

# 432 Class 12 Slides

[github.com/THOMASELOVE/432-2018](https://github.com/THOMASELOVE/432-2018)

2018-02-22

# Setup

```
library(skimr)
library(pROC)
library(ROCR)
library(rms) # note: also loads Hmisc
library(simputation)
library(broom)
library(tidyverse)
```

# Today's Materials

- Logistic Regression and the Framingham Study (conclusion)
- Performing Linear Regression with `ols`
- Hormone Therapy and Baseline LDL in the HERS trial

The HERS trial is described in Vittinghoff et al., especially Chapter 4.

# Logistic Regression and Framingham

# Data Ingest, Cleanup (from Class 10)

```
fram <- read.csv("data/fram_new.csv") %>% tbl_df
set.seed(432001)
fram1 <- fram %>%
  impute_pmm(educ + cigs_day + heart_r ~ age + smoker) %>%
  impute_rlm(bmi + tot_chol ~ sex + age + sbp + heart_r) %>%
  impute_pmm(bp_meds ~ hx_htn + bmi + tot_chol) %>%
  impute_rlm(glucose ~ hx_dm + bmi + tot_chol + age) %>%
  mutate(ed_f = fct_recode(factor(educ),
    "1_Some_HS" = "1", "2_HS_grad" = "2",
    "3_Some_Col" = "3", "4_Col_grad" = "4"))
fram2 <- fram1 %>%
  select(subj, sex, age, smoker, cigs_day, bp_meds,
    hx_stroke, hx_htn, hx_dm, ed_f, tot_chol,
    sbp, dbp, bmi, heart_r, glucose, CHD_10)
```

# The Models We've Fit (predicting CHD\_10)

```
m_01 <- glm(CHD_10 ~ hx_htn, data = fram2,
            family = binomial)

d <- datadist(fram2)
options(datadist = "d")
m_01_lrm <- lrm(CHD_10 ~ hx_htn, data = fram2, x = T, y = T)

m_02 <- glm(CHD_10 ~ hx_htn + tot_chol,
            data = fram2, family = binomial)
m_02_lrm <- lrm(CHD_10 ~ hx_htn + tot_chol, data = fram2,
               x = TRUE, y = TRUE)
```

## Then We Fit a Kitchen Sink Model (Class 11)

```
m_03 <- glm(CHD_10 ~ hx_htn + tot_chol + sex + age +  
             smoker + cigs_day + bp_meds +  
             hx_stroke + hx_dm + ed_f + sbp + dbp +  
             bmi + heart_r + glucose,  
            data = fram2, family = binomial)  
  
d <- datadist(fram2)  
options(datadist = "d")  
m_03_lrm <- lrm(CHD_10 ~ hx_htn + tot_chol + sex + age +  
                smoker + cigs_day + bp_meds +  
                hx_stroke + hx_dm + ed_f + sbp + dbp +  
                bmi + heart_r + glucose,  
               data = fram2, x = TRUE, y = TRUE)
```

# Comparing the Current Models

Model	C statistic	Nagelkerke $R^2$
m_01_1rm	0.614	0.051
m_02_1rm	0.640	0.055
m_03_1rm	0.733	0.159

m\_03\_1rm shows the following predictors as highly significant:

- age, sex, sbp, cigs\_day, glucose, and hx\_stroke

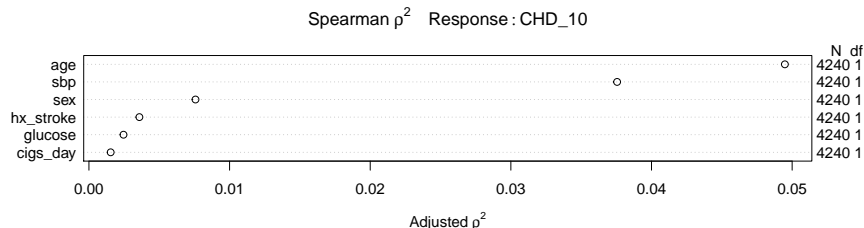


## Fitting a 6-predictor, but still useful model

# What looks useful?

By ANOVA on `m_03_1rm` it looks like `age`, `sex`, `sbp`, `cigs_day`, `glucose`, `hx_stroke` for sure.

```
plot(spearman2(CHD_10 ~ age + sex + sbp + cigs_day +  
              glucose + hx_stroke, data = fram2))
```



