485

F-statistic: 74.59 on 3 and 9 DF, p-value: 1.118e-06

an example where '=' would not work

`model.matrix(fit1)`

```
|      ga gb gc
|  1    1  0  0
|  2    1  0  0
|  3    0  1  0
|  4    0  1  0
|  5    0  1  0
|  6    0  1  0
|  7    0  1  0
|  8    0  0  1
|  9    0  0  1
| 10    0  0  1
| 11    0  0  1
| 12    0  0  1
| attr(,"assign")
| [1] 1 1 1
| attr(,"contrasts")
| attr(,"contrasts")$g
| [1] "contr.treatment"
```

```
summary( fit2 <- lm( y ~ x + g - 1, sdf))
```

```
|  
| Call:
```

```
| lm(formula = y ~ x + g - 1, data = sdf)
```

```
| Residuals:
```

```
|      Min       1Q   Median       3Q      Max  
| -0.7367 -0.3465 -0.1574  0.2736  0.8536
```

```
| Coefficients:
```

```
|      Estimate Std. Error t value Pr(>|t|)  
| x      1.0530     0.1250   8.421 3.01e-05 ***  
| ga     0.9074     0.4421   2.052  0.0742 .  
| gb    -0.2402     0.6746  -0.356  0.7310  
| gc     1.4790     1.0319   1.433  0.1897
```

```
| ---
```

```
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
|  
| Residual standard error: 0.5662 on 8 degrees of freedom  
| Multiple R-squared: 0.9961, Adjusted R-squared: 0.9941  
| F-statistic: 508.3 on 4 and 8 DF, p-value: 1.176e-09
```

```
# an example where '=' would not work
```

```
model.matrix(fit2)
```

```
|      x ga gb gc  
|  1  1  1  0  0  
|  2  2  1  0  0  
|  3  3  0  1  0  
|  4  4  0  1  0  
|  5  5  0  1  0  
|  6  6  0  1  0  
|  7  7  0  1  0  
|  8  6  0  0  1  
|  9  7  0  0  1  
| 10  8  0  0  1  
| 11  9  0  0  1
```

```
| 12 10 0 0 1
| attr(,"assign")
| [1] 1 2 2 2
| attr(,"contrasts")
| attr(,"contrasts")$g
| [1] "contr.treatment"
```

Factor with interaction: g * x

```
sfit2 <- lm( y ~ g * x, sdf)
summary(sfit2)
```

```
|
| Call:
| lm(formula = y ~ g * x, data = sdf)
|
| Residuals:
|      Min       1Q   Median       3Q      Max
| -0.57949 -0.34154 -0.07762  0.29828  0.79094
|
```

```

| Coefficients:
|           Estimate Std. Error t value Pr(>|t|)
| (Intercept)  1.3864    1.4179   0.978   0.366
| gb          -1.3133    1.7596  -0.746   0.484
| gc          -0.5363    2.1597  -0.248   0.812
| x           0.7337    0.8968   0.818   0.445
| gb:x        0.2566    0.9189   0.279   0.789
| gc:x        0.3979    0.9189   0.433   0.680
|
| Residual standard error: 0.6341 on 6 degrees of freedom
| Multiple R-squared:  0.9809,    Adjusted R-squared:  0.9649
| F-statistic: 61.51 on 5 and 6 DF,  p-value: 4.499e-05

```

From which you might conclude that ‘nothing is significant’!

This illustrates that it is often wrong *wrong* **wrong** to form conclusions on the basis of scanning p-values in regression output.

Factor nesting continuous variable: g / x - 1

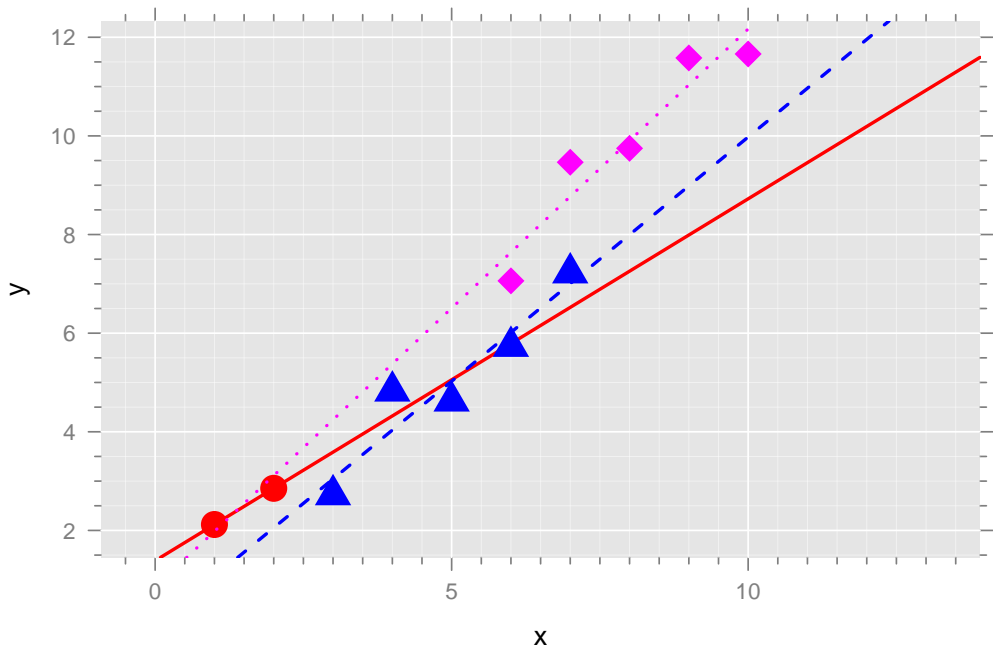

```
sfit3 <- lm( y ~ g / x - 1 , sdf)
summary(sfit3)
```

```
|
| Call:
| lm(formula = y ~ g/x - 1, data = sdf)
|
| Residuals:
|      Min       1Q   Median       3Q      Max
| -0.57949 -0.34154 -0.07762  0.29828  0.79094
|
| Coefficients:
|      Estimate Std. Error t value Pr(>|t|)
| ga      1.38636    1.41789   0.978  0.36595
| gb      0.07309    1.04193   0.070  0.94636
| gc      0.85010    1.62903   0.522  0.62048
| ga:x    0.73374    0.89675   0.818  0.44450
| gb:x    0.99039    0.20052  4.939  0.00261 **
```

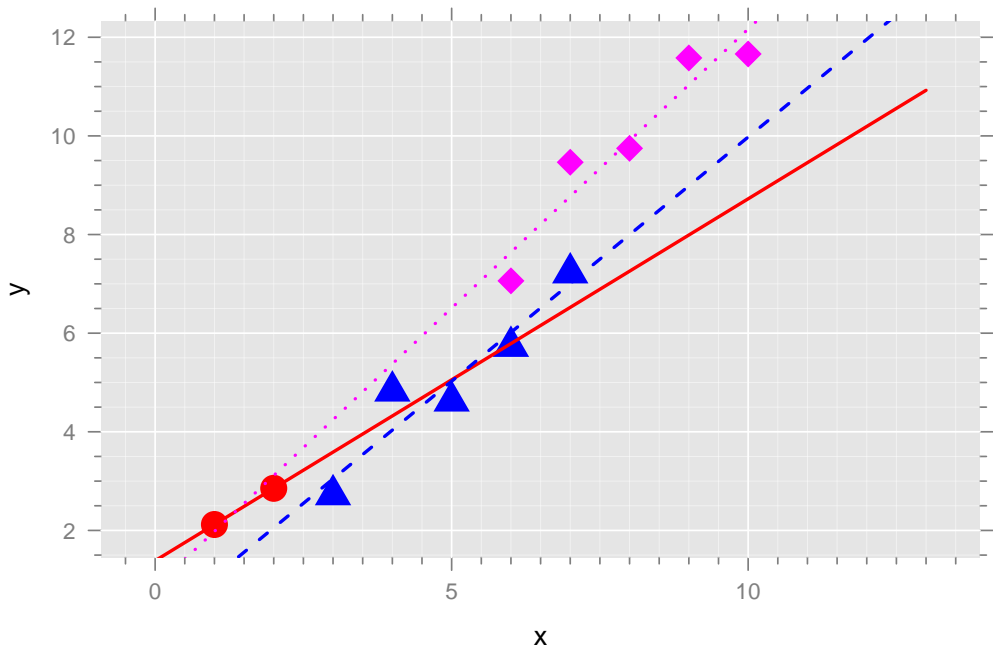
```
| gc:x 1.13165    0.20052    5.644  0.00133 **  
| ---  
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1  
|  
| Residual standard error: 0.6341 on 6 degrees of freedom  
| Multiple R-squared:  0.9963,    Adjusted R-squared:  0.9926  
| F-statistic: 270.2 on 6 and 6 DF,  p-value: 4.985e-07
```

Plotting the fitted model:

```
pred$y2 <- predict( sfit2, newdata = pred)  
xyplot( y ~ x, pred, groups = g, type = c('p','r'))
```

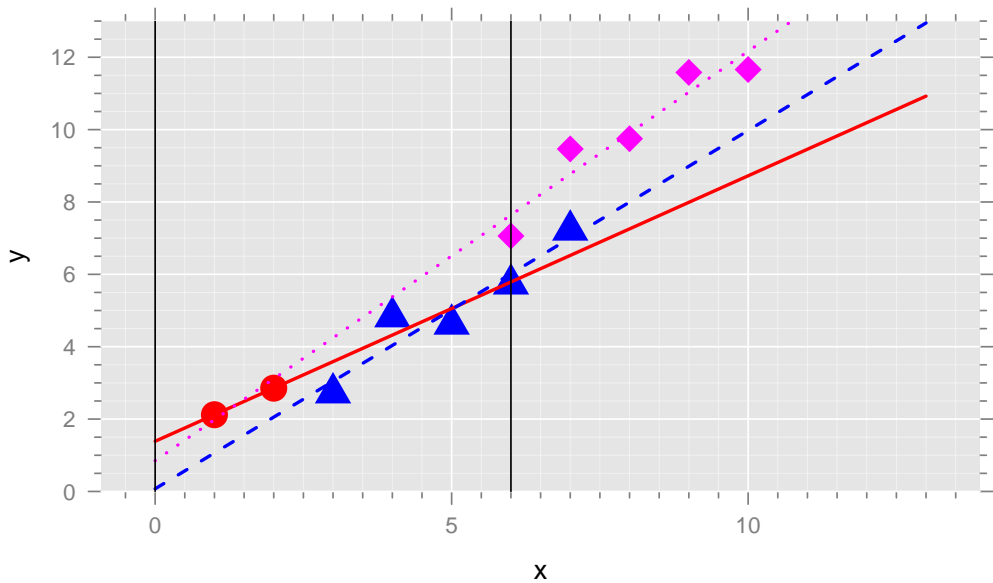


```
# or  
xyplot( y ~ x, pred, groups = g,  
        subscripts = T, y2 = pred$y2, y1 = pred$y1) +  
glayer( panel.xyplot( x, y2[subscripts], ..., type = 'l'))
```



```
# or  
xyplot( y ~ x, pred, groups = g,  
        ylim = c(0,13), auto.key = T,  
        subscripts = T, y2 = pred$y2, y1 = pred$y1) +  
glayer( panel.xyplot( x, y2[subscripts], ..., type = 'l')) +  
layer( panel.abline( v = c(0,6)))
```

a ●
b ▲
c ◆



The model is:

$$E(y|x, g) = \beta_0 + \beta_x x + \beta_{gb} gb + \beta_{gc} gc \\ + \beta_{x:gb} x \times gb + \beta_{x:gc} x \times gc$$

Taking partial derivatives, we see that β_{gb} is the difference between between group 'b' minus group 'a' when $x = 0$. There might not be strong evidence of differences between groups outside the range of the data.

What would happen if we were to explore the difference between group 'b' and group 'c' when $x = 6$:

```
L.bc.6 <- rbind( 'c - b|x=6' =  
                c(0,0,-1,1,-6,6))  
wald( sfit2, L.bc.6)
```

	numDF	denDF	F.value	p.value					
	1	1	6	0.4570639	0.52419				
			Estimate	Std.Error	DF	t-value	p-value	Lower 0.95	Upper
	c - b x=6	2.117587	3.132223	6	0.676065	0.52419	-5.546688	9.78	

We could also do this by reparametrizing:


```
sfit2.x6 <- lm( y ~ I(x-6) * relevel(g,'b'), sdf)
summary(sfit2.x6)
```

```
|
| Call:
```

```
| lm(formula = y ~ I(x - 6) * relevel(g, "b"), data = sdf)
```

```
| Residuals:
```

```
|      Min      1Q   Median      3Q      Max
| -0.57949 -0.34154 -0.07762  0.29828  0.79094
```

```
| Coefficients:
```

```
|
|              Estimate Std. Error t value Pr(>|t|)
| (Intercept)      6.0154    0.3473  17.320 2.37e-06 *
| I(x - 6)         0.9904    0.2005   4.939 0.00261 *
| relevel(g, "b")a -0.2266    4.0750  -0.056 0.95746
| relevel(g, "b")c  1.6246    0.6016   2.701 0.03555 *
| I(x - 6):relevel(g, "b")a -0.2566    0.9189  -0.279 0.78939
```

```
| I(x - 6):relevel(g, "b")c    0.1413    0.2836    0.498    0.63611
| ---
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
|
| Residual standard error: 0.6341 on 6 degrees of freedom
| Multiple R-squared:  0.9809,    Adjusted R-squared:  0.9649
| F-statistic: 61.51 on 5 and 6 DF,  p-value: 4.499e-05
```

Here are some other ways of exploring the model:

```
wald(sfit2, ":")
```

```
|      numDF denDF    F.value p.value
|      :      2      6 0.1890466 0.83249
|      Estimate Std.Error DF t-value p-value Lower 0.95 Upper 0.95
| gb:x 0.256648 0.918898  6  0.27930 0.78939 -1.991815  2.505111
| gc:x 0.397910 0.918898  6  0.43303 0.68013 -1.850553  2.646374
```

```
wald(sfit2, "g")
```

```
|      numDF denDF    F.value p.value
```

	g	4	6	3.965856	0.06566				
		Estimate	Std.Error	DF	t-value	p-value	Lower 0.95	Upper 0	
	gb	-1.313275	1.759556	6	-0.746367	0.48365	-5.618754	2.99220	
	gc	-0.536269	2.159667	6	-0.248311	0.81217	-5.820784	4.74824	
	gb:x	0.256648	0.918898	6	0.279300	0.78939	-1.991815	2.50511	
	gc:x	0.397910	0.918898	6	0.433030	0.68013	-1.850553	2.64637	

```
wald(sfit2, "x")
```

	numDF	denDF	F.value	p.value					
		Estimate	Std.Error	DF	t-value	p-value	Lower 0.95	Upper 0.9	
	x	3	6	18.97152	0.00183				
	x	0.733744	0.896753	6	0.818223	0.44450	-1.460531	2.928020	
	gb:x	0.256648	0.918898	6	0.279300	0.78939	-1.991815	2.505111	
	gc:x	0.397910	0.918898	6	0.433030	0.68013	-1.850553	2.646374	

Using type 2 Anova gives you tests for 'g' and 'x' that assume that higher-order interactions involving 'g' and 'x' are all 0. Note that the error term used is the error term for the full model including interactions. This can lead to inconsistencies with tests based on a model in which interactions has been

dropped.

```
Anova(sfit2) # type 2 anova
```

```
| Anova Table (Type II tests)
```

```
| Response: y
```

	Sum Sq	Df	F value	Pr(>F)	
g	6.2264	2	7.7427	0.0217785	*
x	22.7323	1	56.5365	0.0002865	***
g:x	0.1520	2	0.1890	0.8324941	
Residuals	2.4125	6			

```
| ---
```

```
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
Anova(sfit) # here the gain in degrees of freedom
```

```
| Anova Table (Type II tests)
```

```
| Response: y
```

	Sum Sq	Df	F value	Pr(>F)
x	22.7323	1	70.9133	3.013e-05 ***
g	6.2264	2	9.7116	0.007243 **
Residuals	2.5645	8		

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

outweighs the increase in SSE

3.7 Using Lfx with factors

The 'M' function associated with 'Lfx' can generate code to test for differences between factor levels

```
Lfx(sfit2)
```

```
| list( 1,
| 1 * M(g),
| 1 * x,
| 1 * M(g) * x
```

```
| )
```

The idea is to use the 'Lfx' expression to difference and then to apply to a data frame.

```
dpred <- expand.grid(x = 0:12,  
                    g = levels(sdf$g) ,  
                    g0 = levels(sdf$g))  
  
dim(dpred)
```

```
| [1] 117  3
```

```
some(dpred)
```

```
|      x g g0  
|  4    3 a  a  
|  5    4 a  a  
|  7    6 a  a  
|  9    8 a  a  
| 30    3 c  a  
| 48    8 a  b
```

```
| 70 4 c b
| 91 12 a c
| 92 0 b c
| 100 8 b c
```

we don't need to compare g with g_0 with the same levels and we only need comparisons in one direction (perhaps not)

```
dpred <- subset( dpred, g0 < g)
dim(dpred)
```

```
| [1] 39 3
```

```
some(dpred)
```

```
|      x g g0
| 14 0 b a
| 28 1 c a
| 31 4 c a
| 34 7 c a
| 36 9 c a
```

```
| 38 11 c a
| 69 3 c b
| 71 5 c b
| 72 6 c b
| 75 9 c b
```

```
Lfx(sfit2)
```

```
| list( 1,
| 1 * M(g),
| 1 * x,
| 1 * M(g) * x
| )
```

'difference' $g - g_0$, just like differentiating wrt to g except that 'M' function generates differences

```
Lmat <- Lfx( sfit2,
             list( 0,
                   0 * x,
```



```
1 * M(g,g0),  
1 * x * M(g,g0)  
, dpred)
```

Lmat

	(Intercept)	gb	gc	x	gb:x	gc:x
14	0	0	1	0	0	0
15	0	0	1	0	1	0
16	0	0	1	0	2	0
17	0	0	1	0	3	0
18	0	0	1	0	4	0
19	0	0	1	0	5	0
20	0	0	1	0	6	0
21	0	0	1	0	7	0
22	0	0	1	0	8	0
23	0	0	1	0	9	0
24	0	0	1	0	10	0
25	0	0	1	0	11	0
26	0	0	1	0	12	0

	27	0	0	0	1	0	0
	28	0	0	0	1	0	1
	29	0	0	0	1	0	2
	30	0	0	0	1	0	3
	31	0	0	0	1	0	4
	32	0	0	0	1	0	5
	33	0	0	0	1	0	6
	34	0	0	0	1	0	7
	35	0	0	0	1	0	8
	36	0	0	0	1	0	9
	37	0	0	0	1	0	10
	38	0	0	0	1	0	11
	39	0	0	0	1	0	12
	66	0	0	-1	1	0	0
	67	0	0	-1	1	-1	1
	68	0	0	-1	1	-2	2
	69	0	0	-1	1	-3	3
	70	0	0	-1	1	-4	4
	71	0	0	-1	1	-5	5

	72			0	0	-1	1	-6	6
	73			0	0	-1	1	-7	7
	74			0	0	-1	1	-8	8
	75			0	0	-1	1	-9	9
	76			0	0	-1	1	-10	10
	77			0	0	-1	1	-11	11
	78			0	0	-1	1	-12	12

| attr(,"data")

		x	g	g0
	14	0	b	a
	15	1	b	a
	16	2	b	a
	17	3	b	a
	18	4	b	a
	19	5	b	a
	20	6	b	a
	21	7	b	a
	22	8	b	a
	23	9	b	a

	24	10	b	a
	25	11	b	a
	26	12	b	a
	27	0	c	a
	28	1	c	a
	29	2	c	a
	30	3	c	a
	31	4	c	a
	32	5	c	a
	33	6	c	a
	34	7	c	a
	35	8	c	a
	36	9	c	a
	37	10	c	a
	38	11	c	a
	39	12	c	a
	66	0	c	b
	67	1	c	b
	68	2	c	b

```
| 69 3 c b
| 70 4 c b
| 71 5 c b
| 72 6 c b
| 73 7 c b
| 74 8 c b
| 75 9 c b
| 76 10 c b
| 77 11 c b
| 78 12 c b
```

```
wald(sfit2, Lmat)
```

```
|      numDF denDF F.value p.value
| 1         4      6 63.0834 5e-05
|      Estimate Std.Error DF t-value  p-value Lower 0.95 Upper 0.95
| 14 -0.536269  2.159667 6  -0.248311 0.81217  -5.820784  4.748246
| 15 -0.279621  1.759556 6  -0.158915 0.87895  -4.585100  4.025859
| 16 -0.022973  1.793506 6  -0.012809 0.99020  -4.411523  4.365578
```

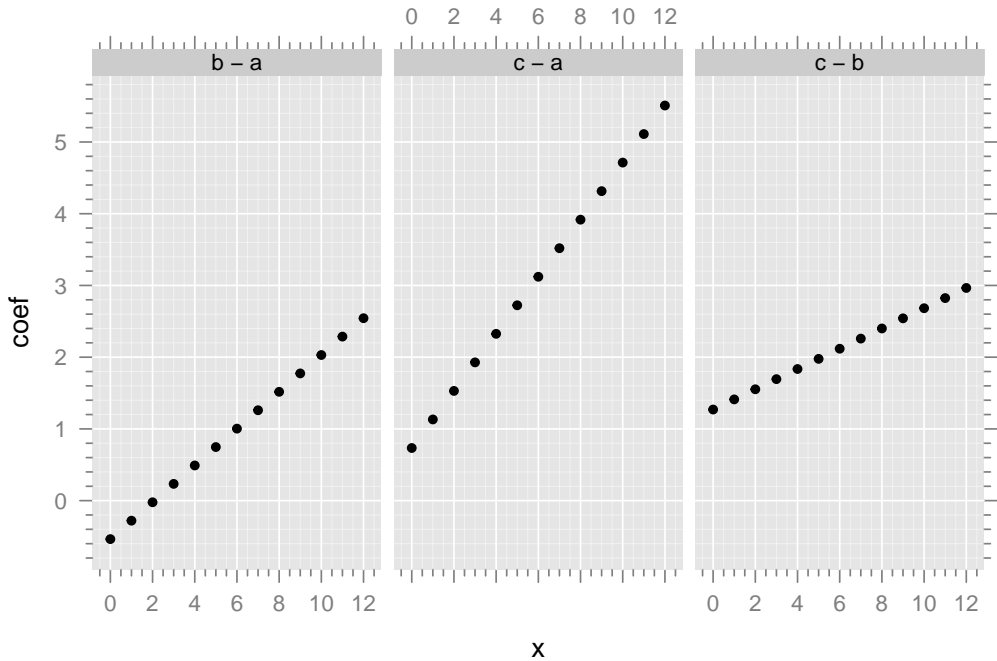
	17	0.233675	2.241882	6	0.104232	0.92038	-5.252013	5.719364
	18	0.490323	2.919616	6	0.167941	0.87215	-6.653720	7.634366
	19	0.746971	3.702841	6	0.201729	0.84679	-8.313553	9.807496
	20	1.003619	4.537251	6	0.221195	0.83228	-10.098634	12.105872
	21	1.260267	5.399168	6	0.233419	0.82320	-11.951020	14.471555
	22	1.516915	6.277271	6	0.241652	0.81710	-13.843013	16.876843
	23	1.773563	7.165612	6	0.247510	0.81277	-15.760057	19.307183
	24	2.030211	8.060806	6	0.251862	0.80955	-17.693872	21.754294
	25	2.286859	8.960801	6	0.255207	0.80709	-19.639432	24.213150
	26	2.543507	9.864282	6	0.257850	0.80514	-21.593523	26.680537
	27	0.733744	0.896753	6	0.818223	0.44450	-1.460531	2.928020
	28	1.131654	0.200520	6	5.643597	0.00133	0.641000	1.622309
	29	1.529565	0.982344	6	1.557057	0.17046	-0.874144	3.933273
	30	1.927475	1.891702	6	1.018910	0.34756	-2.701354	6.556304
	31	2.325385	2.807281	6	0.828341	0.43918	-4.543783	9.194554
	32	2.723296	3.724495	6	0.731185	0.49222	-6.390215	11.836806
	33	3.121206	4.642375	6	0.672330	0.52640	-8.238276	14.480688
	34	3.519116	5.560591	6	0.632867	0.55016	-10.087161	17.125393
	35	3.917027	6.479001	6	0.604573	0.56761	-11.936518	19.770572

	36	4.314937	7.397532	6	0.583294	0.58094	-13.786173	22.416047
	37	4.712847	8.316145	6	0.566711	0.59146	-15.636026	25.061720
	38	5.110757	9.234814	6	0.553423	0.59997	-17.486018	27.707533
	39	5.508668	10.153525	6	0.542537	0.60700	-19.336112	30.353448
	66	1.270013	1.748093	6	0.726513	0.49488	-3.007418	5.547443
	67	1.411275	1.944114	6	0.725922	0.49522	-3.345801	6.168351
	68	1.552537	2.159667	6	0.718878	0.49924	-3.731978	6.837052
	69	1.693800	2.389472	6	0.708859	0.50501	-4.153028	7.540628
	70	1.835062	2.629796	6	0.697796	0.51143	-4.599817	8.269941
	71	1.976324	2.878004	6	0.686700	0.51792	-5.065898	9.018547
	72	2.117587	3.132223	6	0.676065	0.52419	-5.546688	9.781861
	73	2.258849	3.391102	6	0.666111	0.53010	-6.038878	10.556576
	74	2.400111	3.653649	6	0.656908	0.53561	-6.540046	11.340269
	75	2.541374	3.919128	6	0.648454	0.54070	-7.048388	12.131136
	76	2.682636	4.186982	6	0.640709	0.54539	-7.562539	12.927811
	77	2.823898	4.456781	6	0.633618	0.54971	-8.081452	13.729249
	78	2.965161	4.728193	6	0.627123	0.55368	-8.604311	14.534633

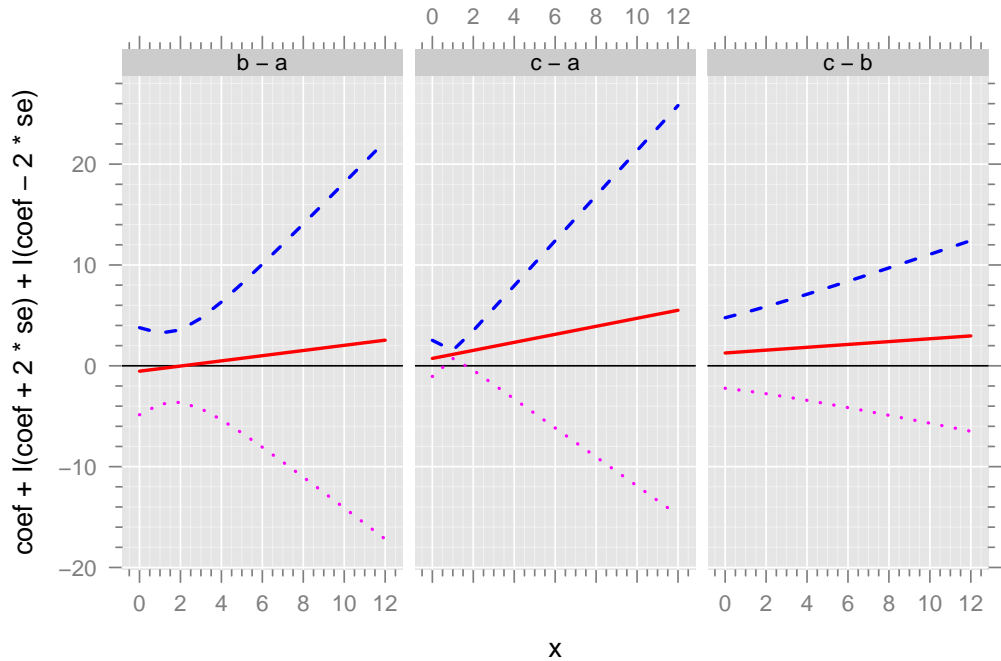
```
ww <- as.data.frame(wald(sfit2, Lmat))  
head(ww)
```

		coef	se	U2	L2	x	g	g0
	14	-0.5362685	2.159667	3.783066	-4.855603	0	b	a
	15	-0.2796206	1.759556	3.239492	-3.798733	1	b	a
	16	-0.0229726	1.793506	3.564039	-3.609984	2	b	a
	17	0.2336754	2.241882	4.717440	-4.250089	3	b	a
	18	0.4903233	2.919616	6.329555	-5.348909	4	b	a
	19	0.7469713	3.702841	8.152652	-6.658710	5	b	a

```
ww$gap <- with(ww, paste( g, '-', g0))  
xyplot( coef ~ x | gap, ww)
```

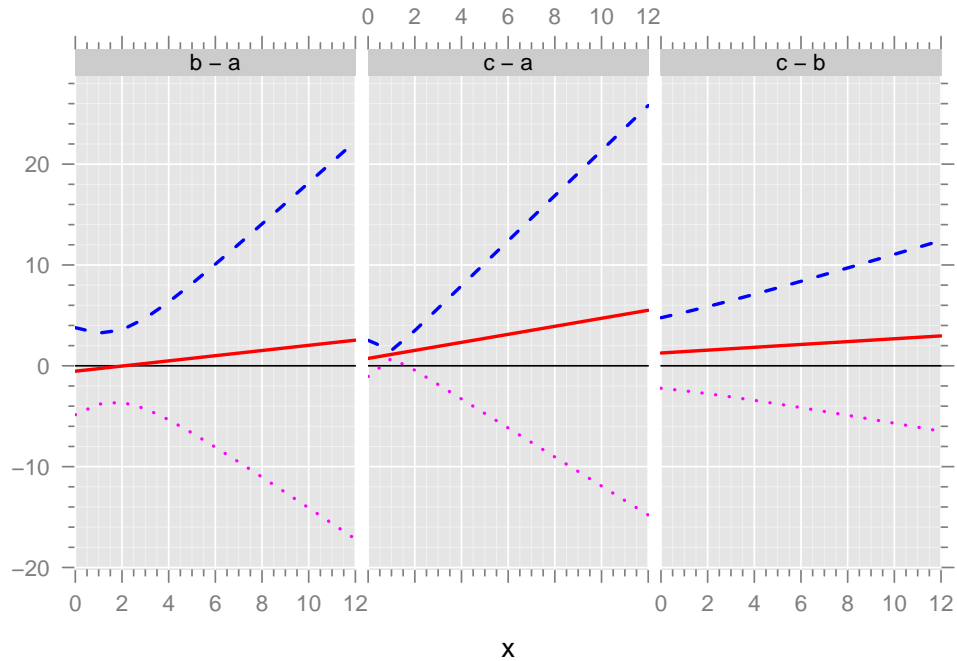
```
td( col = c('black', 'blue', 'blue'), lty = 1, lwd = 2)
xyplot( coef + I(coef+2*se) + I(coef-2*se) ~ x | gap,
        ww, type = 'l') +
layer_( panel.abline( h = 0))
```



OR

```
xyplot( coef +I(coef+2*se) + I(coef-2*se) ~ x | gap,  
        ww, type = 'l',  
        ylab = "Estimated difference plus or minus 2 SEs",  
        xlim = c(0,12)) +  
layer_( panel.abline( h = 0))
```

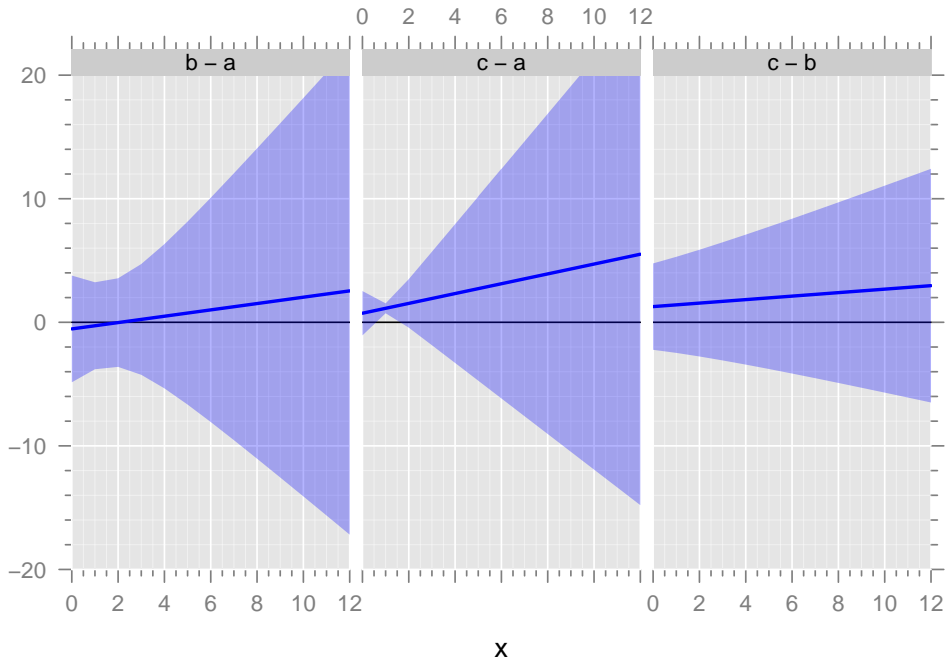
Estimated difference plus or minus 2 SEs



OR

```
gd(col = 'blue')
xyplot( coef ~ x | gap, ww, type = 'l',
        ylab = list("differences between response levels", cex = 1.3),
        xlim = c(0,12),
        ylim = c(-20,20),
        upper = ww$coef + 2* ww$se,
        lower = ww$coef - 2* ww$se,
        subscripts = T) +
layer(panel.fit(...)) +
layer_( panel.abline( h = 0))
```

differences between response levels



Note that if the significant gap between ‘c’ and ‘b’ around $x = 6$ is a question inspired by the data and not a ‘prior’ hypothesis, then some adjustment should be made for **multiplicity**.

3.7.1 Exercises

1. Explore how to modify the appearance of the ‘strips’, i.e. where it says ‘c - b’
2. Plot approximate ‘95% confidence bands’ for each group with ± 2 SEs
3. Plot approximate ‘95% prediction bands’ for each group.

reset the random seed:

```
set.seed(NULL)
```

3.8 Using WHO regions as predictors of Life Expectancy


```
fitr <- lm( LifeExp ~
            (smoke + I(smoke^2)) * region + hiv + special, dd)
summary(fitr)
```

```
|
| Call:
| lm(formula = LifeExp ~ (smoke + I(smoke^2)) * region + hiv +
|     special, data = dd)
|
| Residuals:
|      Min       1Q   Median       3Q      Max
| -10.3209  -3.1528   0.1424   3.0536  10.7776
|
| Coefficients:
|
|             Estimate Std. Error t value Pr(>|t|)
| (Intercept)  5.645e+01  1.512e+00  37.324 < 2e-16 ***
| smoke        9.687e-03  1.343e-02   0.721 0.472152
| I(smoke^2)   1.503e-05  1.811e-05   0.830 0.408190
```

regionAMR	1.079e+01	2.846e+00	3.792	0.000226	***
regionEMR	-1.088e+00	3.106e+00	-0.350	0.726834	
regionEUR	2.211e+01	3.982e+00	5.551	1.5e-07	***
regionSEAR	1.146e+01	3.977e+00	2.881	0.004634	**
regionWPR	8.297e+00	7.772e+00	1.068	0.287671	
hiv	-2.405e-01	1.112e-01	-2.162	0.032401	*
special	-1.006e+01	2.958e+00	-3.402	0.000885	***
smoke:regionAMR	1.876e-02	1.693e-02	1.108	0.269968	
smoke:regionEMR	1.857e-02	1.532e-02	1.212	0.227560	
smoke:regionEUR	-9.910e-03	1.430e-02	-0.693	0.489629	
smoke:regionSEAR	2.485e-03	2.294e-02	0.108	0.913918	
smoke:regionWPR	4.084e-03	2.122e-02	0.192	0.847700	
I(smoke^2):regionAMR	-3.295e-05	1.988e-05	-1.658	0.099715	.
I(smoke^2):regionEMR	-2.432e-05	1.844e-05	-1.319	0.189551	
I(smoke^2):regionEUR	-1.559e-05	1.817e-05	-0.858	0.392392	
I(smoke^2):regionSEAR	-2.559e-05	2.434e-05	-1.051	0.295049	
I(smoke^2):regionWPR	-1.788e-05	1.938e-05	-0.923	0.357846	

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1					

```
|
| Residual standard error: 4.556 on 132 degrees of freedom
|   (42 observations deleted due to missingness)
| Multiple R-squared:  0.8045,    Adjusted R-squared:  0.7764
| F-statistic: 28.6 on 19 and 132 DF,  p-value: < 2.2e-16
```

```
wald(fitr,":")
```

```
|   numDF denDF  F.value p.value
|   :     10   132 3.198898 0.00104
|
|           Estimate Std.Error DF  t-value  p-value
| smoke:regionAMR    0.018757 0.016932 132  1.107787 0.26997
| smoke:regionEMR    0.018572 0.015320 132  1.212307 0.22756
| smoke:regionEUR   -0.009910 0.014304 132 -0.692837 0.48963
| smoke:regionSEAR   0.002485 0.022944 132  0.108305 0.91392
| smoke:regionWPR    0.004084 0.021225 132  0.192431 0.84770
| I(smoke^2):regionAMR -0.000033 0.000020 132 -1.657885 0.09972
| I(smoke^2):regionEMR -0.000024 0.000018 132 -1.318706 0.18955
| I(smoke^2):regionEUR -0.000016 0.000018 132 -0.858101 0.39239
```

I(smoke^2):regionSEAR	-0.000026	0.000024	132	-1.051283	0.29505
I(smoke^2):regionWPR	-0.000018	0.000019	132	-0.922703	0.35785
	Upper	0.95			
smoke:regionAMR	0.052251				
smoke:regionEMR	0.048876				
smoke:regionEUR	0.018384				
smoke:regionSEAR	0.047871				
smoke:regionWPR	0.046069				
I(smoke^2):regionAMR	0.000006				
I(smoke^2):regionEMR	0.000012				
I(smoke^2):regionEUR	0.000020				
I(smoke^2):regionSEAR	0.000023				
I(smoke^2):regionWPR	0.000020				

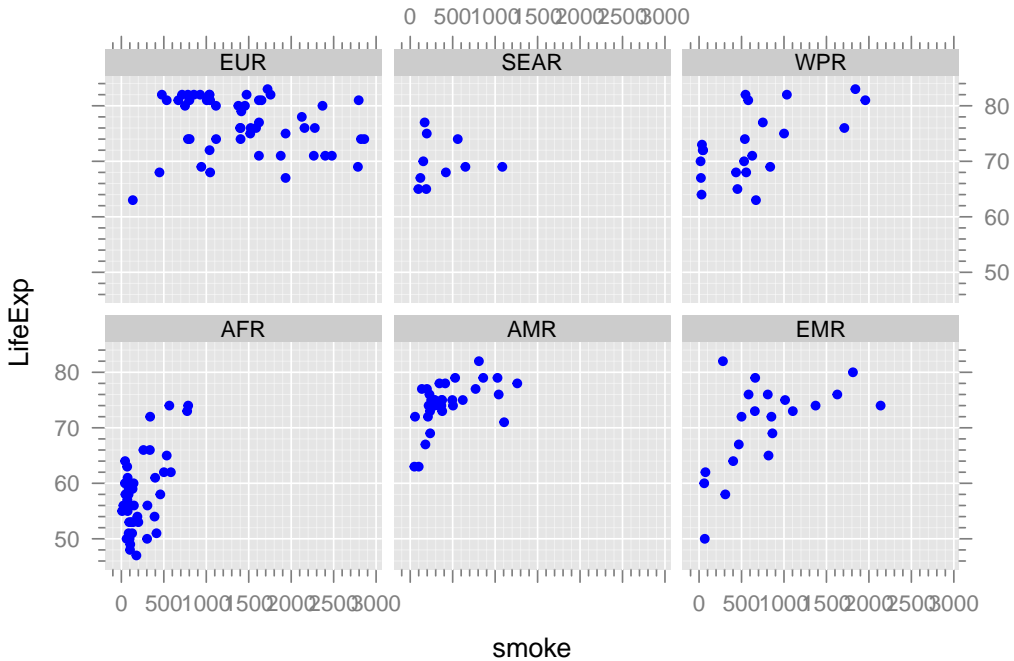
```
wald(fitr,"2):")
```

	numDF	denDF	F.value	p.value					
2):	5	132	2.093536	0.07012					
			Estimate	Std.Error	DF	t-value	p-value	L	

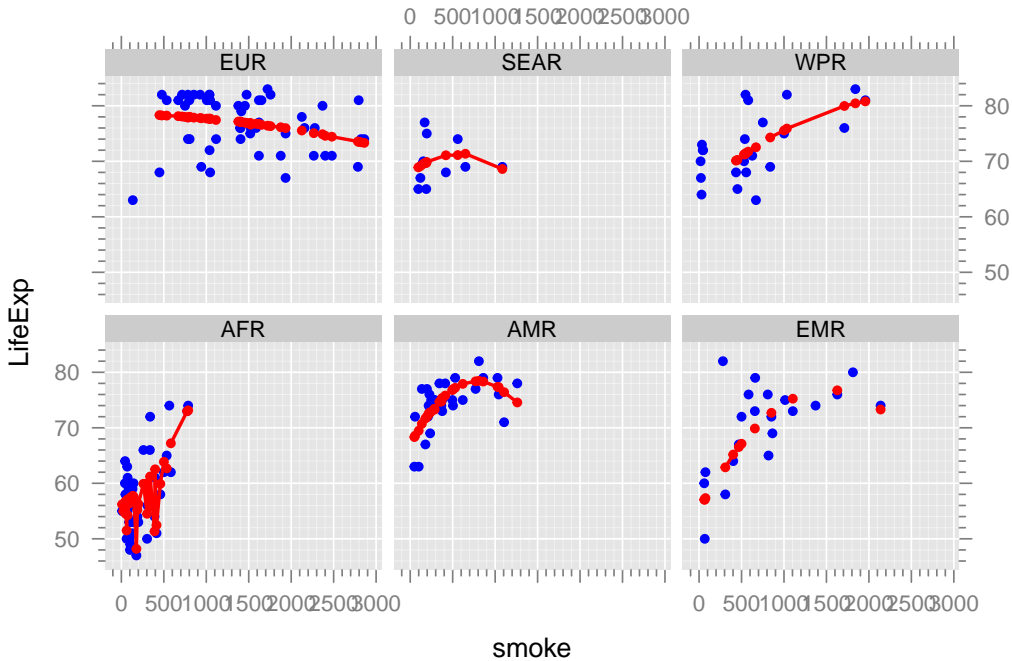
	I(smoke^2):regionAMR	-3.3e-05	2.0e-05	132	-1.657885	0.09972	-
	I(smoke^2):regionEMR	-2.4e-05	1.8e-05	132	-1.318706	0.18955	-
	I(smoke^2):regionEUR	-1.6e-05	1.8e-05	132	-0.858101	0.39239	-
	I(smoke^2):regionSEAR	-2.6e-05	2.4e-05	132	-1.051283	0.29505	-
	I(smoke^2):regionWPR	-1.8e-05	1.9e-05	132	-0.922703	0.35785	-
		Upper	0.95				
	I(smoke^2):regionAMR	6.0e-06					
	I(smoke^2):regionEMR	1.2e-05					
	I(smoke^2):regionEUR	2.0e-05					
	I(smoke^2):regionSEAR	2.3e-05					
	I(smoke^2):regionWPR	2.0e-05					

Using data values for prediction instead of creating a separate prediction data frame: This can work with curvilinear models if the data is sufficiently dense.

```
dd$yq <- predict( fitr, dd)
xyplot( LifeExp ~ smoke | region, dd)
```



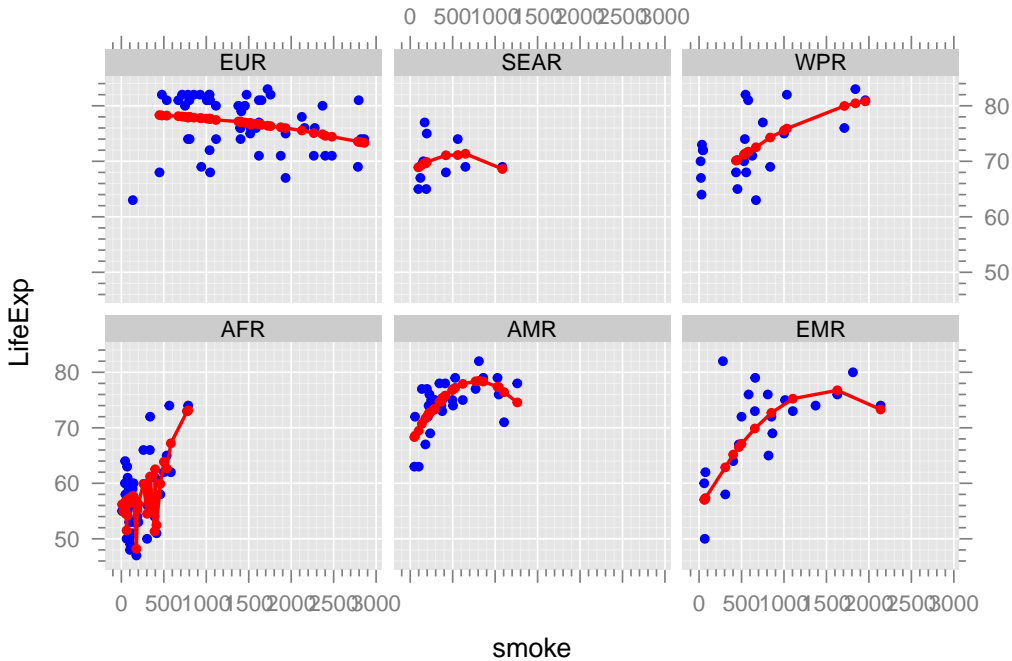
```
# reorder for nice lines:  
dd <- dd[order(dd$region,dd$smoke),]  
xyplot( LifeExp ~ smoke | region,  
        dd, subtitles = T, yq = dd$yq) +  
layer( panel.xyplot( x, yq[subscripts],...,  
                    type = 'b', col = 'red'))
```



presents a problem because of missing hiv values.

Try again keeping non-missing data together to avoid interrupting lines

```
dd <- dd[order(is.na(dd$yq),dd$region,dd$smoke),]  
xyplot( LifeExp ~ smoke | region,  
        dd, subtitles = T, yq = dd$yq) +  
  layer( panel.xyplot( x, yq[subscripts],...,  
                      type = 'b', col = 'red'))
```



(Intercept)	5.088e+01	7.646e+00	6.654	1.12
smoke	-1.832e-02	5.188e-02	-0.353	0.72
I(smoke^2)	4.173e-05	7.885e-05	0.529	0.59
regionAMR	-1.375e+01	1.544e+01	-0.890	0.37
regionEMR	6.606e+00	1.782e+01	0.371	0.71
regionEUR	-8.512e+00	1.948e+01	-0.437	0.66
regionSEAR	1.698e+02	8.098e+01	2.097	0.03
regionWPR	-6.176e+00	2.585e+01	-0.239	0.81
log(HE)	1.527e+00	1.679e+00	0.910	0.36
hiv	-3.930e-01	1.003e-01	-3.917	0.00
special	-1.296e+01	2.343e+00	-5.532	2.13
smoke:regionAMR	6.425e-02	8.185e-02	0.785	0.43
smoke:regionEMR	2.158e-02	6.486e-02	0.333	0.74
smoke:regionEUR	2.975e-02	5.868e-02	0.507	0.61
smoke:regionSEAR	-1.262e+00	6.146e-01	-2.053	0.04
smoke:regionWPR	1.699e-02	7.631e-02	0.223	0.82
I(smoke^2):regionAMR	-5.165e-05	9.443e-05	-0.547	0.58
I(smoke^2):regionEMR	-4.252e-05	8.567e-05	-0.496	0.62
I(smoke^2):regionEUR	-4.762e-05	7.938e-05	-0.600	0.54

I(smoke^2):regionSEAR	1.839e-03	8.954e-04	2.054	0.04
I(smoke^2):regionWPR	-3.620e-05	8.313e-05	-0.435	0.66
smoke:log(HE)	5.178e-03	9.762e-03	0.530	0.59
I(smoke^2):log(HE)	-6.228e-06	1.370e-05	-0.454	0.65
regionAMR:log(HE)	3.852e+00	2.780e+00	1.386	0.16
regionEMR:log(HE)	-1.179e+00	3.844e+00	-0.307	0.75
regionEUR:log(HE)	3.195e+00	2.932e+00	1.090	0.27
regionSEAR:log(HE)	-3.376e+01	1.688e+01	-2.000	0.04
regionWPR:log(HE)	2.831e+00	4.291e+00	0.660	0.51
smoke:regionAMR:log(HE)	-1.044e-02	1.392e-02	-0.750	0.45
smoke:regionEMR:log(HE)	-2.242e-03	1.204e-02	-0.186	0.85
smoke:regionEUR:log(HE)	-6.670e-03	1.044e-02	-0.639	0.52
smoke:regionSEAR:log(HE)	2.685e-01	1.292e-01	2.077	0.04
smoke:regionWPR:log(HE)	-4.551e-03	1.300e-02	-0.350	0.72
I(smoke^2):regionAMR:log(HE)	6.664e-06	1.592e-05	0.419	0.67
I(smoke^2):regionEMR:log(HE)	5.200e-06	1.448e-05	0.359	0.72
I(smoke^2):regionEUR:log(HE)	6.979e-06	1.376e-05	0.507	0.61
I(smoke^2):regionSEAR:log(HE)	-3.900e-04	1.869e-04	-2.087	0.03
I(smoke^2):regionWPR:log(HE)	5.384e-06	1.424e-05	0.378	0.70

```
| ---  
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1  
|  
| Residual standard error: 3.194 on 111 degrees of freedom  
|   (45 observations deleted due to missingness)  
| Multiple R-squared:  0.9154,    Adjusted R-squared:  0.8873  
| F-statistic: 32.48 on 37 and 111 DF,  p-value: < 2.2e-16
```

```
length(coef(fitrhe))
```

```
| [1] 38
```

Should have > 380 observations using Harrell's rules of thumb for valid regression

```
wald(fitrhe, ":")
```

```
|   numDF denDF F.value p.value  
|   :    27   111 1.37843 0.12542  
|  
|               Estimate   Std.Error DF   t-value  
| smoke:regionAMR         0.064254  0.081849 111  0.785035
```

	smoke:regionEMR	0.021575	0.064860	111	0.332641
	smoke:regionEUR	0.029747	0.058678	111	0.506947
	smoke:regionSEAR	-1.261784	0.614556	111	-2.053164
	smoke:regionWPR	0.016993	0.076313	111	0.222673
	I(smoke^2):regionAMR	-0.000052	0.000094	111	-0.546953
	I(smoke^2):regionEMR	-0.000043	0.000086	111	-0.496344
	I(smoke^2):regionEUR	-0.000048	0.000079	111	-0.599894
	I(smoke^2):regionSEAR	0.001839	0.000895	111	2.054470
	I(smoke^2):regionWPR	-0.000036	0.000083	111	-0.435470
	smoke:log(HE)	0.005178	0.009762	111	0.530439
	I(smoke^2):log(HE)	-0.000006	0.000014	111	-0.454460
	regionAMR:log(HE)	3.851782	2.780032	111	1.385517
	regionEMR:log(HE)	-1.179181	3.844186	111	-0.306744
	regionEUR:log(HE)	3.194526	2.932055	111	1.089518
	regionSEAR:log(HE)	-33.758269	16.879974	111	-1.999901
	regionWPR:log(HE)	2.830507	4.290773	111	0.659673
	smoke:regionAMR:log(HE)	-0.010442	0.013922	111	-0.750002
	smoke:regionEMR:log(HE)	-0.002242	0.012041	111	-0.186234
	smoke:regionEUR:log(HE)	-0.006670	0.010440	111	-0.638910

	smoke:regionSEAR:log(HE)	0.268460	0.129242	111	2.077190
	smoke:regionWPR:log(HE)	-0.004551	0.012996	111	-0.350166
	I(smoke^2):regionAMR:log(HE)	0.000007	0.000016	111	0.418627
	I(smoke^2):regionEMR:log(HE)	0.000005	0.000014	111	0.359028
	I(smoke^2):regionEUR:log(HE)	0.000007	0.000014	111	0.507212
	I(smoke^2):regionSEAR:log(HE)	-0.000390	0.000187	111	-2.087196
	I(smoke^2):regionWPR:log(HE)	0.000005	0.000014	111	0.378247
		Lower 0.95	Upper 0.95		
	smoke:regionAMR	-0.097934	0.226443		
	smoke:regionEMR	-0.106950	0.150100		
	smoke:regionEUR	-0.086528	0.146022		
	smoke:regionSEAR	-2.479567	-0.044001		
	smoke:regionWPR	-0.134226	0.168211		
	I(smoke^2):regionAMR	-0.000239	0.000135		
	I(smoke^2):regionEMR	-0.000212	0.000127		
	I(smoke^2):regionEUR	-0.000205	0.000110		
	I(smoke^2):regionSEAR	0.000065	0.003614		
	I(smoke^2):regionWPR	-0.000201	0.000129		
	smoke:log(HE)	-0.014165	0.024521		

I(smoke^2):log(HE)	-0.000033	0.000021
regionAMR:log(HE)	-1.657038	9.360602
regionEMR:log(HE)	-8.796691	6.438330
regionEUR:log(HE)	-2.615536	9.004588
regionSEAR:log(HE)	-67.207064	-0.309474
regionWPR:log(HE)	-5.671945	11.332960
smoke:regionAMR:log(HE)	-0.038029	0.017146
smoke:regionEMR:log(HE)	-0.026102	0.021617
smoke:regionEUR:log(HE)	-0.027359	0.014018
smoke:regionSEAR:log(HE)	0.012358	0.524561
smoke:regionWPR:log(HE)	-0.030303	0.021202
I(smoke^2):regionAMR:log(HE)	-0.000025	0.000038
I(smoke^2):regionEMR:log(HE)	-0.000024	0.000034
I(smoke^2):regionEUR:log(HE)	-0.000020	0.000034
I(smoke^2):regionSEAR:log(HE)	-0.000760	-0.000020
I(smoke^2):regionWPR:log(HE)	-0.000023	0.000034

```
wald(fitrhe, "2")
```

	numDF	denDF	F.value	p.value					
	2	12	111	0.7935878	0.65615				
						Estimate	Std.Error	DF	t-value
I(smoke^2)						0.000042	0.000079	111	0.529253
I(smoke^2):regionAMR						-0.000052	0.000094	111	-0.546953
I(smoke^2):regionEMR						-0.000043	0.000086	111	-0.496344
I(smoke^2):regionEUR						-0.000048	0.000079	111	-0.599894
I(smoke^2):regionSEAR						0.001839	0.000895	111	2.054470
I(smoke^2):regionWPR						-0.000036	0.000083	111	-0.435470
I(smoke^2):log(HE)						-0.000006	0.000014	111	-0.454460
I(smoke^2):regionAMR:log(HE)						0.000007	0.000016	111	0.418627
I(smoke^2):regionEMR:log(HE)						0.000005	0.000014	111	0.359028
I(smoke^2):regionEUR:log(HE)						0.000007	0.000014	111	0.507212
I(smoke^2):regionSEAR:log(HE)						-0.000390	0.000187	111	-2.087196
I(smoke^2):regionWPR:log(HE)						0.000005	0.000014	111	0.378247
						Lower 0.95	Upper 0.95		
I(smoke^2)						-0.000115	0.000198		
I(smoke^2):regionAMR						-0.000239	0.000135		
I(smoke^2):regionEMR						-0.000212	0.000127		

	I(smoke^2):regionEUR	-0.000205	0.000110
	I(smoke^2):regionSEAR	0.000065	0.003614
	I(smoke^2):regionWPR	-0.000201	0.000129
	I(smoke^2):log(HE)	-0.000033	0.000021
	I(smoke^2):regionAMR:log(HE)	-0.000025	0.000038
	I(smoke^2):regionEMR:log(HE)	-0.000024	0.000034
	I(smoke^2):regionEUR:log(HE)	-0.000020	0.000034
	I(smoke^2):regionSEAR:log(HE)	-0.000760	-0.000020
	I(smoke^2):regionWPR:log(HE)	-0.000023	0.000034

```
wald(fitrhe, "2|:.*:") # quadratic terms and 3 and
```

	numDF	denDF	F.value	p.value		
	2 :.*:	17	111	1.286303 0.21443		
			Estimate	Std.Error	DF	t-value
	I(smoke^2)		0.000042	0.000079	111	0.529253
	I(smoke^2):regionAMR		-0.000052	0.000094	111	-0.546953
	I(smoke^2):regionEMR		-0.000043	0.000086	111	-0.496344
	I(smoke^2):regionEUR		-0.000048	0.000079	111	-0.599894

	I(smoke ²):regionSEAR	0.001839	0.000895	111	2.054470
	I(smoke ²):regionWPR	-0.000036	0.000083	111	-0.435470
	I(smoke ²):log(HE)	-0.000006	0.000014	111	-0.454460
	smoke:regionAMR:log(HE)	-0.010442	0.013922	111	-0.750002
	smoke:regionEMR:log(HE)	-0.002242	0.012041	111	-0.186234
	smoke:regionEUR:log(HE)	-0.006670	0.010440	111	-0.638910
	smoke:regionSEAR:log(HE)	0.268460	0.129242	111	2.077190
	smoke:regionWPR:log(HE)	-0.004551	0.012996	111	-0.350166
	I(smoke ²):regionAMR:log(HE)	0.000007	0.000016	111	0.418627
	I(smoke ²):regionEMR:log(HE)	0.000005	0.000014	111	0.359028
	I(smoke ²):regionEUR:log(HE)	0.000007	0.000014	111	0.507212
	I(smoke ²):regionSEAR:log(HE)	-0.000390	0.000187	111	-2.087196
	I(smoke ²):regionWPR:log(HE)	0.000005	0.000014	111	0.378247
		Lower 0.95	Upper 0.95		
	I(smoke ²)	-0.000115	0.000198		
	I(smoke ²):regionAMR	-0.000239	0.000135		
	I(smoke ²):regionEMR	-0.000212	0.000127		
	I(smoke ²):regionEUR	-0.000205	0.000110		
	I(smoke ²):regionSEAR	0.000065	0.003614		

I(smoke^2):regionWPR	-0.000201	0.000129
I(smoke^2):log(HE)	-0.000033	0.000021
smoke:regionAMR:log(HE)	-0.038029	0.017146
smoke:regionEMR:log(HE)	-0.026102	0.021617
smoke:regionEUR:log(HE)	-0.027359	0.014018
smoke:regionSEAR:log(HE)	0.012358	0.524561
smoke:regionWPR:log(HE)	-0.030303	0.021202
I(smoke^2):regionAMR:log(HE)	-0.000025	0.000038
I(smoke^2):regionEMR:log(HE)	-0.000024	0.000034
I(smoke^2):regionEUR:log(HE)	-0.000020	0.000034
I(smoke^2):regionSEAR:log(HE)	-0.000760	-0.000020
I(smoke^2):regionWPR:log(HE)	-0.000023	0.000034

higher way interaction

```
fitr2 <- lm( LifeExp ~
              (smoke + log(HE) + region)^2 + hiv + special, dd)
summary(fitr2)
```

|

Call:

```
lm(formula = LifeExp ~ (smoke + log(HE) + region)^2 + hiv + spec  
    data = dd)
```

Residuals:

Min	1Q	Median	3Q	Max
-6.9873	-1.8511	0.1576	1.7496	8.8413

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)	
(Intercept)	4.534e+01	3.590e+00	12.628	< 2e-16	***
smoke	1.726e-02	4.899e-03	3.522	0.000594	***
log(HE)	2.623e+00	8.903e-01	2.947	0.003816	**
regionAMR	6.004e+00	6.061e+00	0.991	0.323748	
regionEMR	3.025e+00	8.003e+00	0.378	0.706085	
regionEUR	-4.070e+00	7.195e+00	-0.566	0.572584	
regionSEAR	7.659e+00	6.956e+00	1.101	0.272966	
regionWPR	-2.520e+00	5.886e+00	-0.428	0.669332	
hiv	-4.235e-01	9.424e-02	-4.494	1.55e-05	***

special	-1.352e+01	2.183e+00	-6.195	7.35e-09	***
smoke:log(HE)	-6.207e-04	6.668e-04	-0.931	0.353725	
smoke:regionAMR	-1.142e-02	4.176e-03	-2.735	0.007128	**
smoke:regionEMR	-8.776e-03	4.180e-03	-2.099	0.037745	*
smoke:regionEUR	-1.372e-02	3.758e-03	-3.651	0.000379	***
smoke:regionSEAR	-1.321e-02	4.980e-03	-2.652	0.009019	**
smoke:regionWPR	-1.100e-02	4.111e-03	-2.675	0.008450	**
log(HE):regionAMR	8.114e-01	1.180e+00	0.687	0.493061	
log(HE):regionEMR	3.281e-01	1.700e+00	0.193	0.847248	
log(HE):regionEUR	2.389e+00	1.156e+00	2.067	0.040730	*
log(HE):regionSEAR	7.592e-01	1.455e+00	0.522	0.602617	
log(HE):regionWPR	2.073e+00	1.159e+00	1.789	0.075940	.

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 3.254 on 128 degrees of freedom
(45 observations deleted due to missingness)

Multiple R-squared: 0.8988, Adjusted R-squared: 0.883

F-statistic: 56.83 on 20 and 128 DF, p-value: < 2.2e-16

```
wald(fitr2, ':')
```

```
|      numDF denDF  F.value p.value  
|      :      11   128 2.478905 0.00752  
|  
|              Estimate  Std.Error DF  t-value  p-value Low  
| smoke:log(HE)         -0.000621 0.000667 128 -0.930769 0.35373 -0.  
| smoke:regionAMR      -0.011419 0.004176 128 -2.734757 0.00713 -0.  
| smoke:regionEMR      -0.008776 0.004180 128 -2.099411 0.03774 -0.  
| smoke:regionEUR      -0.013722 0.003758 128 -3.651365 0.00038 -0.  
| smoke:regionSEAR     -0.013207 0.004980 128 -2.651798 0.00902 -0.  
| smoke:regionWPR      -0.010997 0.004111 128 -2.674936 0.00845 -0.  
| log(HE):regionAMR    0.811401 1.180355 128 0.687421 0.49306 -1.  
| log(HE):regionEMR    0.328123 1.699935 128 0.193021 0.84725 -3  
| log(HE):regionEUR    2.389164 1.155748 128 2.067201 0.04073 0.  
| log(HE):regionSEAR   0.759205 1.454596 128 0.521935 0.60262 -2.  
| log(HE):regionWPR    2.073361 1.158788 128 1.789249 0.07594 -0.  
|  
|              Upper 0.95  
| smoke:log(HE)         0.000699
```



```

| smoke:regionAMR    -0.003157
| smoke:regionEMR    -0.000505
| smoke:regionEUR    -0.006286
| smoke:regionSEAR   -0.003352
| smoke:regionWPR    -0.002862
| log(HE):regionAMR  3.146934
| log(HE):regionEMR  3.691734
| log(HE):regionEUR  4.676008
| log(HE):regionSEAR 3.637371
| log(HE):regionWPR  4.366221

```

```
wald(fitr2, 'HE):|:log')
```

```

|           numDF denDF   F.value p.value
| HE):|:log      6   128 0.9991023 0.42891
|
|           Estimate Std.Error DF  t-value  p-value Low
| smoke:log(HE)    -0.000621 0.000667 128 -0.930769 0.35373 -0.
| log(HE):regionAMR  0.811401 1.180355 128  0.687421 0.49306 -1.
| log(HE):regionEMR  0.328123 1.699935 128  0.193021 0.84725 -3.

```

	log(HE):regionEUR	2.389164	1.155748	128	2.067201	0.04073	0.
	log(HE):regionSEAR	0.759205	1.454596	128	0.521935	0.60262	-2.
	log(HE):regionWPR	2.073361	1.158788	128	1.789249	0.07594	-0.
		Upper	0.95				
	smoke:log(HE)	0.000699					
	log(HE):regionAMR	3.146934					
	log(HE):regionEMR	3.691734					
	log(HE):regionEUR	4.676008					
	log(HE):regionSEAR	3.637371					
	log(HE):regionWPR	4.366221					

```
fitr3 <- lm( LifeExp ~
              region* smoke + log(HE)+ hiv + special, dd)
summary(fitr3)
```

```
|
| Call:
| lm(formula = LifeExp ~ region * smoke + log(HE) + hiv + special,
|     data = dd)
```

Residuals:

Min	1Q	Median	3Q	Max
-7.016	-1.863	0.165	1.627	8.700

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)	
(Intercept)	41.364150	1.472822	28.085	< 2e-16	***
regionAMR	8.712382	1.438295	6.057	1.31e-08	***
regionEMR	3.799335	1.759763	2.159	0.032631	*
regionEUR	9.842506	1.887832	5.214	6.84e-07	***
regionSEAR	10.122815	1.839269	5.504	1.83e-07	***
regionWPR	7.460923	1.969657	3.788	0.000229	***
smoke	0.010666	0.002712	3.933	0.000134	***
log(HE)	3.688028	0.314332	11.733	< 2e-16	***
hiv	-0.501073	0.079030	-6.340	3.25e-09	***
special	-14.766685	2.013362	-7.334	1.91e-11	***
regionAMR:smoke	-0.009904	0.003175	-3.120	0.002216	**
regionEMR:smoke	-0.007235	0.003068	-2.358	0.019822	*

```

| regionEUR:smoke    -0.011760    0.002835    -4.148 5.93e-05 ***
| regionSEAR:smoke  -0.009663    0.004494    -2.150 0.033335 *
| regionWPR:smoke   -0.008164    0.003110    -2.625 0.009666 **
| ---
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
|
| Residual standard error: 3.254 on 134 degrees of freedom
|   (45 observations deleted due to missingness)
| Multiple R-squared:  0.894, Adjusted R-squared:  0.883
| F-statistic: 80.76 on 14 and 134 DF,  p-value: < 2.2e-16

```

```
wald(fitr3, ":")
```

```

|   numDF denDF F.value p.value
|   :      5   134 4.25484 0.00127
|
|           Estimate Std.Error DF  t-value  p-value Lower
| regionAMR:smoke   -0.009904 0.003175  134 -3.119787 0.00222 -0.01
| regionEMR:smoke   -0.007235 0.003068  134 -2.357952 0.01982 -0.01
| regionEUR:smoke   -0.011760 0.002835  134 -4.147578 0.00006 -0.01

```

	regionSEAR:smoke	-0.009663	0.004494	134	-2.150191	0.03334	-0.01
	regionWPR:smoke	-0.008164	0.003110	134	-2.625207	0.00967	-0.01
		Upper	0.95				
	regionAMR:smoke	-0.003625					
	regionEMR:smoke	-0.001166					
	regionEUR:smoke	-0.006152					
	regionSEAR:smoke	-0.000775					
	regionWPR:smoke	-0.002013					

```
wald(fitr3, 'region')
```

		numDF	denDF	F.value	p.value				
	region	10	134	8.121058	<.00001				
				Estimate	Std.Error	DF	t-value	p-value	Lower
	regionAMR			8.712382	1.438295	134	6.057439	<.00001	5.86
	regionEMR			3.799335	1.759763	134	2.159003	0.03263	0.31
	regionEUR			9.842506	1.887832	134	5.213655	<.00001	6.10
	regionSEAR			10.122815	1.839269	134	5.503717	<.00001	6.48
	regionWPR			7.460923	1.969657	134	3.787931	0.00023	3.56

	regionAMR:smoke	-0.009904	0.003175	134	-3.119787	0.00222	-0.01
	regionEMR:smoke	-0.007235	0.003068	134	-2.357952	0.01982	-0.01
	regionEUR:smoke	-0.011760	0.002835	134	-4.147578	0.00006	-0.01
	regionSEAR:smoke	-0.009663	0.004494	134	-2.150191	0.03334	-0.01
	regionWPR:smoke	-0.008164	0.003110	134	-2.625207	0.00967	-0.01
		Upper 0.95					
	regionAMR	11.557078					
	regionEMR	7.279840					
	regionEUR	13.576310					
	regionSEAR	13.760568					
	regionWPR	11.356561					
	regionAMR:smoke	-0.003625					
	regionEMR:smoke	-0.001166					
	regionEUR:smoke	-0.006152					
	regionSEAR:smoke	-0.000775					
	regionWPR:smoke	-0.002013					

```
wald(fitr3, 'HE')
```

	numDF	denDF	F.value	p.value				
HE	1	134	137.6607	<.00001				
			Estimate	Std.Error	DF	t-value	p-value	Lower 0.95 Upper
log(HE)			3.688028	0.314332	134	11.73289	<.00001	3.066333 4.309

Anova(fitr3)

```

Anova Table (Type II tests)

Response: LifeExp

      Sum Sq   Df  F value    Pr(>F)
region    634.67    5   11.9873 1.339e-09 ***
smoke        6.95    1    0.6561  0.41936
log(HE)   1457.70    1  137.6607 < 2.2e-16 ***
hiv       425.67    1   40.1992 3.250e-09 ***
special   569.62    1   53.7926 1.910e-11 ***
region:smoke 225.27    5    4.2548  0.00127 **
Residuals 1418.94  134
---

```

| Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```
anova(fitr3)
```

| Analysis of Variance Table

| Response: LifeExp

	Df	Sum Sq	Mean Sq	F value	Pr(>F)	
region	5	8942.0	1788.40	168.8905	< 2.2e-16	***
smoke	1	141.2	141.20	13.3348	0.0003725	***
log(HE)	1	1788.2	1788.22	168.8740	< 2.2e-16	***
hiv	1	337.6	337.64	31.8856	9.375e-08	***
special	1	537.9	537.94	50.8014	5.711e-11	***
region:smoke	5	225.3	45.05	4.2548	0.0012701	**
Residuals	134	1418.9	10.59			

| ---

| Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1


```
wald(fitr3, 'smoke')
```

	numDF	denDF	F.value	p.value				
smoke	6	134	3.655058	0.00216				
			Estimate	Std.Error	DF	t-value	p-value	Lower
smoke			0.010666	0.002712	134	3.933292	0.00013	0.00
regionAMR:smoke			-0.009904	0.003175	134	-3.119787	0.00222	-0.01
regionEMR:smoke			-0.007235	0.003068	134	-2.357952	0.01982	-0.01
regionEUR:smoke			-0.011760	0.002835	134	-4.147578	0.00006	-0.01
regionSEAR:smoke			-0.009663	0.004494	134	-2.150191	0.03334	-0.01
regionWPR:smoke			-0.008164	0.003110	134	-2.625207	0.00967	-0.01
			Upper	0.95				
smoke			0.016029					
regionAMR:smoke			-0.003625					
regionEMR:smoke			-0.001166					
regionEUR:smoke			-0.006152					
regionSEAR:smoke			-0.000775					
regionWPR:smoke			-0.002013					

```
library(p3d)
Plot3d(LifeExp ~ smoke + HE | region, dd)
```

```
|   region   col
|  1   AFR   blue
|  2   AMR  green
|  3   EMR orange
|  4   EUR magenta
|  5  SEAR  cyan
|  6   WPR   red
```

```
| Use left mouse to rotate, middle mouse (or scroll) to zoom, right
```

```
Fit3d( fitr3, other.vars=list(hiv=0,special=0))
```

```
| Warning in log(HE): NaNs produced
```

```
#Id3d(par=2, labels = dd$country)
par3d(windowRect=c(10,10,700,700))
rgl.snapshot('regions.png')
```

3.8.1 Exercise

1. Explore this data further producing informative graphs.

4 Exploring Regression Using R

The following is an example of exploring data using regression in R. But it's very unrealistic.

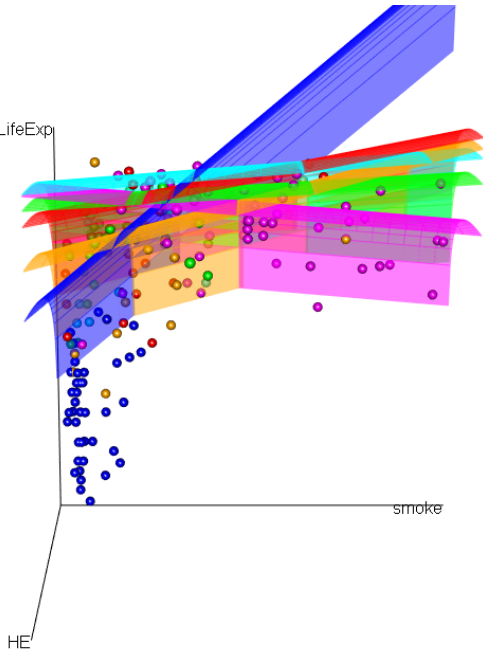
Real data analysis does not start with a neat rectangular data set.

Hadley Wickham: “Data analysis is the process by which data becomes understanding, knowledge and insight”

The process involves much more than running regressions:

- subject matter understanding
- getting data

LifeExp



smoke

HE

- tidying the data
- formulating research questions
- transforming data to variables for analysis
- exploratory visualization
- deciding on a starting model
 - not too big, not too small
 - includes key variables based on subject matter and questions
- modeling fitting
- model diagnostics
- refining the model: dropping some terms and adding others
- formulating parameter functions for estimation and testing
- interpreting results
- GO BACK and iterate a varying number of previous steps in various orders

4.1 Interactive 3D

```
Init3d(cex=1)
ds <- dd
ds$Life <- ds$LE
ds$Cigarettes <- ds$smoke
ds$Health <- ds$HE
ds$area <- ds$region
ds$area <- tr(ds$region,
              c("AFR", "AMR", "EMR", "EUR", "SEAR", "WPR"),
              c("Africa", "South Asia", "Other")[c(1,3,3,3,2,3)])
```

```
Plot3d( Life ~ Cigarettes + Health | region, ds)
```

	region	col
	1 AFR	blue
	2 AMR	green
	3 EMR	orange
	4 EUR	magenta

```
| 5 SEAR cyan  
| 6 WPR red
```

```
| Use left mouse to rotate, middle mouse (or scroll) to zoom, right
```

```
fg()  
spinto()  
Axes3d()  
# Id3d()  
fit <- lm( Life ~ Cigarettes, ds)  
summary(fit)
```

```
|  
| Call:  
| lm(formula = Life ~ Cigarettes, data = ds)  
|  
| Residuals:  
|      Min      1Q   Median      3Q      Max  
| -19.2997 -5.7258  0.6864  6.5300 14.9811
```

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	6.508e+01	8.560e-01	76.03	< 2e-16 ***
Cigarettes	6.915e-03	8.547e-04	8.09	7.99e-14 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 7.969 on 183 degrees of freedom
(9 observations deleted due to missingness)

Multiple R-squared: 0.2635, Adjusted R-squared: 0.2594

F-statistic: 65.46 on 1 and 183 DF, p-value: 7.987e-14

wald(fit)

	numDF	denDF	F.value	p.value
	2	183	7194.457	<.00001

	Estimate	Std.Error	DF	t-value	p-value	Lower	0.95
(Intercept)	65.075840	0.855974	183	76.025515	<.00001	63.386994	

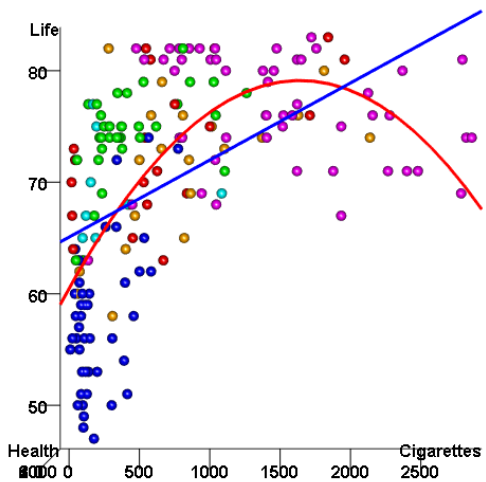

```
| Cigarettes    0.006915 0.000855  183  8.090493 <.00001  0.005228
```

```
Fit3d(fit, lwd = 3)
fitsq <- lm( Life ~ Cigarettes+I(Cigarettes^2), ds)
Fit3d(fitsq, lwd = 3, col = 'red')
spin(0,0,0)
# Id3d(pad=1)
# Id3d("Canada")
# Id3d("United States")
par3d(windowRect=c(10,10,700,700))
rgl.snapshot('quadsmoke.png')
Pop3d(2)
```

4.1.1 Controlling for Health

```
spin(-90,0,0)

fitlin <- lm( Life ~ Cigarettes + Health, ds)
summary(fitlin)
```



Call:

```
lm(formula = Life ~ Cigarettes + Health, data = ds)
```

Residuals:

Min	1Q	Median	3Q	Max
-17.960	-4.309	1.161	5.304	11.772

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)	
(Intercept)	6.368e+01	7.464e-01	85.313	< 2e-16	***
Cigarettes	4.312e-03	7.658e-04	5.631	6.84e-08	***
Health	3.125e-03	3.585e-04	8.717	1.90e-15	***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 6.633 on 179 degrees of freedom

```
| (12 observations deleted due to missingness)
| Multiple R-squared: 0.4786, Adjusted R-squared: 0.4728
| F-statistic: 82.16 on 2 and 179 DF, p-value: < 2.2e-16
```

```
fith <- lm( Life ~ Cigarettes + Health + log( Health),ds)
Fit3d(fitlin, col = 'pink')
Fit3d(fith, col = 'red')
```

```
| Warning in log(Health): NaNs produced
```

```
Pop3d(2)
```

4.2 A more interesting model?

1. Health Expenditures
2. Proportion provided through government

```
ds$propGovt <- with(ds, govt/total) # proportion of health exp.
                                     # from Govt
```

```
Plot3d( Life ~ Health + propGovt |area, ds)
```

```
|           area   col
|    1   Africa  blue
|    2   Other   green
|    3 South Asia orange
```

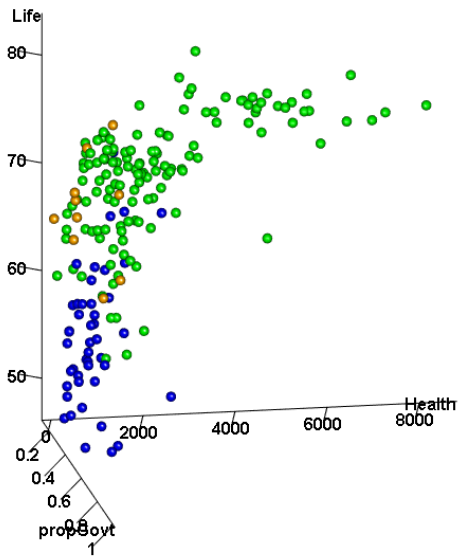
```
| Use left mouse to rotate, middle mouse (or scroll) to zoom, right
```

```
fg()
Axes3d()
spin(-10,15,0)
par3d(windowRect=c(10,10,700,700))
rgl.snapshot('health-pgovt.png')
```

Try something that looks sensible:

The relationship between Life Expectance and Health Expenditures per capita and the proportion of health expenditures funnelled through the government

```
ds$propGovt <- with(ds, govt/total)
               # proportion of health exp. from Govt
```



```
fit <- lm( Life ~ (Health + log(Health) + propGovt) * area , ds,
          na.action = na.exclude)
summary(fit)
```

```
|
| Call:
| lm(formula = Life ~ (Health + log(Health) + propGovt) * area,
|     data = ds, na.action = na.exclude)
|
| Residuals:
|      Min       1Q   Median       3Q      Max
| -13.4282  -1.9480   0.1869   2.1919  14.7809
|
| Coefficients:
|
|             Estimate Std. Error t value Pr(>|t|)
| (Intercept)   39.041364    5.426332    7.195 1.68e-1
| Health       -0.002672    0.003847   -0.695 0.48822
| log(Health)    2.368302    1.239444    1.911 0.05764
```

propGovt	16.083451	3.854716	4.172	4.71e-0
areaOther	4.316456	6.528366	0.661	0.50934
areaSouth Asia	15.629164	15.611788	1.001	0.31813
Health:areaOther	0.002218	0.003872	0.573	0.56736
Health:areaSouth Asia	0.006672	0.015873	0.420	0.67473
log(Health):areaOther	2.454985	1.393231	1.762	0.07977
log(Health):areaSouth Asia	1.029188	4.155759	0.248	0.80468
propGovt:areaOther	-17.211452	4.343678	-3.962	0.00010
propGovt:areaSouth Asia	-21.993937	8.781094	-2.505	0.01315

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 4.007 on 178 degrees of freedom

(4 observations deleted due to missingness)

Multiple R-squared: 0.817, Adjusted R-squared: 0.8057

F-statistic: 72.24 on 11 and 178 DF, p-value: < 2.2e-16

```
Plot3d( Life ~ Health + propGovt | area, ds)
```



```
|           area    col
|  1     Africa   blue
|  2      Other   green
|  3 South Asia orange
```

```
| Use left mouse to rotate, middle mouse (or scroll) to zoom, right
```

```
Fit3d( fit)
```

```
| Warning in log(Health): NaNs produced
```

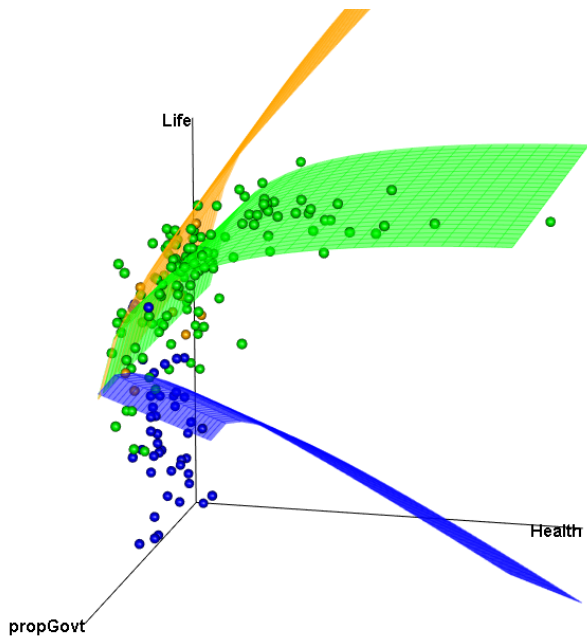
```
spin(13,15,10)
par3d(windowRect=c(10,10,700,700))
rgl.snapshot('health-pgovt-fit.png')
```

Question: Is this model too big for the data?

Should we drop Health expenditures?

None of the coefficients relating to Health are significant.

Can we conclude that “Health” does not add to the predictive power of this



model?

4.2.0.1 Type II SS Type II sums of squares are slightly less prone to massive misinterpretation since each test $\%>\%$ satisfies the POM.

```
Anova(fit) # Type II:
```

```
| Anova Table (Type II tests)
|
| Response: Life
|
|           Sum Sq  Df F value    Pr(>F)
| Health           19.55   1  1.2175 0.2713384
| log(Health)     942.29   1 58.6882 1.142e-12 ***
| propGovt        24.05   1  1.4979 0.2226053
| area           2187.86   2 68.1331 < 2.2e-16 ***
| Health:area       6.63   2  0.2065 0.8136300
| log(Health):area  50.69   2  1.5785 0.2091650
| propGovt:area    269.57   2  8.3948 0.0003281 ***
| Residuals      2857.93 178
```

```
| ---
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
| # slightly less prone to massive misinterpretation
```

4.2.1 Simultaneous tests of groups of coefficients:

```
wald(fit, "Health")
```

```
|          numDF denDF F.value p.value
| Health         6   178 29.4052 <.00001
|
|          Estimate Std.Error DF  t-value  p-value
| Health          -0.002672 0.003847 178 -0.694585 0.484
| log(Health)         2.368302 1.239444 178  1.910778 0.057
| Health:areaOther    0.002218 0.003872 178  0.573005 0.571
| Health:areaSouth Asia 0.006672 0.015873 178  0.420359 0.674
| log(Health):areaOther 2.454985 1.393231 178  1.762080 0.080
| log(Health):areaSouth Asia 1.029188 4.155759 178  0.247653 0.807
|
|          Upper 0.95
```

Health	0.004919
log(Health)	4.814197
Health:areaOther	0.009858
Health:areaSouth Asia	0.037995
log(Health):areaOther	5.204360
log(Health):areaSouth Asia	9.230082

Note:

In the above, note that the overall evidence is **VERY strong** although individual p-values not even significant

I cannot sufficiently stress the importance of the principle this illustrates.

100% of beginning graduates students in statistics programs will fall into the trap of mis-interpreting p-values in regression output. It's as bad a professional error as a doctor amputating the wrong leg.

4.3 Two valid tests:

The Likelihood Ratio Test and the Wald test:

Null model:

```
fit0 <- lm( Life ~ propGovt * area , ds,  
           na.action = na.exclude)
```

OR you can use 'update':

```
fit0 <- update( fit, . ~ propGovt * area)  
summary(fit0)
```

```
|  
| Call:  
| lm(formula = Life ~ propGovt + area + propGovt:area, data = ds,  
|     na.action = na.exclude)  
|  
| Residuals:  
|      Min      1Q   Median      3Q      Max  
| -16.8782  -3.3650   0.6224   3.8491  17.9132  
|  
| Coefficients:
```

	Estimate	Std. Error	t value	Pr(> t)	
(Intercept)	48.387	2.668	18.133	< 2e-16	***
propGovt	19.124	5.023	3.807	0.000191	***
areaOther	20.277	3.153	6.431	1.06e-09	***
areaSouth Asia	19.891	5.055	3.935	0.000118	***
propGovt:areaOther	-9.998	5.631	-1.776	0.077451	.
propGovt:areaSouth Asia	-16.757	9.671	-1.733	0.084828	.

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 5.561 on 184 degrees of freedom
(4 observations deleted due to missingness)
Multiple R-squared: 0.6356, Adjusted R-squared: 0.6257
F-statistic: 64.19 on 5 and 184 DF, p-value: < 2.2e-16

1) LRT:

```
anova( fit, fit0)
```

Analysis of Variance Table

```

|
| Model 1: Life ~ (Health + log(Health) + propGovt) * area
| Model 2: Life ~ propGovt + area + propGovt:area
|   Res.Df    RSS Df Sum of Sq    F    Pr(>F)
| 1     178 2857.9
| 2     184 5690.7 -6    -2832.7 29.405 < 2.2e-16 ***
| ---
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

2) Wald test:

- Doesn't require fitting a new model – works for linear parameters, not necessarily so good for non-linear parameters
- Tests the simultaneous hypotheses that ALL coefficients are = to 0 simultaneously – which is equivalent to dropping Health entirely:

```
wald(fit, "Health")
```

```

|           numDF denDF F.value p.value
| Health      6    178 29.4052 <.00001

```


	Estimate	Std.Error	DF	t-value	p-value
Health	-0.002672	0.003847	178	-0.694585	0.4831
log(Health)	2.368302	1.239444	178	1.910778	0.0587
Health:areaOther	0.002218	0.003872	178	0.573005	0.5691
Health:areaSouth Asia	0.006672	0.015873	178	0.420359	0.6731
log(Health):areaOther	2.454985	1.393231	178	1.762080	0.0801
log(Health):areaSouth Asia	1.029188	4.155759	178	0.247653	0.8111
	Upper 0.95				
Health	0.004919				
log(Health)	4.814197				
Health:areaOther	0.009858				
Health:areaSouth Asia	0.037995				
log(Health):areaOther	5.204360				
log(Health):areaSouth Asia	9.230082				

Note that the F-values are identical – which works in the case of OLS regression with normal error but rarely otherwise.

4.4 Explore interactions

There many approaches to simplifying a model. The most widespread is to trim down non-significant interactions.

If you do this it is vital to never drop a group of terms unless you either:

1. do it one term at a time making sure that you observe the principle of marginality (see the Appendix for more on the principle of marginality) as you go along, or
2. you only drop groups of terms when you have tested them **as a group**.

We could refit and use LRTs or we can use Wald tests:

```
wald(fit, ":") # Use REGULAR EXPRESSION matching
```

```
|      numDF denDF  F.value p.value  
|      :      6   178 4.077738 0.00074  
|  
|              Estimate   Std.Error DF  t-value  p-  
| Health:areaOther        0.002218 0.003872 178  0.573005 0.  
| Health:areaSouth Asia    0.006672 0.015873 178  0.420359 0.
```

	log(Health):areaOther	2.454985	1.393231	178	1.762080	0.
	log(Health):areaSouth Asia	1.029188	4.155759	178	0.247653	0.
	propGovt:areaOther	-17.211452	4.343678	178	-3.962414	0.
	propGovt:areaSouth Asia	-21.993937	8.781094	178	-2.504692	0.
		Lower 0.95	Upper 0.95			
	Health:areaOther	-0.005422	0.009858			
	Health:areaSouth Asia	-0.024651	0.037995			
	log(Health):areaOther	-0.294391	5.204360			
	log(Health):areaSouth Asia	-7.171707	9.230082			
	propGovt:areaOther	-25.783184	-8.639720			
	propGovt:areaSouth Asia	-39.322380	-4.665493			

to test whether there's any evidence of interaction

Explore how regular expressions work:

?regex

Testing all interactions that involve 'Health'

```
wald(fit, "Health.*:")
```

	numDF	denDF	F.value	p.value				
Health.*:	4	178	3.611567	0.0074				
			Estimate	Std.Error	DF	t-value	p-val	
Health:areaOther			0.002218	0.003872	178	0.573005	0.567	
Health:areaSouth Asia			0.006672	0.015873	178	0.420359	0.674	
log(Health):areaOther			2.454985	1.393231	178	1.762080	0.079	
log(Health):areaSouth Asia			1.029188	4.155759	178	0.247653	0.804	
			Upper	0.95				
Health:areaOther			0.009858					
Health:areaSouth Asia			0.037995					
log(Health):areaOther			5.204360					
log(Health):areaSouth Asia			9.230082					

Testing all interactions that involve 'Govt' (always to check to make sure that you captured exactly the right terms)

```
wald(fit, "Govt:")
```

```
|          numDF denDF  F.value p.value
| Govt:         2   178 8.394828 0.00033
|
|              Estimate Std.Error DF  t-value  p-value
| propGovt:areaOther    -17.21145 4.343678 178 -3.962414 0.0001
| propGovt:areaSouth Asia -21.99394 8.781094 178 -2.504692 0.0131
|
|              Upper 0.95
| propGovt:areaOther    -8.639720
| propGovt:areaSouth Asia -4.665493
```

4.4.1 Some comments on reading a model

```
summary(fit)
```

4.4.1.1 Table of coefficients and p-values:

```
|
| Call:
```

```
lm(formula = Life ~ (Health + log(Health) + propGovt) * area,  
    data = ds, na.action = na.exclude)
```

Residuals:

Min	1Q	Median	3Q	Max
-13.4282	-1.9480	0.1869	2.1919	14.7809

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	39.041364	5.426332	7.195	1.68e-1
Health	-0.002672	0.003847	-0.695	0.48822
log(Health)	2.368302	1.239444	1.911	0.05764
propGovt	16.083451	3.854716	4.172	4.71e-0
areaOther	4.316456	6.528366	0.661	0.50934
areaSouth Asia	15.629164	15.611788	1.001	0.31813
Health:areaOther	0.002218	0.003872	0.573	0.56736
Health:areaSouth Asia	0.006672	0.015873	0.420	0.67473
log(Health):areaOther	2.454985	1.393231	1.762	0.07977
log(Health):areaSouth Asia	1.029188	4.155759	0.248	0.80468

```

| propGovt:areaOther          -17.211452    4.343678   -3.962 0.00010
| propGovt:areaSouth Asia    -21.993937    8.781094   -2.505 0.01315
| ---
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
|
| Residual standard error: 4.007 on 178 degrees of freedom
|   (4 observations deleted due to missingness)
| Multiple R-squared:  0.817, Adjusted R-squared:  0.8057
| F-statistic: 72.24 on 11 and 178 DF,  p-value: < 2.2e-16

```

Problems and limitations:

1. Except for very simple models this is generally misleading and not meaningful
2. Very few know how to interpret these correctly, even statisticians
3. The p-value answers how much evidence is there that this term adds to the model when all other terms are already in the model.
4. Only one degree of freedom per coefficient: never asks whether groups of terms are significant which is essential with categorical factors with 3 or more levels.

- Often it's meaningless to change one term keeping others constant, e.g. x if x^2 is also in the model.
- Terms that are marginal to higher order interactions have a specific conditional interpretation that is generally just a very small and arbitrary part of the picture.
- The interpretation of tests does not respect the principle of marginality.

```
anova(fit)
```

4.4.1.2 Type I (sequential) Tests and Sums of Squares

```
| Analysis of Variance Table
```

```
|
```

```
| Response: Life
```

```
|           Df Sum Sq Mean Sq  F value    Pr(>F)
| Health           1  6071.7   6071.7  378.1620 < 2.2e-16 ***
| log(Health)      1  4091.9   4091.9  254.8571 < 2.2e-16 ***
| propGovt         1    14.8     14.8    0.9238  0.3377716
```



```

| area                2 2187.9 1093.9 68.1331 < 2.2e-16 ***
| Health:area        2   79.0   39.5  2.4611 0.0882394 .
| log(Health):area   2   44.2   22.1  1.3773 0.2549413
| propGovt:area      2  269.6  134.8  8.3948 0.0003281 ***
| Residuals          178 2857.9   16.1
| ---
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

The results depend on the order of the terms.

```
anova(update(fit, . ~ area * (propGovt + log(Health) + Health)))
```

```

| Analysis of Variance Table
|

```

```

| Response: Life
|

```

```

|           Df Sum Sq Mean Sq  F value    Pr(>F)
| area       2  9077.6  4538.8 282.6878 < 2.2e-16 ***
| propGovt   1   718.7   718.7  44.7647 2.783e-10 ***
| log(Health) 1  2566.2  2566.2 159.8331 < 2.2e-16 ***
| Health     1     3.8     3.8   0.2359  0.627813

```

```

| area:propGovt      2  160.9    80.4    5.0101  0.007642 **
| area:log(Health)   2  225.3   112.7    7.0166  0.001166 **
| area:Health        2    6.6     3.3     0.2065  0.813630
| Residuals          178 2857.9   16.1
| ---
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

Notes:

1. Each p-values asks whether there's evidence that each terms adds **to the previous terms** listed in the model.
2. Interpretation of tests respects marginality since interactions are listed after their included main effects and sub-interactions.
3. Factors with multiple degrees of freedom are tested jointly.

Anova(fit)

4.4.1.3 Type II Tests and Sums of Squares

Anova Table (Type II tests)

Response: Life

	Sum Sq	Df	F value	Pr(>F)	
Health	19.55	1	1.2175	0.2713384	
log(Health)	942.29	1	58.6882	1.142e-12	***
propGovt	24.05	1	1.4979	0.2226053	
area	2187.86	2	68.1331	< 2.2e-16	***
Health:area	6.63	2	0.2065	0.8136300	
log(Health):area	50.69	2	1.5785	0.2091650	
propGovt:area	269.57	2	8.3948	0.0003281	***
Residuals	2857.93	178			

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Notes:

1. Terms within a 'level' are each added last respecting marginality. e.g. each main effect is added last among main effects but not including higher-order interactions that contain the effect.

2. Main effects are interpretable as test of significance under assumption that there are no interactions.

4.4.1.4 Type III Tests and Sums of Squares

1. Popularized by SAS and SPSS
2. No universal definition so can be misleading
3. With interactions, main effects are averages over levels of interacting variables which can be misleading if groups sizes are unequal.
4. Loved by many researchers, deprecated by most statisticians ... like pie charts.

```
Anova(fit, type = 3)    # Type III Anova: Very popular but ....!
```

```
| Anova Table (Type III tests)
|
| Response: Life
|
|           Sum Sq  Df F value    Pr(>F)
| (Intercept)   831.13   1  51.7651 1.678e-11 ***
| Health         7.75    1  0.4824 0.4882213
```

```

|   log(Health)           58.62    1  3.6511 0.0576404 .
|   propGovt             279.52    1 17.4090 4.706e-05 ***
|   area                 18.52    2  0.5766 0.5628569
|   Health:area          6.63     2  0.2065 0.8136300
|   log(Health):area    50.69    2  1.5785 0.2091650
|   propGovt:area       269.57    2  8.3948 0.0003281 ***
|   Residuals           2857.93 178
|   ---
|   Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

4.4.1.5 Alternative – or supplementary – approaches Answer specific questions.

e.g. None of the above are equivalent to Wald test for ‘OVERALL SIGNIFICANCE’:

```
wald(fit, "Health")
```

```

|           numDF denDF F.value p.value
|   Health      6   178 29.4052 <.00001

```

	Estimate	Std.Error	DF	t-value	p-value
Health	-0.002672	0.003847	178	-0.694585	0.484
log(Health)	2.368302	1.239444	178	1.910778	0.058
Health:areaOther	0.002218	0.003872	178	0.573005	0.571
Health:areaSouth Asia	0.006672	0.015873	178	0.420359	0.675
log(Health):areaOther	2.454985	1.393231	178	1.762080	0.080
log(Health):areaSouth Asia	1.029188	4.155759	178	0.247653	0.811
	Upper 0.95				
Health	0.004919				
log(Health)	4.814197				
Health:areaOther	0.009858				
Health:areaSouth Asia	0.037995				
log(Health):areaOther	5.204360				
log(Health):areaSouth Asia	9.230082				

```
wald(fit, "area")
```

	numDF	denDF	F.value	p.value
area	8	178	20.09158	<.00001

	Estimate	Std.Error	DF	t-value	p-
areaOther	4.316456	6.528366	178	0.661185	0.
areaSouth Asia	15.629164	15.611788	178	1.001113	0.
Health:areaOther	0.002218	0.003872	178	0.573005	0.
Health:areaSouth Asia	0.006672	0.015873	178	0.420359	0.
log(Health):areaOther	2.454985	1.393231	178	1.762080	0.
log(Health):areaSouth Asia	1.029188	4.155759	178	0.247653	0.
propGovt:areaOther	-17.211452	4.343678	178	-3.962414	0.
propGovt:areaSouth Asia	-21.993937	8.781094	178	-2.504692	0.
	Lower 0.95	Upper 0.95			
areaOther	-8.566498	17.199409			
areaSouth Asia	-15.178841	46.437168			
Health:areaOther	-0.005422	0.009858			
Health:areaSouth Asia	-0.024651	0.037995			
log(Health):areaOther	-0.294391	5.204360			
log(Health):areaSouth Asia	-7.171707	9.230082			
propGovt:areaOther	-25.783184	-8.639720			
propGovt:areaSouth Asia	-39.322380	-4.665493			

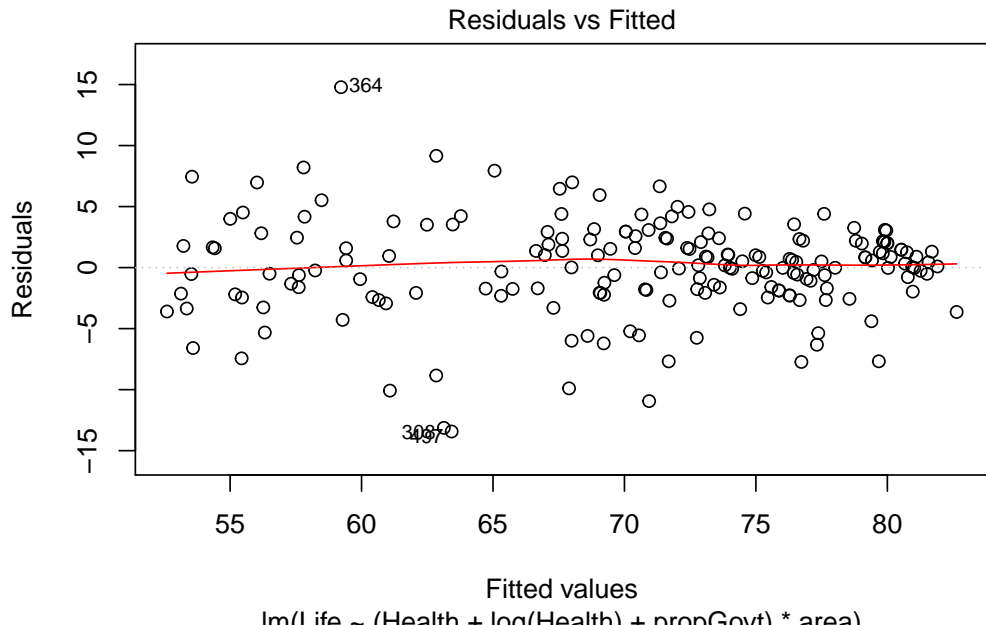
```
wald(fit, "propGovt")
```

```
|          numDF denDF  F.value p.value
| propGovt      3   178 6.095867 0.00057
|
|          Estimate Std.Error DF  t-value  p-value
| propGovt          16.08345 3.854716 178  4.172409 0.0000
| propGovt:areaOther -17.21145 4.343678 178 -3.962414 0.0001
| propGovt:areaSouth Asia -21.99394 8.781094 178 -2.504692 0.0131
|
|          Upper 0.95
| propGovt          23.690273
| propGovt:areaOther -8.639720
| propGovt:areaSouth Asia -4.665493
```

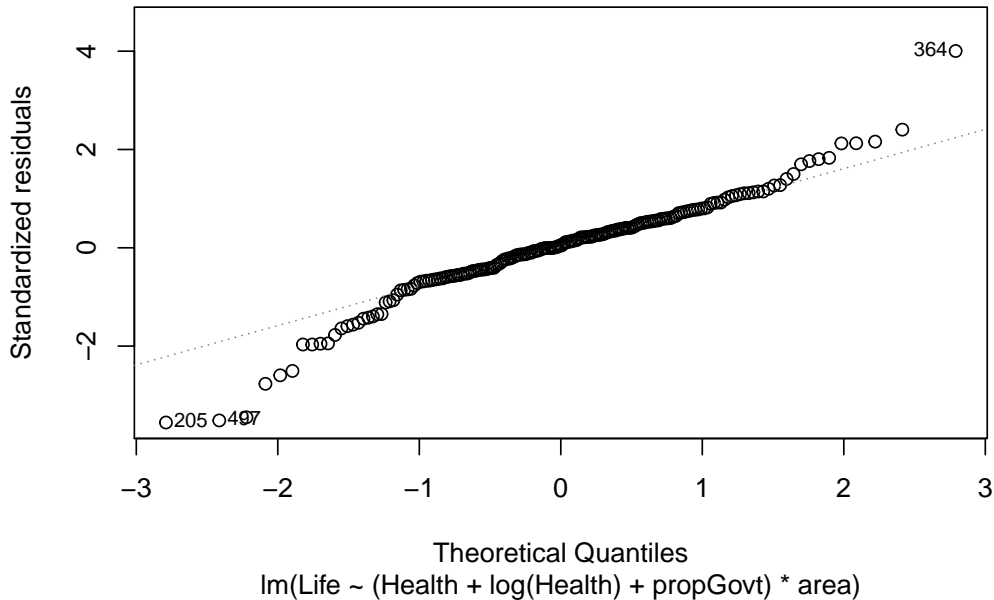
4.5 Regression diagnostics – quick

```
plot(fit)
```

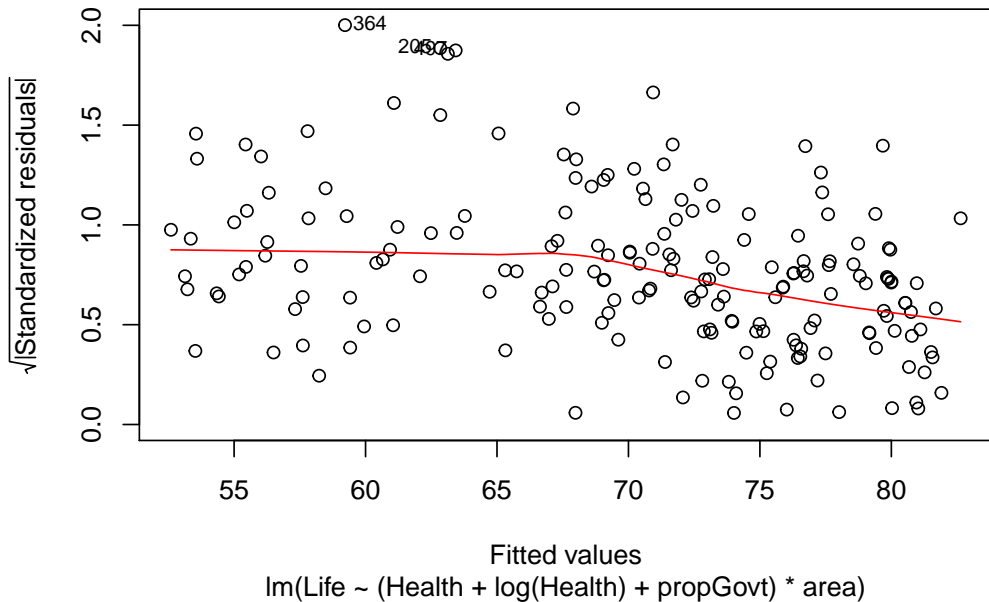

4.5.0.1 Traditional



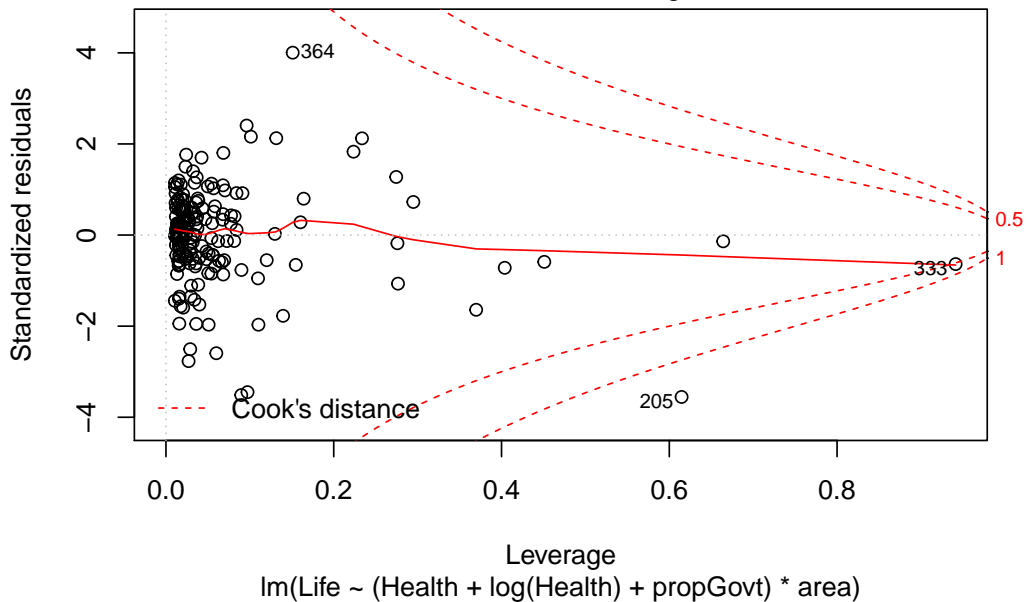
Normal Q-Q



Scale-Location



Residuals vs Leverage



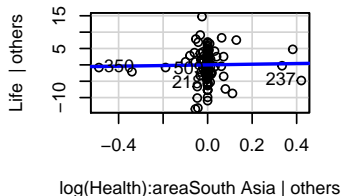
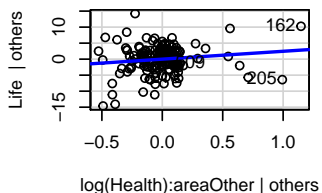
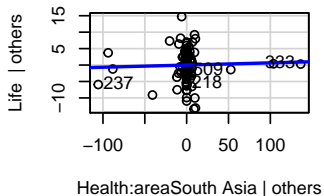
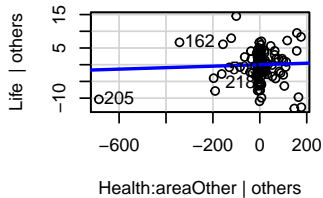
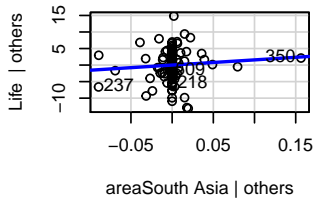
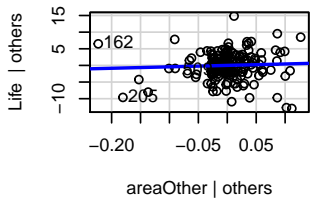
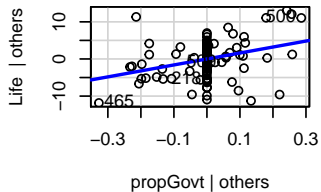
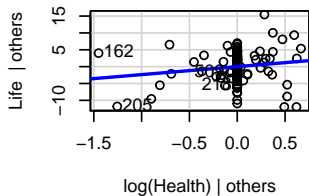
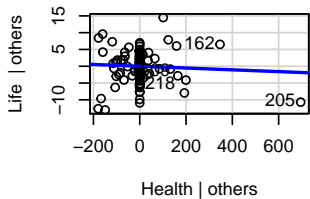
produces 4 plots

- 1) resid ~ fit
- 2) normal quantiles of residuals – Why would this matter??? GEQ. What can it mean if observed residuals are not normal? Clue: Why would you expect them to be normal anyways?
- 3) scale-location for heteroscedasticity
- 4) Residual vs leverage plot – will see deeper meaning of this plot

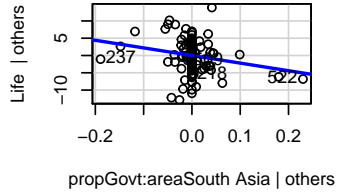
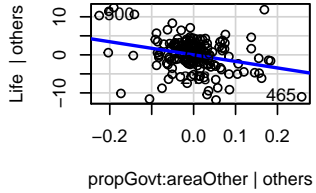
points with high Cook's distance might have strong influence on fitted vals

Note: added-variable plots = partial residual leverage plots

```
avPlots(fit) # look at these as if they
```

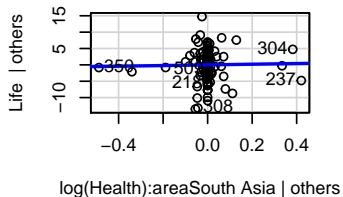
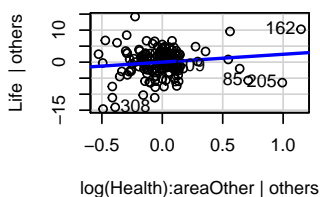
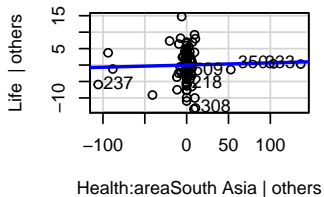
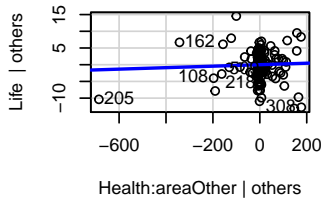
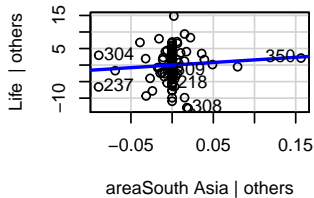
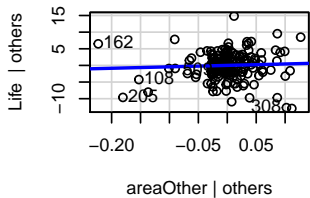
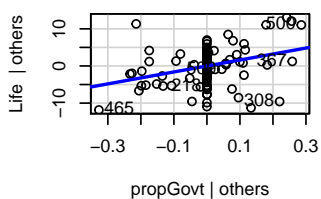
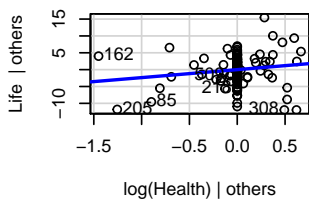
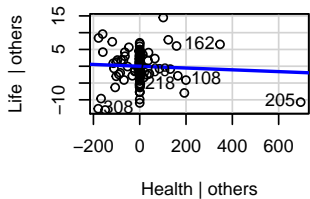


Added-Variable Plots

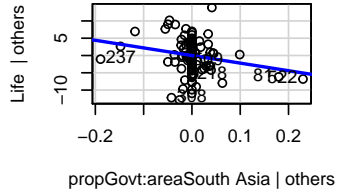
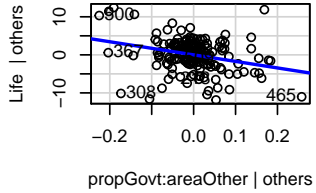


```
# were simple regression plots
```

```
avPlots(fit, id =list(n = 3))
```

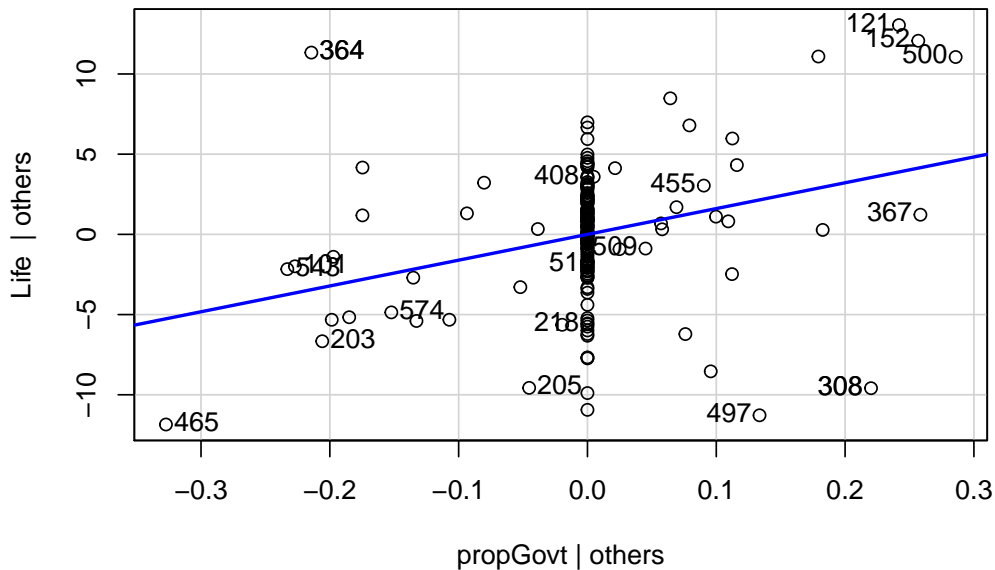



Added-Variable Plots



```
avPlot(fit, 'propGovt', id =list(n = 10))
```

Added-Variable Plot: propGovt



Interactively, you can use:

```
avPlot(fit, 'propGovt', id.method = "identify")
```

4.5.1 Visualize fit for diagnostics

In 3D

```
Plot3d( Life ~ Health + propGovt | area, ds)
```

```
|           area   col
|  1     Africa  blue
|  2     Other   green
|  3 South Asia orange
```

```
| Use left mouse to rotate, middle mouse (or scroll) to zoom, right
```

```
Fit3d( fit , resid = T)
```

```
| Warning in log(Health): NaNs produced
```

```
# Id3d() # outliers?
```

2D

```
summary(ds)
```

```
|          country          iso3      region  HealthExpPC.G
| Afghanistan      : 1  AFG      : 1  AFR :46  Min.    :  2.
| Albania          : 1  AGO      : 1  AMR :35  1st Qu.: 42.
| Algeria          : 1  ALB      : 1  EMR :22  Median  : 183.
| Andorra          : 1  AND      : 1  EUR :53  Mean    : 782.
| Angola           : 1  ARE      : 1  SEAR:11 3rd Qu.: 650.
| Antigua and Barbuda: 1  ARG      : 1  WPR :27  Max.    :7696.
| (Other)          :188  (Other):188      NA's    :3
| HealthExpPC.Tot.ppp HealthExpPC.Govt.ppp HealthExpPC.Tot.exch
| Min.    : 16.99      Min.    :  3.61      Min.    : 13.9
| 1st Qu.: 166.09     1st Qu.: 79.92     1st Qu.: 90.7
| Median  : 518.83     Median  : 286.72    Median  : 333.8
| Mean    :1114.08     Mean    : 770.12     Mean    :1094.2
```

	3rd Qu.:1333.26	3rd Qu.: 926.80	3rd Qu.: 971.4	
	Max. :8607.88	Max. :5794.45	Max. :9120.8	
	NA's :4	NA's :4	NA's :3	
	total	govt	private	sex
	Min. : 16.99	Min. : 3.61	Min. : 0.51	BTSX:194
	1st Qu.: 166.09	1st Qu.: 79.92	1st Qu.: 57.10	FMLE: 0
	Median : 518.83	Median : 286.72	Median : 187.50	MLE : 0
	Mean :1114.08	Mean : 770.12	Mean : 343.96	
	3rd Qu.:1333.26	3rd Qu.: 926.80	3rd Qu.: 475.42	
	Max. :8607.88	Max. :5794.45	Max. :4653.69	
	NA's :4	NA's :4	NA's :4	
	lifeexp.Birth	lifeexp.At60	smoking.tobacco.current	smoking
	Min. :47.00	Min. :11.00	Min. : 4.00	Min.
	1st Qu.:64.00	1st Qu.:17.00	1st Qu.:14.00	1st Qu.
	Median :72.50	Median :19.00	Median :23.00	Median
	Mean :70.01	Mean :19.36	Mean :22.63	Mean
	3rd Qu.:76.00	3rd Qu.:22.00	3rd Qu.:29.00	3rd Qu.
	Max. :83.00	Max. :26.00	Max. :57.00	Max.
			NA's :47	NA's

	smoking.cig.current	smoking.cig.daily	Pop.Total	Pop.M
	Min. : 4.0	Min. : 2.00	Min. : 1	Min.
	1st Qu.:12.0	1st Qu.: 8.25	1st Qu.: 1696	1st Qu.
	Median :21.0	Median :16.00	Median : 7790	Median
	Mean :21.2	Mean :17.22	Mean : 36360	Mean
	3rd Qu.:29.0	3rd Qu.:23.75	3rd Qu.: 24535	3rd Qu.
	Max. :57.0	Max. :55.00	Max. :1390000	Max.
	NA's :46	NA's :48		NA's
	Pop.pCntUnder15	Pop.pCntOver60	Pop.pCntAnnGrowth	consumption.c
	Min. :13.12	Min. : 0.81	Min. :-9.100	Min. : 9.
	1st Qu.:18.72	1st Qu.: 5.20	1st Qu.: -2.300	1st Qu.: 179.
	Median :28.65	Median : 8.53	Median :-1.300	Median : 529.
	Mean :28.73	Mean :11.16	Mean :-1.453	Mean : 730.
	3rd Qu.:37.75	3rd Qu.:16.69	3rd Qu.: -0.500	3rd Qu.:1039.
	Max. :49.99	Max. :31.92	Max. : 0.800	Max. :2861.
				NA's :9
	hiv_prev15_49	LifeExp	LE	smoke
	Min. : 0.000	Min. :47.00	Min. :47.00	Min. : 9.0
	1st Qu.: 0.200	1st Qu.:64.00	1st Qu.:64.00	1st Qu.: 179.0

Median	: 0.400	Median	:72.50	Median	:72.50	Median	: 529.0
Mean	: 1.817	Mean	:70.01	Mean	:70.01	Mean	: 730.1
3rd Qu.	: 1.200	3rd Qu.	:76.00	3rd Qu.	:76.00	3rd Qu.	:1039.0
Max.	:26.000	Max.	:83.00	Max.	:83.00	Max.	:2861.0
NA's	:40					NA's	:9

	HE		hiv		special		yq
Min.	: 16.99	Min.	: 0.000	Min.	:0.00000	Min.	:48
1st Qu.	: 166.09	1st Qu.	: 0.200	1st Qu.	:0.00000	1st Qu.	:60
Median	: 518.83	Median	: 0.400	Median	:0.00000	Median	:72
Mean	:1114.08	Mean	: 1.817	Mean	:0.01546	Mean	:69
3rd Qu.	:1333.26	3rd Qu.	: 1.200	3rd Qu.	:0.00000	3rd Qu.	:76
Max.	:8607.88	Max.	:26.000	Max.	:1.00000	Max.	:80
NA's	:4	NA's	:40			NA's	:42

	Life		Cigarettes		Health		area
Min.	:47.00	Min.	: 9.0	Min.	: 16.99	Africa	:
1st Qu.	:64.00	1st Qu.	: 179.0	1st Qu.	: 166.09	Other	:1
Median	:72.50	Median	: 529.0	Median	: 518.83	South Asia:	
Mean	:70.01	Mean	: 730.1	Mean	:1114.08		
3rd Qu.	:76.00	3rd Qu.	:1039.0	3rd Qu.	:1333.26		

	Max.	:83.00	Max.	:2861.0	Max.	:8607.88
			NA's	:9	NA's	:4
	propGovt					
	Min.	:0.1296				
	1st Qu.	:0.4561				
	Median	:0.6070				
	Mean	:0.5940				
	3rd Qu.	:0.7478				
	Max.	:0.9989				
	NA's	:4				

Create a prediction data frame with values for which you want to predict model with observed values for Health and area but controlling for propGovt

4.5.1.1 3 ways:

1. easiest: use data BUT need to control for propGovt
2. Generate cartesian product of values of predictors: but hard to generate conditional ranges
3. Create prediction data set with original data augmented by extra points

```
ds <- sortdf( ds, ~ Health)
pred1 <- rbind(ds,NA,ds,NA, ds,NA, ds,NA, ds, NA)
      # to set five values for predicted propGovt
pred1$propGovt <- rep(seq(.1,.9,by=.2), each = nrow(ds)+1)

pred1$Life.fit <- predict(fit, newdata = pred1)
```

4.5.1.2 1. add predicted values to data frame In 'panels':

```
xyplot(Life ~ Health | area, ds)
```

0 2000 4000 6000 8000

Africa

Other

South Asia

Life

80

70

60

50

0

2000

4000

6000

8000

0

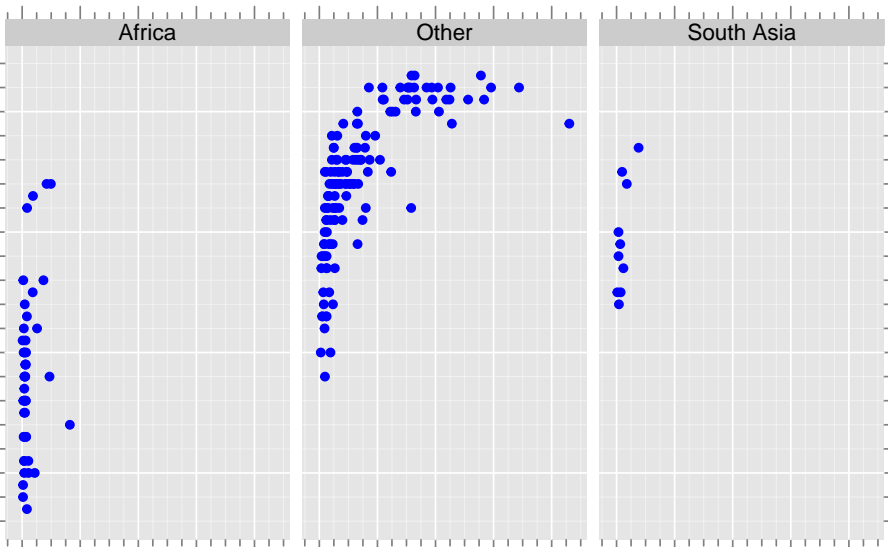
2000

4000

6000

8000

Health



```
gd() # ggplot2 look-alike
```

```
xyplot(Life ~ Health | area, ds)
```

0 2000 4000 6000 8000

Africa

Other

South Asia

Life

80

70

60

50

0

2000

4000

6000

8000

0

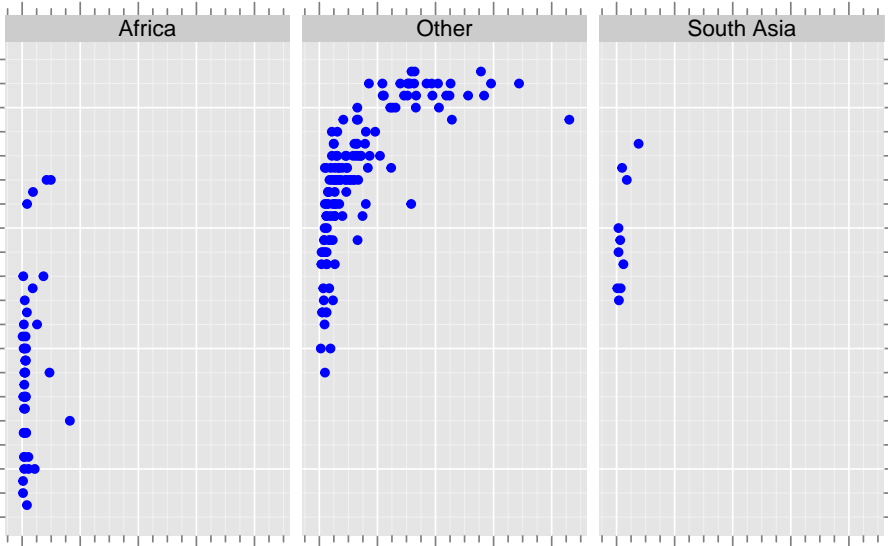
2000

4000

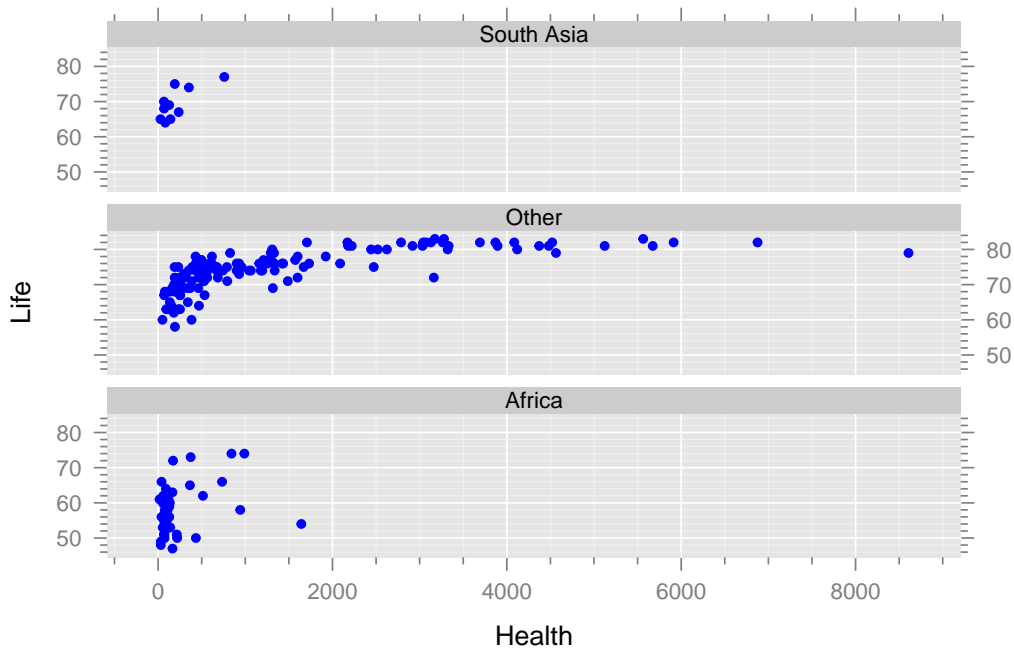
6000

8000

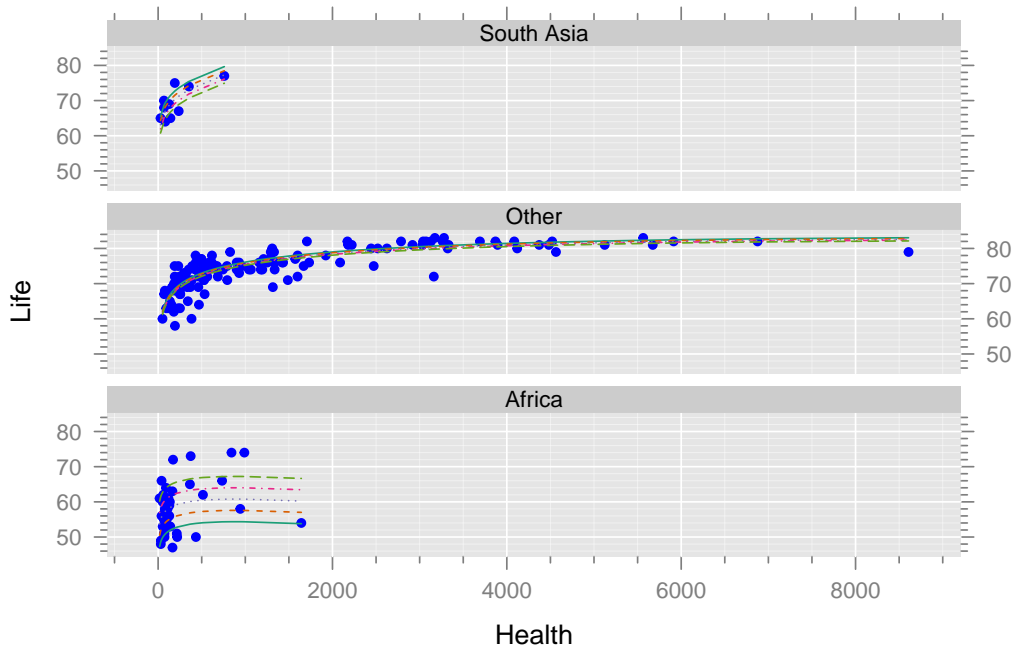
Health



```
xyplot(Life ~ Health | area, ds, layout = c(1,3))
```

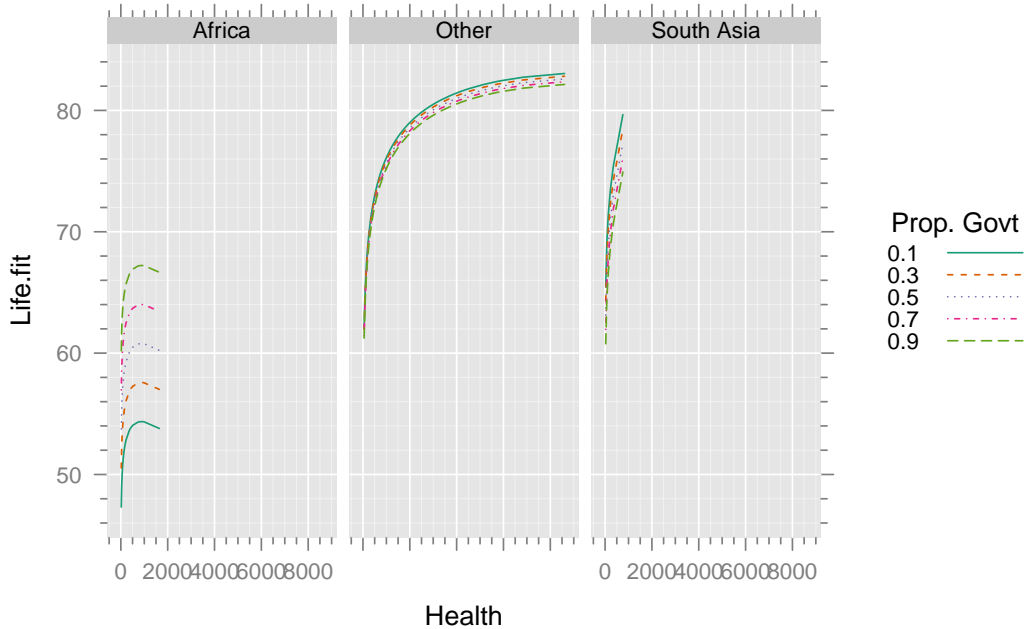



```
xyplot(Life ~ Health | area, ds, layout = c(1,3)) +  
  xyplot( Life.fit ~ Health | area, pred1, groups = propGovt,  
         type = 'l')
```



```
gd(lwd = 2)
(p <- xyplot(Life.fit ~ Health | area,
             pred1, groups = propGovt,
             type = 'l',
             auto.key = list(space='right',
                             lines = T, points = F,
                             title = 'Prop. Govt',
                             cex.title = 1)))
```

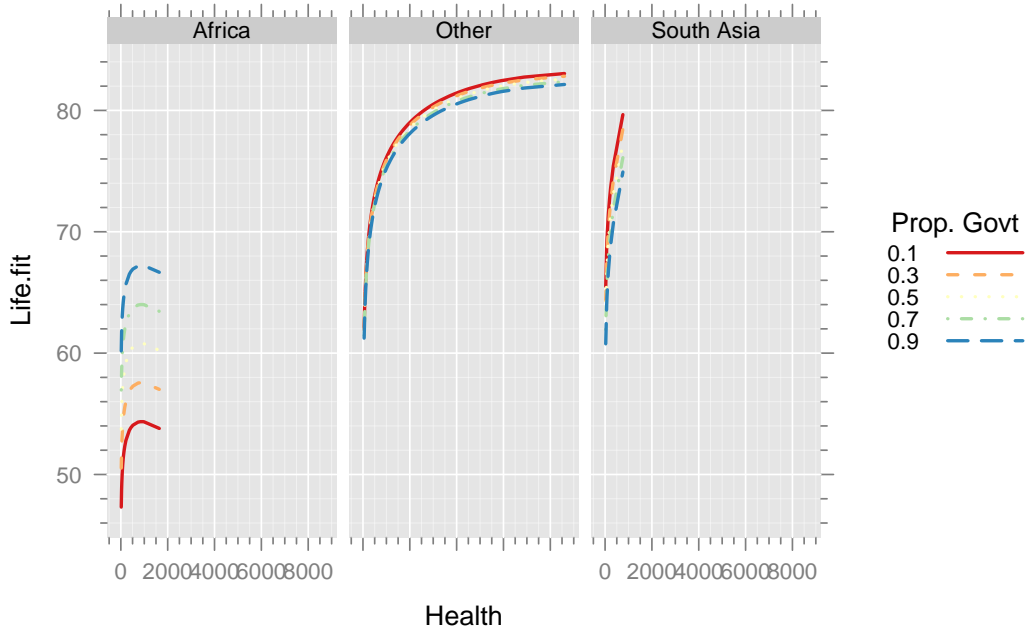
0 20004006008000



Some colour palettes you can choose from. `display.brewer.all()` Note that the first group consists of 'progressive' palettes, the second group of categorical palettes (one is 'paired') and the third group of 'bipolar' palettes. Note that yellow often doesn't work for lines that blend into the background so you might have to avoid palettes that include yellow for some purposes.

```
gd(lwd = 2, col = brewer.pal(5,"Spectral"))  
p # replots with new parameters
```

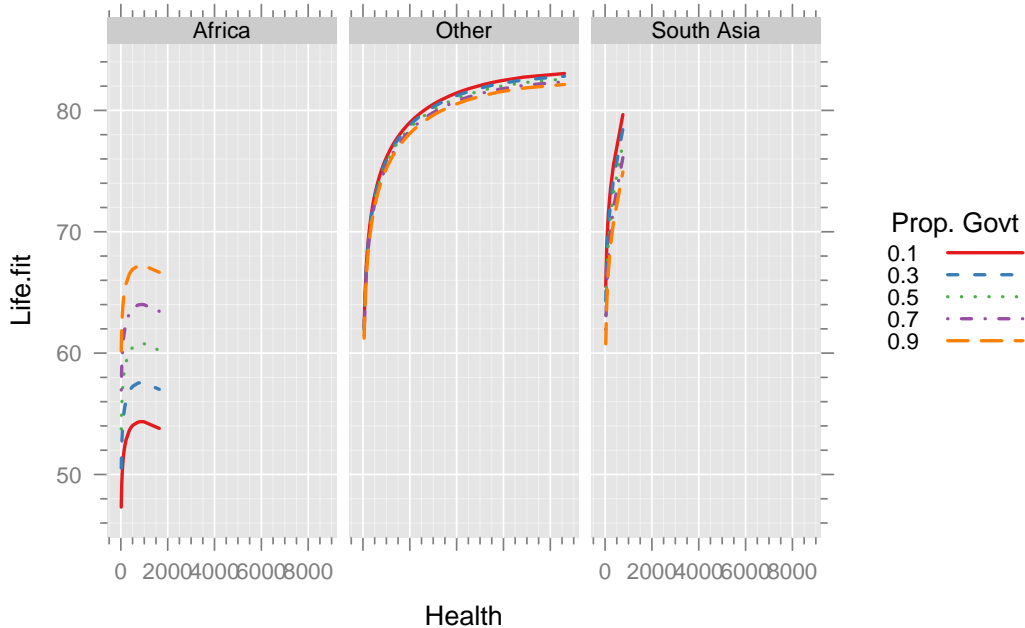
0 20004006008000



```
gd(lwd = 2, col = brewer.pal(5,"Set1"))
```

```
p
```

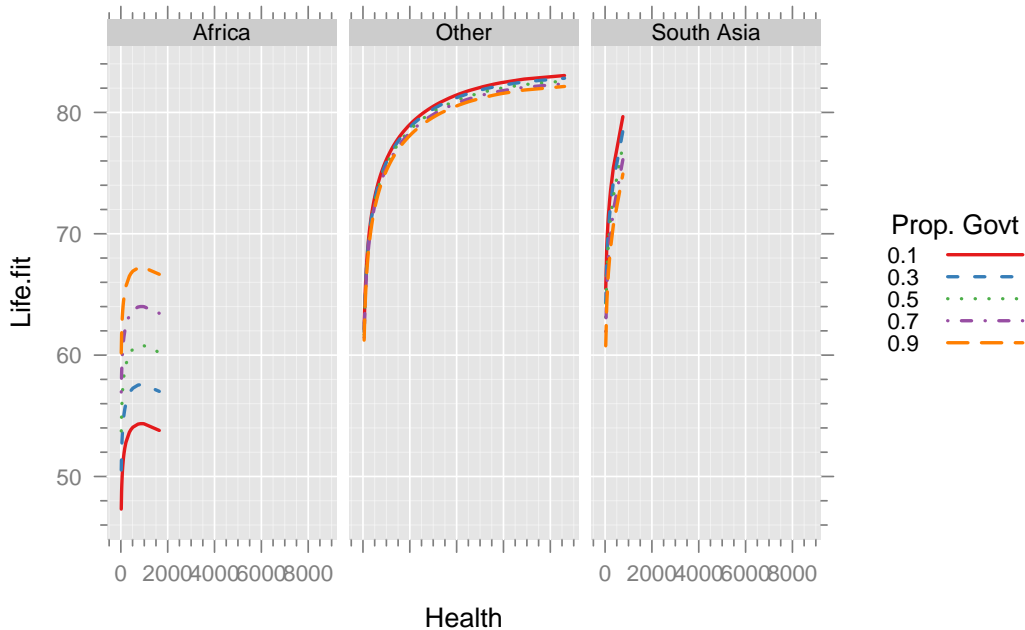
0 2000400060008000




```
gd(lty = 1)
```

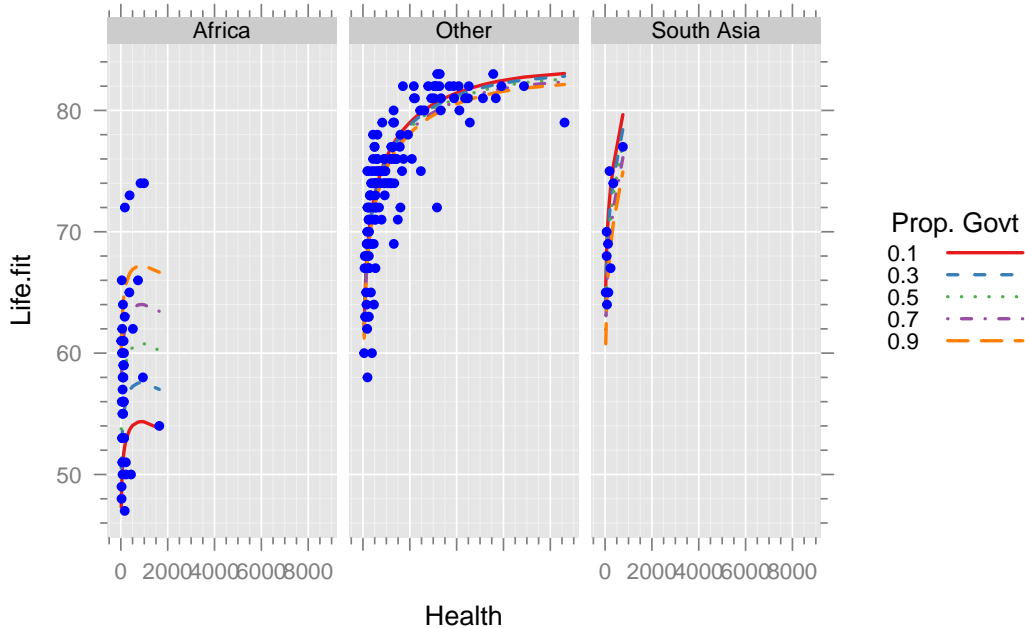
```
p
```

0 200040060008000



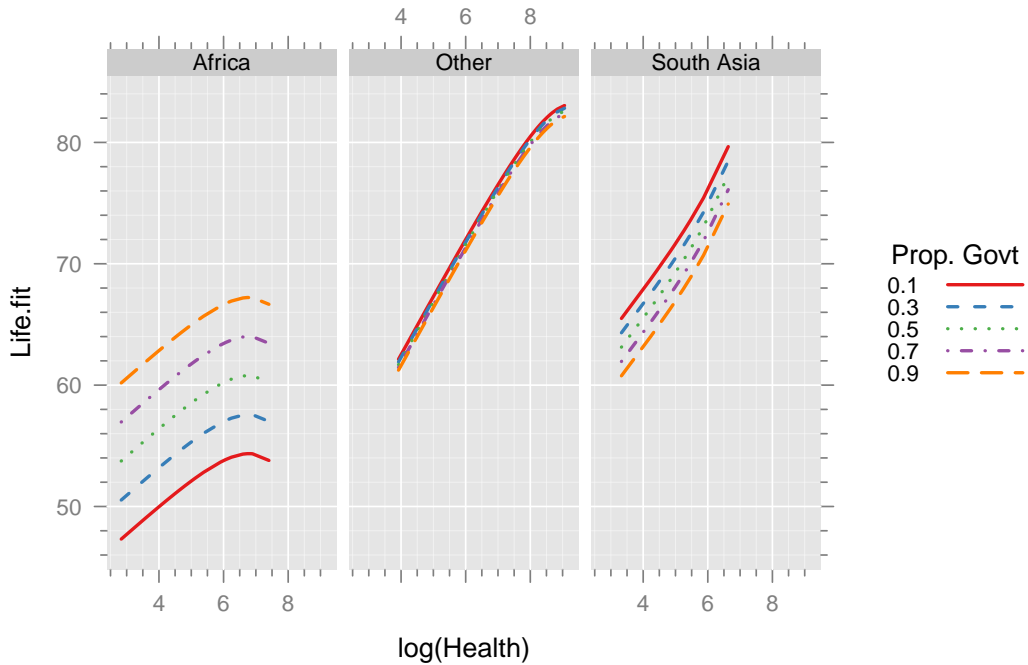
```
p + xyplot(Life ~ Health | area, ds)
```

0 20004006008000

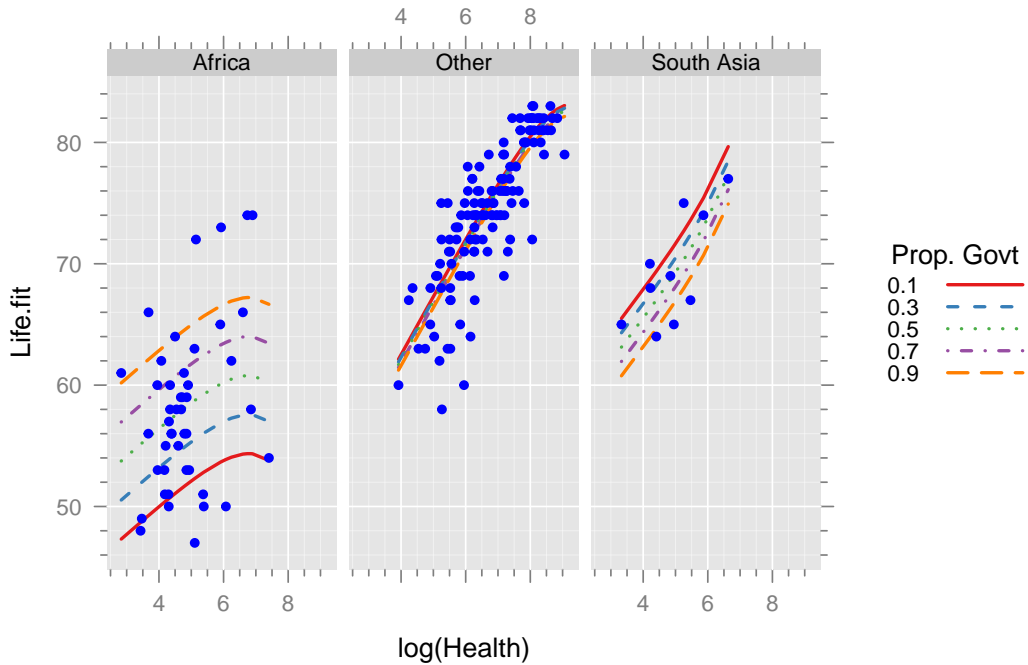


Probably better with $\log(\text{Health})$

```
(p <- xyplot(Life.fit ~ log(Health) | area,  
            pred1, groups = propGovt,  
            type = 'l',  
            auto.key = list(space = 'right', lines = T,  
                            points = F, title = 'Prop. Govt',  
                            cex.title = 1)))
```



```
p + xyplot(Life ~ log(Health) | area, ds)
```

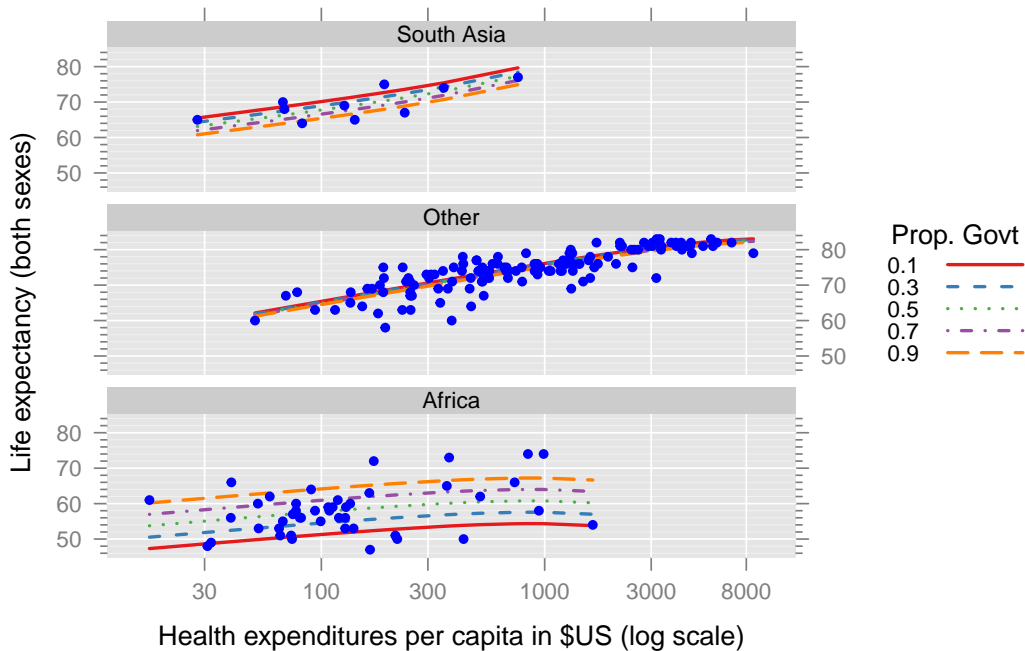


BUT NEVER NEVER USE axes that are not meaningful

Your work will be written off as incomprehensible!

Also: use meaningful labels

```
update(p,  
  xlab = "Health expenditures per capita in $US (log scale)",  
  ylab = "Life expectancy (both sexes)",  
  layout = c(1,3),  
  scales = list( x = list(  
    at = log(c(30,100,300,1000,3000,8000)),  
    labels = c(30,100,300,1000,3000,8000))) +  
  xyplot(Life ~ log(Health) | area, ds)
```



Dropping some observations: BEWARE

Two ways:

- 1) Drop from data set and refit
- 2) Add parameters for dummy variables for observations to drop and refit

Suppose we want to drop “Equatorial Guinea”

CAUTION: this needs good reflection BUT we often should approach this like a sensitivity analysis: e.g. “would it make a big difference if I dropped this point?”

```
ds$EqG <- 1*(ds$country == "Equatorial Guinea")

fit2 <- lm( Life ~
            (Health + log(Health) + propGovt) * area + EqG, ds,
            na.action = na.exclude)
```

OR

```
fit2 <- update(fit, . ~ . + EqG)
summary(fit2)
```

Call:

```
lm(formula = Life ~ Health + log(Health) + propGovt + area +  
    EqG + Health:area + log(Health):area + propGovt:area, data =  
    na.action = na.exclude)
```

Residuals:

Min	1Q	Median	3Q	Max
-13.3962	-1.8221	0.1869	2.1919	10.5391

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	50.031832	6.036711	8.288	2.78e-1
Health	0.012027	0.005459	2.203	0.02888
log(Health)	-0.386600	1.412966	-0.274	0.78470
propGovt	15.122468	3.735031	4.049	7.69e-0
areaOther	-6.674012	6.982150	-0.956	0.34044
areaSouth Asia	4.638695	15.383108	0.302	0.76335
EqG	-22.943658	6.239158	-3.677	0.00031

```
| Health:areaOther          -0.012481    0.005475   -2.279 0.02384
| Health:areaSouth Asia    -0.008027    0.015854   -0.506 0.61329
| log(Health):areaOther     5.209887    1.541017    3.381 0.00088
| log(Health):areaSouth Asia 3.784089    4.086121    0.926 0.35566
| propGovt:areaOther        -16.250469    4.206623   -3.863 0.00015
| propGovt:areaSouth Asia  -21.032953    8.491622   -2.477 0.01419
```

```
| ---
```

```
| Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
| Residual standard error: 3.873 on 177 degrees of freedom
| (4 observations deleted due to missingness)
```

```
| Multiple R-squared:  0.83, Adjusted R-squared:  0.8185
```

```
| F-statistic: 72.01 on 12 and 177 DF,  p-value: < 2.2e-16
```

```
Fit3d(fit2, other.vars = list( EqG = 0))
```

```
| Warning in log(Health): NaNs produced
```

```
spin(-2,18,10)
```

```
par3d(windowRect=c(10,10,700,700))
```

```
rgl.snapshot('dropEqG.png')
```

Figure X: Note the change in the fitted surface for Africa when Equatorial Guinea is dropped from the model.

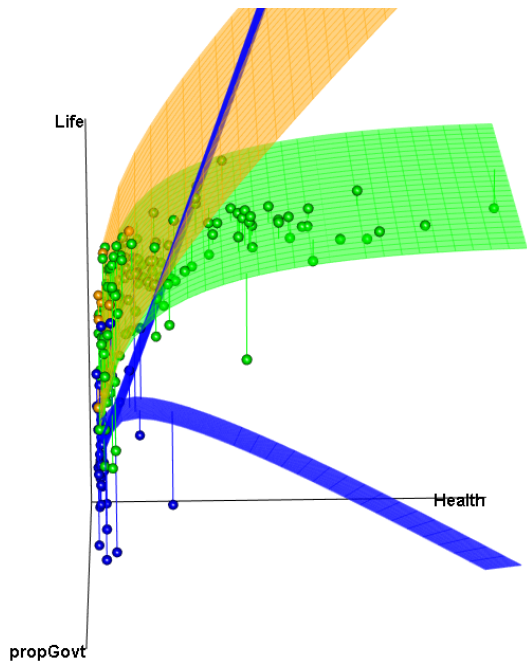
4.6 Asking questions:

4.6.1 Can we simplify the model?

Often this process focuses on the initial regression parameter and asks which ones ‘can we drop’? Often starting with highest order interactions and working in.

BEWARE THE PRINCIPLE OF MARGINALITY

In general (i.e. 99.9% of the time) DO NOT eliminate a term without also eliminating all higher-order terms to which the term is *marginal*. e.g. ‘Health’ is marginal to ‘Health:area’, and ‘Health:area’ would be marginal to ‘Health:propGovt:area’. Otherwise the resulting model loses invariance with respect to changes of origins of interacting variables.



Here's a model that seems to tell a different story but it's perfectly equivalent. The individual terms estimate different aspects of the model, but the models as a whole are equivalent and produce the same fitted values.

```
fit2.eq <- lm( Life ~
              area/(Health + log(Health) + propGovt) + EqG -1,
              ds, na.action = na.exclude)
anova(fit2, fit2.eq)
```

```
| Analysis of Variance Table
```

```
| Model 1: Life ~ Health + log(Health) + propGovt + area + EqG + H
```

```
| log(Health):area + propGovt:area
```

```
| Model 2: Life ~ area/(Health + log(Health) + propGovt) + EqG - 1
```

```
| Res.Df    RSS Df    Sum of Sq F Pr(>F)
```

```
| 1      177 2655.1
```

```
| 2      177 2655.1  0 -5.2751e-11
```



```
AIC( fit2, fit2.eq)
```

```
|           df      AIC  
|  fit2      14 1068.266  
|  fit2.eq  14 1068.266
```

```
summary(fit2)
```

```
|  
| Call:  
| lm(formula = Life ~ Health + log(Health) + propGovt + area +  
|     EqG + Health:area + log(Health):area + propGovt:area, data =  
|     na.action = na.exclude)  
|  
| Residuals:  
|      Min      1Q  Median      3Q      Max  
| -13.3962 -1.8221  0.1869  2.1919 10.5391  
|  
| Coefficients:
```

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	50.031832	6.036711	8.288	2.78e-1
Health	0.012027	0.005459	2.203	0.02888
log(Health)	-0.386600	1.412966	-0.274	0.78470
propGovt	15.122468	3.735031	4.049	7.69e-0
areaOther	-6.674012	6.982150	-0.956	0.34044
areaSouth Asia	4.638695	15.383108	0.302	0.76335
EqG	-22.943658	6.239158	-3.677	0.00031
Health:areaOther	-0.012481	0.005475	-2.279	0.02384
Health:areaSouth Asia	-0.008027	0.015854	-0.506	0.61329
log(Health):areaOther	5.209887	1.541017	3.381	0.00088
log(Health):areaSouth Asia	3.784089	4.086121	0.926	0.35566
propGovt:areaOther	-16.250469	4.206623	-3.863	0.00015
propGovt:areaSouth Asia	-21.032953	8.491622	-2.477	0.01419

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 3.873 on 177 degrees of freedom
 (4 observations deleted due to missingness)

```
| Multiple R-squared: 0.83, Adjusted R-squared: 0.8185  
| F-statistic: 72.01 on 12 and 177 DF, p-value: < 2.2e-16
```

```
summary(fit2.eq)
```

```
|  
| Call:  
| lm(formula = Life ~ area/(Health + log(Health) + propGovt) +  
|     EqG - 1, data = ds, na.action = na.exclude)
```

```
| Residuals:  
|      Min      1Q  Median      3Q      Max  
| -13.3962 -1.8221  0.1869  2.1919 10.5391
```

```
| Coefficients:  
|  
|           Estimate Std. Error t value Pr(>|t|)  
| areaAfrica      5.003e+01  6.037e+00   8.288 2.78e-1  
| areaOther       4.336e+01  3.508e+00  12.358 < 2e-1  
| areaSouth Asia  5.467e+01  1.415e+01   3.864 0.00015
```

EqG	-2.294e+01	6.239e+00	-3.677	0.00031
areaAfrica:Health	1.203e-02	5.459e-03	2.203	0.02888
areaOther:Health	-4.536e-04	4.217e-04	-1.075	0.28363
areaSouth Asia:Health	4.000e-03	1.488e-02	0.269	0.78843
areaAfrica:log(Health)	-3.866e-01	1.413e+00	-0.274	0.78470
areaOther:log(Health)	4.823e+00	6.150e-01	7.842	4.00e-1
areaSouth Asia:log(Health)	3.397e+00	3.834e+00	0.886	0.37674
areaAfrica:propGovt	1.512e+01	3.735e+00	4.049	7.69e-0
areaOther:propGovt	-1.128e+00	1.935e+00	-0.583	0.56072
areaSouth Asia:propGovt	-5.910e+00	7.626e+00	-0.775	0.43935

| ---

| Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

| Residual standard error: 3.873 on 177 degrees of freedom
 | (4 observations deleted due to missingness)

| Multiple R-squared: 0.9972, Adjusted R-squared: 0.997

| F-statistic: 4880 on 13 and 177 DF, p-value: < 2.2e-16

Anova(fit2)

Anova Table (Type II tests)

Response: Life

	Sum Sq	Df	F value	Pr(>F)	
Health	12.01	1	0.8003	0.3722085	
log(Health)	763.67	1	50.9096	2.39e-11	***
propGovt	19.56	1	1.3039	0.2550477	
area	1845.55	2	61.5164	< 2.2e-16	***
EqG	202.85	1	13.5230	0.0003124	***
Health:area	79.23	2	2.6410	0.0741004	.
log(Health):area	171.81	2	5.7268	0.0038909	**
propGovt:area	240.45	2	8.0147	0.0004655	***
Residuals	2655.08	177			

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Anova(fit2.eq)

| Anova Table (Type II tests)

| Response: Life

	Sum Sq	Df	F value	Pr(>F)	
area	944894	3	20997.0169	< 2.2e-16	***
EqG	203	1	13.5230	0.0003124	***
area:Health	91	3	2.0274	0.1117604	
area:log(Health)	935	3	20.7877	1.385e-11	***
area:propGovt	260	3	5.7778	0.0008611	***
Residuals	2655	177			

| ---

| Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

wald(fit2, "th:|th):")

	numDF	denDF	F.value	p.value
th: th):	4	177	3.143431	0.01582

	Estimate	Std.Error	DF	t-value	p-value
Health:areaOther	-0.012481	0.005475	177	-2.279349	0.0241
Health:areaSouth Asia	-0.008027	0.015854	177	-0.506274	0.6137
log(Health):areaOther	5.209887	1.541017	177	3.380811	0.0008
log(Health):areaSouth Asia	3.784089	4.086121	177	0.926083	0.3521
	Upper 0.95				
Health:areaOther	-0.001675				
Health:areaSouth Asia	0.023261				
log(Health):areaOther	8.251017				
log(Health):areaSouth Asia	11.847874				

Question: Under what conditions would two seemingly different models produce exactly the same fit (i.e. predicted values of Y)?

4.6.2 Asking specific questions

What question does each coefficient answer and how can we get answers to the questions we want?

PRINCIPLE

Except with very simple models, raw regression output generally answers few meaningful questions AND most important questions are rarely answered by raw regression output

Interpreting β s:

Each term involving 'area' is a comparison with the **REFERENCE LEVEL** when **ALL VARIABLES IN HIGHER ORDER INTERACTING TERMS** are set to 0.

Each term involving 'area' is a comparison with the **REFERENCE LEVEL** (Africa because it's the level that isn't showing) when **ALL VARIABLES IN HIGHER ORDER INTERACTING TERMS** are set to 0.

The model is:

$$\begin{aligned}
Y &= \beta_0 + \beta_1 Health + \beta_2 \ln(Health) + \beta_3 propGovt \\
&+ \beta_4 area_{Other} + \beta_5 area_{SouthAsia} \\
&+ \beta_6 EqG \\
&+ \beta_7 Health \times area_{Other} + \beta_8 Health \times area_{SouthAsia} \\
&+ \beta_9 \ln(Health) \times area_{Other} + \beta_{10} \ln(Health) \times area_{SouthAsia} \\
&+ \beta_{11} propGovt \times area_{Other} + \beta_{12} propGovt \times area_{SouthAsia} \\
&+ \varepsilon
\end{aligned}$$

where $\varepsilon \sim N(0, \sigma^2)$ independently of predictors.

Note: Here we encounter the vital difference between assumptions that can be checked and assumptions that can't be checked. We are assuming that

- 1) errors are normal,
- 2) they have the same variance for each observation, and
- 3) they are independent of predictors.

The first two assumptions can be checked with diagnostics, the third, in general,

cannot. In econometrics it's recognized as a key assumption related to the 'exogeneity' of the predictors and the causal interpretation of the model. For a further treatment of this question see Murnane and Willett (2010) and Pearl and Mackenzie (2018).

4.7 Understanding coefficients

Q: What does β_1 mean?

A: It's the expected change in Y when you change *Health* by one unit keeping all other terms constant — **BUT THAT'S IMPOSSIBLE**

We'll have more luck with β_3 : It's the expected change in Y when you change *progGovt* by one unit keeping all other terms constant, i.e. when all variables that interact with *propGovt* are equal to 0.

i.e. when $area_{Other} = area_{SouthAsia} = 0$

i.e. **in Africa**

So we have a clear interpretation: it's the expected change in Life Expectancy

using our model to compare a hypothetical country where health expenditures are entirely supported by the government with a hypothetical country in which they are entirely private **in Africa**.

This is not obvious to a casual user of regression and most surely not to most clients and readers of academic journals.

4.7.1 How can we get answers to meaningful questions?

You're in luck. Calculus comes in handy.

What's the 'effect' (a very misused word and I'm still looking for a better one) of increasing Health expenditures by 1 dollar?

$$\begin{aligned}
\frac{\partial E(Y)}{\partial Health} &= 0 \times \beta_0 + \beta_1 + \beta_2 \frac{1}{Health} + 0\beta_3 \\
&+ 0 \times \beta_4 + 0 \times \beta_5 \\
&+ 0 \times \beta_6 \\
&+ \beta_7 area_{Other} + \beta_8 area_{SouthAsia} \\
&+ \beta_9 \frac{area_{Other}}{Health} + \beta_{10} \frac{area_{SouthAsia}}{Health} \\
&+ 0 \times \beta_{11} + 0 \times \beta_{12} \\
&= L\beta
\end{aligned}$$

where

$$L = \begin{bmatrix} 0 & 1 & \frac{1}{Health} & 0 & 0 & 0 & 0 & area_{Other} & area_{SouthAsia} & \frac{area_{Other}}{Health} & \frac{area_{SouthAsia}}{Health} & 0 \end{bmatrix}$$

and

$$\beta = \begin{bmatrix} \beta_0 \\ \beta_1 \\ \beta_2 \\ \beta_3 \\ \beta_4 \\ \beta_5 \\ \beta_6 \\ \beta_7 \\ \beta_8 \\ \beta_9 \\ \beta_{10} \\ \beta_{11} \\ \beta_{12} \end{bmatrix}$$

which we can estimate with

$$\hat{\eta} = L\hat{\beta}$$

NOTE: This does not seem to depend on *propGovt*! Is this reasonable? What should we do about that?

Exercise: Explore what could be done with *propGovt*.

$\hat{\beta}$ is obtained with:

```
coef(fit2)
```

```
|          (Intercept)                Health
|          50.031832427                0.012026938
|          log(Health)                propGovt
|          -0.386599991                15.122467573
|          areaOther                areaSouth Asia
|          -6.674012321                4.638695485
|          EqG                Health:areaOther
|          -22.943657897                -0.012480505
|          Health:areaSouth Asia    log(Health):areaOther
|          -0.008026658                5.209886574
|          log(Health):areaSouth Asia    propGovt:areaOther
|          3.784089239                -16.250468580
|          propGovt:areaSouth Asia
|          -21.032953424
```



```
|           Estimate Std.Error DF   t-value  p-value Lower 0.95 Upper 0.
| [1,] 0.047779 0.005796 177 8.243213 <.00001 0.036341 0.059218
```

How could we mass produce this?

```
ex <- expression( cbind( 0,1,1/Health,0,0,0,0,
                          area == "Other", area == "South Asia",
                          (area == "Other")/Health,
                          (area == "South Asia")/Health,
                          0,0))
```

ex

```
| expression(cbind(0, 1, 1/Health, 0, 0, 0, 0, area == "Other",
|           area == "South Asia", (area == "Other")/Health, (area ==
|           "South Asia")/Health, 0, 0))
```

```
with( list(Health=100, area = "South Asia"), eval(ex))
```

```
|           [,1] [,2] [,3] [,4] [,5] [,6] [,7] [,8] [,9] [,10] [,11] [,
| [1,]      0    1 0.01    0    0    0    0    0    1    0    0.01
```



```
pred <- expand.grid( Health = seq(30,4000,10),  
                    area = levels(ds$area))
```

```
head(pred)
```

```
|      Health  area  
|    1      30 Africa  
|    2      40 Africa  
|    3      50 Africa  
|    4      60 Africa  
|    5      70 Africa  
|    6      80 Africa
```

```
tail(pred)
```

```
|           Health      area  
|   1189    3950 South Asia  
|   1190    3960 South Asia  
|   1191    3970 South Asia  
|   1192    3980 South Asia
```

```
| 1193 3990 South Asia
| 1194 4000 South Asia
```

```
dim(pred)
```

```
| [1] 1194 2
```

```
head(with(pred, eval(ex)))
```

```
|      [,1] [,2]      [,3] [,4] [,5] [,6] [,7] [,8] [,9] [,10] [,11]
| [1,] 0    1 0.03333333 0    0    0    0    0    0    0
| [2,] 0    1 0.02500000 0    0    0    0    0    0    0
| [3,] 0    1 0.02000000 0    0    0    0    0    0    0
| [4,] 0    1 0.01666667 0    0    0    0    0    0    0
| [5,] 0    1 0.01428571 0    0    0    0    0    0    0
| [6,] 0    1 0.01250000 0    0    0    0    0    0    0
```

```
ww <- wald(fit2, with(pred, eval(ex)))
str(ww)
```

```
| List of 1
```

```
| $ :List of 7
| ..$ anova :List of 4
| .. ..$ numDF : int 6
| .. ..$ denDF : int 177
| .. ..$ F-value: num [1, 1] 33.2
| .. ..$ p-value: num [1, 1] 1.3e-26
| ..$ estimate:Classes 'data.frame.lab' and 'data.frame': 1194
| .. ..$ Estimate : num [1:1194] -0.00086 0.00236 0.00429 0.005
| .. ..$ Std.Error : num [1:1194] 0.0424 0.0306 0.0236 0.0189 0.
| .. ..$ DF : num [1:1194] 177 177 177 177 177 177 177 17
| .. ..$ t-value : num [1:1194] -0.0203 0.0772 0.1822 0.2954 0
| .. ..$ p-value : num [1:1194] 0.984 0.939 0.856 0.768 0.677
| .. ..$ Lower 0.95: num [1:1194] -0.0844 -0.058 -0.0422 -0.0317
| .. ..$ Upper 0.95: num [1:1194] 0.0827 0.0628 0.0508 0.0429 0.
| .. ..- attr(*, "labs")= chr [1:2] "" ""
| ..$ coef : num [1:1194] -0.00086 0.00236 0.00429 0.00558 0.
| ..$ L : num [1:1194, 1:13] 0 0 0 0 0 0 0 0 0 0 0 ...
| ..$ se : num [1:1194] 0.0424 0.0306 0.0236 0.0189 0.0156
| ..$ L.full : num [1:6, 1:13] 0 0 0 0 0 ...
```

```
| ..$ L.rank : int 6  
| - attr(*, "class")= chr "wald"
```

```
head(as.data.frame(ww))
```

```
|          coef          se          U2          L2  
| 1 -0.0008597286 0.04235567 0.08385161 -0.08557107  
| 2  0.0023619380 0.03061080 0.06358353 -0.05885966  
| 3  0.0042949379 0.02357817 0.05145127 -0.04286140  
| 4  0.0055836046 0.01890309 0.04338979 -0.03222258  
| 5  0.0065040807 0.01557660 0.03765728 -0.02464912  
| 6  0.0071944378 0.01309439 0.03338321 -0.01899433
```

```
pred <- cbind( pred, as.data.frame(ww))
```

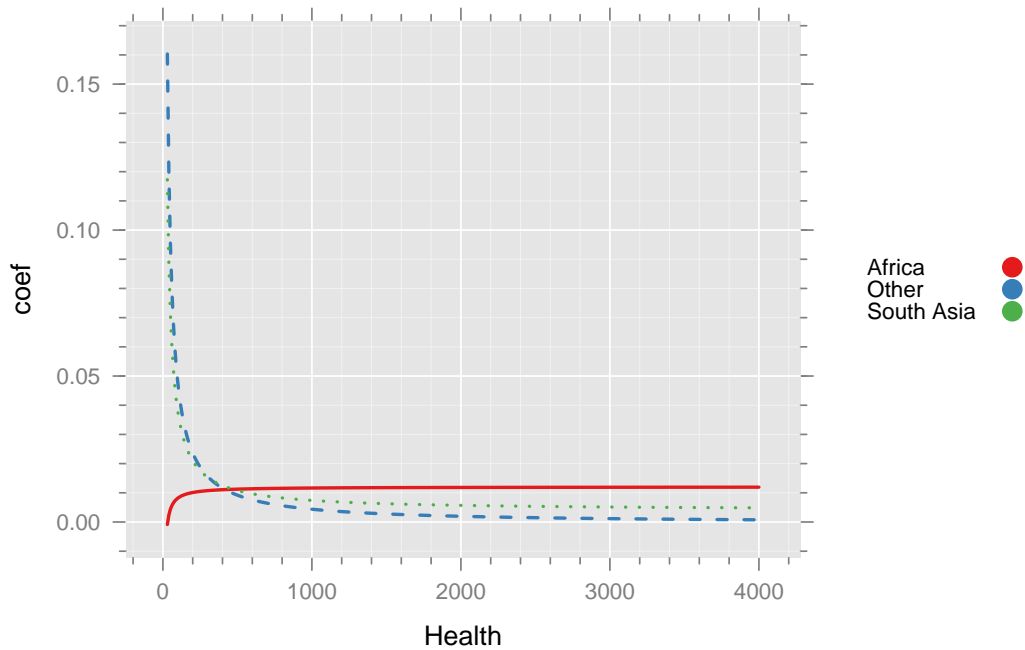
```
head(pred)
```

```
| Health area          coef          se          U2          L2  
| 1      30 Africa -0.0008597286 0.04235567 0.08385161 -0.08557107  
| 2      40 Africa  0.0023619380 0.03061080 0.06358353 -0.05885966  
| 3      50 Africa  0.0042949379 0.02357817 0.05145127 -0.04286140
```

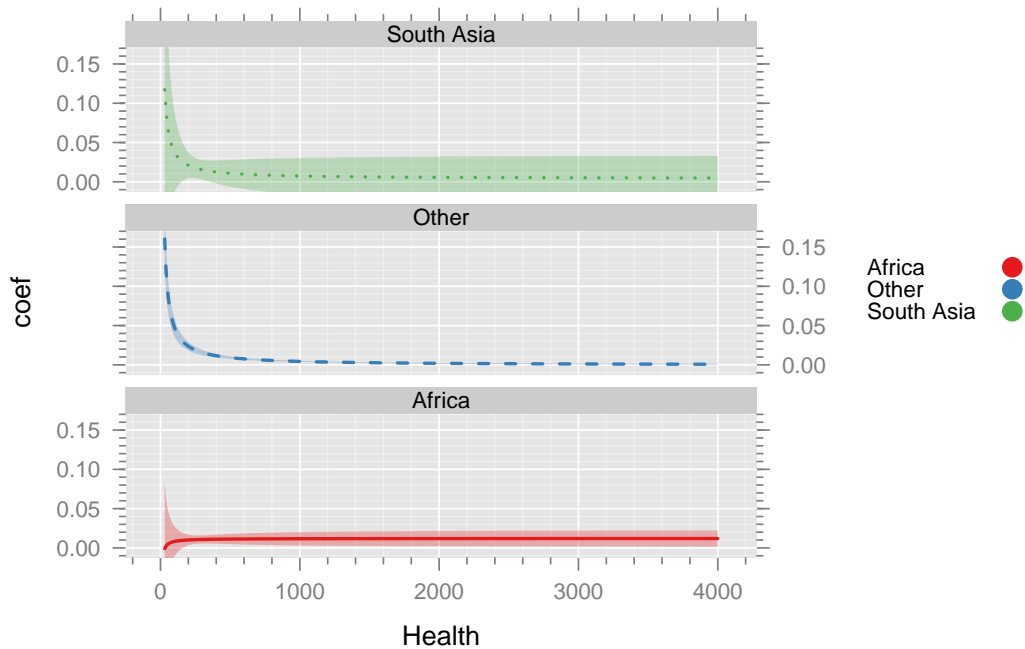
	4	60	Africa	0.0055836046	0.01890309	0.04338979	-0.03222258
	5	70	Africa	0.0065040807	0.01557660	0.03765728	-0.02464912
	6	80	Africa	0.0071944378	0.01309439	0.03338321	-0.01899433

```
gd(lwd=2)
```

```
xyplot(coef ~ Health, pred, groups = area, type = 'l',  
        auto.key=list(space = 'right'))
```

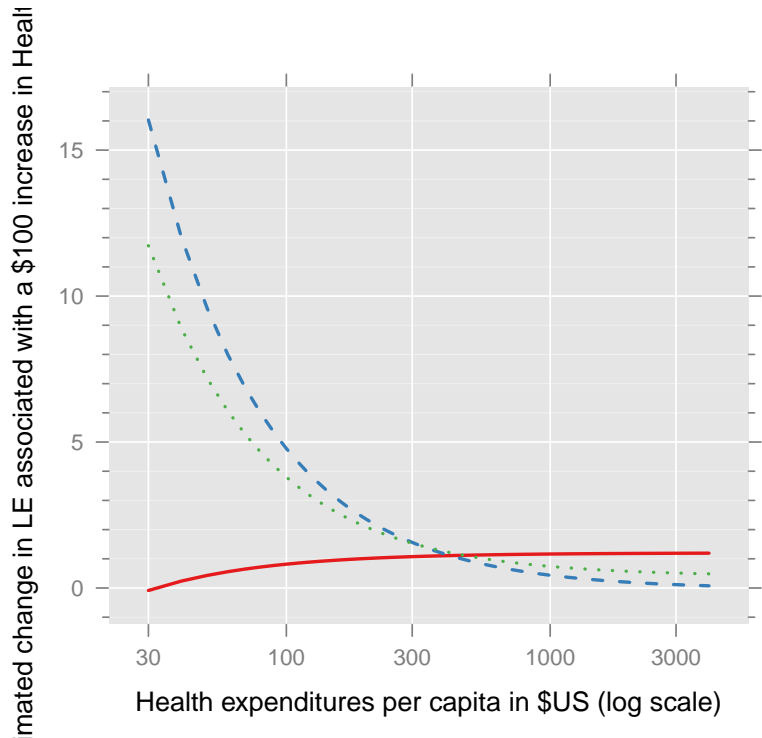


```
xyplot(coef ~ Health|area, pred, groups = area, type = 'l',
       auto.key = list(space='right'),
       subscripts = T,
       lower = pred$coef - 2* pred$se,
       upper = pred$coef + 2* pred$se,
       layout = c(1,3)) +
glayer( gpanel.fit(...))
```



Make labels and axes interpretable for presentation:

```
xyplot(I(100*coef) ~ log(Health), pred, groups = area, type = 'l',
       auto.key = list(space = 'right', lines = T, points = F),
       ylab =
"Estimated change in LE associated with a $100 increase in Health Ex
       xlab = "Health expenditures per capita in $US (log scale)",
       scales = list(x = list(
         at = log(c(30, 100, 300, 1000, 3000, 8000)),
         labels = c(30, 100, 300, 1000, 3000, 8000))))
```



Africa
Other
South Asia

Limit plot to ranges in each Area:

1. create a small data set with ranges

```
dsr <- ds
```

1. max within each area

```
dsr$max <- with(dsr, capply( Health, area, max, na.rm = T))
```

2. min within each area

```
dsr$min <- with(dsr, capply( Health, area, min, na.rm = T))
```

3. summary data frame with variables that are 'area invariant'

```
dsr <- up(dsr, ~ area) # keeps 'area' invariant variables only  
dsr
```

	sex	area	max	min
Africa	BTSX	Africa	1642.71	16.99
Other	BTSX	Other	8607.88	50.47
South Asia	BTSX	South Asia	759.66	27.86

4. merge back into pred

```
predr <- merge(pred, dsr[,c('area', 'max', 'min')], all.x = T)
```

5. keep values of Health that are within range

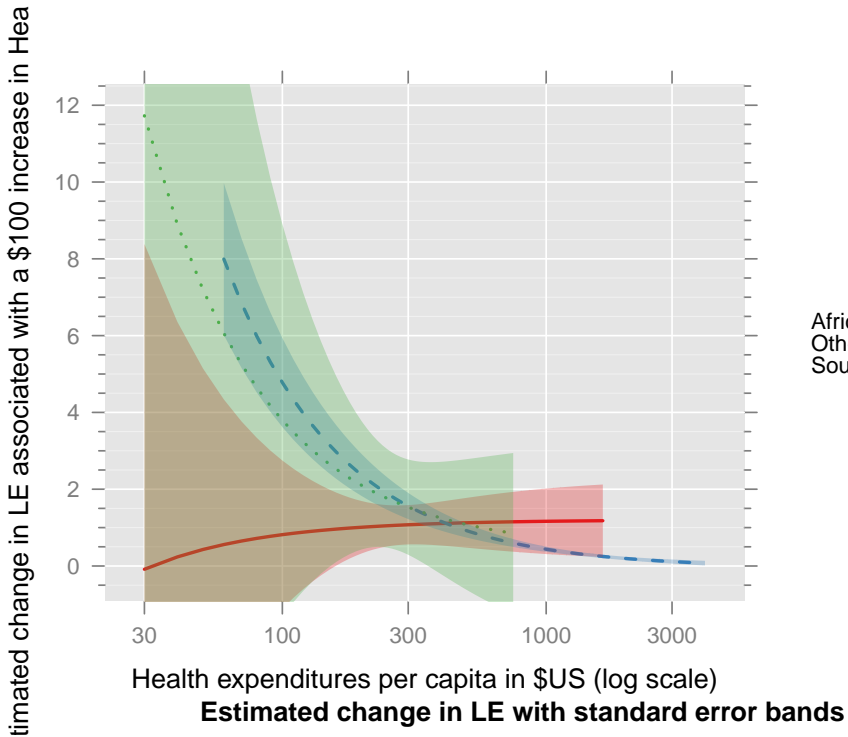
```
predr <- subset( predr, (Health <= max) & ( Health >= min))  
head(predr)
```

	area	Health	coef	se	U2	L2	
	1	Africa	30	-0.0008597286	0.04235567	0.08385161	-0.08557107
	2	Africa	40	0.0023619380	0.03061080	0.06358353	-0.05885966
	3	Africa	50	0.0042949379	0.02357817	0.05145127	-0.04286140
	4	Africa	60	0.0055836046	0.01890309	0.04338979	-0.03222258
	5	Africa	70	0.0065040807	0.01557660	0.03765728	-0.02464912
	6	Africa	80	0.0071944378	0.01309439	0.03338321	-0.01899433

4.7.2 Plotting fitted values and bands

If you define arguments **fit**, **lower** and **upper** in 'xyplot', they will be available to 'gpanel.fit' to draw the fitted line and confidence or predictions bands.

```
xyplot(I(100*coef) ~ log(Health), predr, groups = area, type = 'l',
       auto.key = list(space = 'right', lines = T, points = F),
       ylab =
"Estimated change in LE associated with a $100 increase in Health Ex
lower = 100*(predr$coef - 2* predr$se),
upper = 100*(predr$coef + 2* predr$se),
sub = "Estimated change in LE with standard error bands",
xlab = "Health expenditures per capita in $US (log scale)",
scales = list(x = list(
  at = log(c(30,100,300,1000,3000,8000)),
  labels = c(30,100,300,1000,3000,8000)))) +
glayer(gpanel.fit(...))
```



```
xyplot(I(100*coef) ~ log(Health)|area, predr, groups = area,
       type = 'l',
       auto.key=list(space='right', lines = T, points = F),
       ylab = "Additional years of Life Expectancy",
       xlim = c(-5,15),
       layout = c(1,3),
       lower = 100*(predr$coef - predr$se),
       upper = 100*(predr$coef + predr$se),
       xlab = "Health expenditures per capita in $US (log scale)",
       scales = list( x = list(
         at = log(c(30,100,300,1000,3000,8000)),
         labels = c(30,100,300,1000,3000,8000)))) +
glayer(gpanel.fit(...))
```

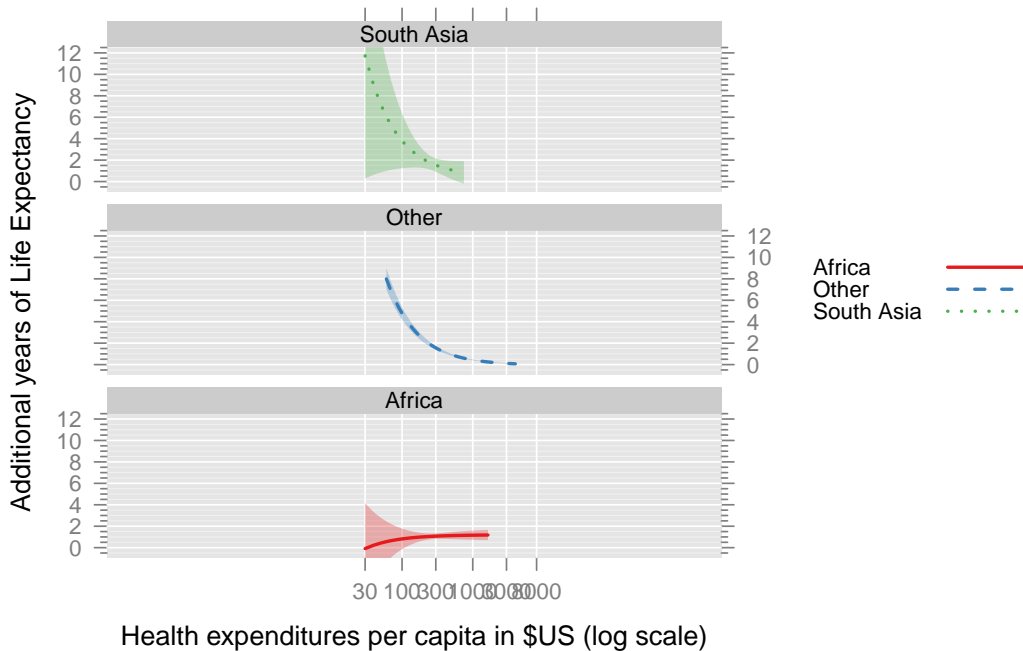


Figure: Additional years of life expectancy associated with a \$100 (U.S.) increase in health expenditures per capita per year, in three world regions as a function of the current level of health expenditures. The bands show the standard error of estimation.

4.7.2.1 Exercises

1. Redo the above plot showing relationship of LE with a 1% increase in Health Expenditures.
2. What happens if you introduce the possibility of interaction between health expenditures and *propGovt*?
3. How do things change if we do a regression that gives more weight to larger countries? How should we do this?
4. Compare Africa and South Asia: Is there evidence of a difference between LE adjusted for Health Expenditures and *propGovt*. Prepare an appropriate plot.
5. Same for South Asia and “Other”.

4.7.3 In the future:

We will explore the ‘Lfx’ function in ‘spida2’ and the ‘sc’ function for generalized splines generated by the ‘gsp’ function.

5 Appendices

5.1 Notes on the Principle of Marginality

The principle of marginality (POM) came up in class recently and it occurred to me that there might be confusion arising from the distinction between the requirements for:

- 1) A model to satisfy the POM, and
- 2) A null hypothesis specified by setting a set of terms to zero to satisfy the POM.

In a linear model with main effects and interactions of various orders, a model satisfies the POM if, for any interaction in the model, all included lower-order

interactions and main effects are also in the model. In other words, the model is closed under taking *margins* where we think of A:B as a margin of A:B:C, for example. Note that the intercept is considered a 0-th order effect and must be included if anything else is included.

A hypothesis that sets some parameters to 0 in a model that satisfies the POM, itself satisfies the POM provided the resulting H_0 model also satisfies the POM.

That's why the requirements for the set of terms that are set to 0 seems to be the reverse of the requirements for a model. For any given term set to zero in the hypothesis, all higher-order terms in the model that include the given term must also be set to zero. Otherwise, the null model would include those higher-order terms without including the given term which would result in a null hypothesis that violates the POM.

Thus the requirement for the set of terms set to zero is the 'reverse' of the requirement for models. The set of terms set to zero must be closed under taking interactions that are in the full model.

Note that the main significance of the POM is that predicted values (\hat{Y}) for a model that satisfies the POM are invariant under location-scale transformations

of numerical variables and non-singular recodings of categorical variables.

Hypotheses that satisfy the POM result in a test statistic, hence p-value, that is invariant under those same transformations and recodings.

So, if you stick to models and hypotheses that satisfy the POM, you don't need to worry that your conclusions would have been different if you had measured temperature in degrees Celsius instead of Fahrenheit or whether you had used a different category as a reference level for a factor.

Wald tests performed by specifying a regular expression to be matched in a model's terms will usually satisfy the POM because, if a regular expression matches a term, it will also match higher-order terms that contain that term.

This is true for tests of interactions with e.g. `wald(fit, ":")` for all interactions or `wald(fit, ':.*:')` for two-way and higher-order interactions, and for test of any particular effect, e.g. `wald(fit, 'X')` provided one checks that the regular expression does not match unintended terms.

Of course, you will be interested in estimating many parameters whose values do depend on the units used and the reference level. But those will be parameters

addressing specific questions. For example in a model $Y \sim X * G$ where X is continuous and G has three levels: A, B and C, you may want to estimate the difference in the rate of change with respect to X (which does depend on the units of X) comparing levels C and B. The exact hypothesis to test whether this is a particular value will not satisfy the POM because the estimate will depend on the units of X and the coding of the factor G . This is okay because it obeys the principle of “you know what you are doing”. The general rule is:

Never violate the POM unless you know what you are doing!

References

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