## New Model 4

```
m_04 <- glm(CHD_10 ~ rcs(age, 5) + rcs(sbp, 3) + sex +  
            hx_stroke + glucose + cigs_day,  
            data = fram2, family = binomial)  
  
dd <- datadist(fram2)  
options(datadist = "dd")  
  
m_04_lrm <- lrm(CHD_10 ~ rcs(age, 5) + rcs(sbp, 3) + sex +  
               hx_stroke + glucose + cigs_day,  
               data = fram2, x = TRUE, y = TRUE)
```

## m\_04\_lrm

```
> m_04_lrm
```

```
Logistic Regression Model
```

```
lrm(formula = CHD_10 ~ rcs(age, 5) + rcs(sbp, 3) + sex + hx_stroke +  
      glucose + cigs_day, data = fram2, x = TRUE, y = TRUE)
```

		Model Likelihood Ratio Test		Discrimination Indexes		Rank Discrim. Indexes	
Obs	4240	LR chi2	401.44	R2	0.158	C	0.731
0	3596	d.f.	10	g	1.041	Dxy	0.461
1	644	Pr(> chi2)	<0.0001	gr	2.833	gamma	0.461
max  deriv	2e-06			gp	0.120	tau-a	0.119
				Brier	0.115		

	Coef	S.E.	Wald Z	Pr(> Z )
Intercept	-8.7201	2.4092	-3.62	0.0003
age	0.0732	0.0576	1.27	0.2037
age'	0.2871	0.3539	0.81	0.4172
age''	-1.1057	0.9608	-1.15	0.2498
age'''	1.4723	0.9640	1.53	0.1267
sbp	0.0147	0.0066	2.23	0.0256
sbp'	0.0030	0.0080	0.37	0.7080
sex=M	0.4935	0.0973	5.07	<0.0001
hx_stroke	1.0514	0.4345	2.42	0.0155
glucose	0.0076	0.0016	4.69	<0.0001
cigs_day	0.0208	0.0039	5.39	<0.0001

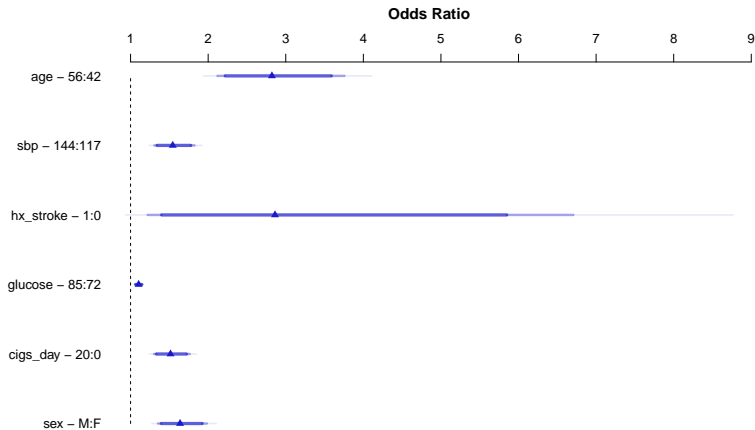
# Validating our Summary Statistics

```
set.seed(432329); validate(m_04_lrm)[1:4,]
```

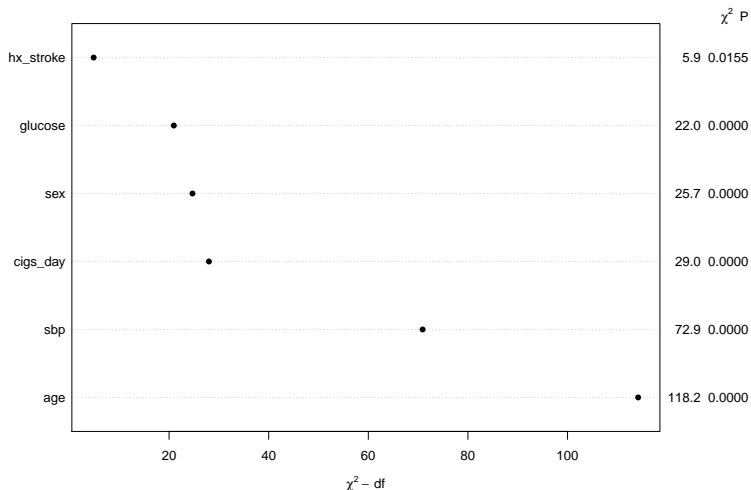
	index.orig	training	test	optimism
Dxy	0.4611110	0.4672261	0.45678154	0.010444568
R2	0.1575417	0.1629811	0.15375377	0.009227324
Intercept	0.0000000	0.0000000	-0.04796015	0.047960154
Slope	1.0000000	1.0000000	0.96803833	0.031961665

	index.corrected	n
Dxy	0.45066647	40
R2	0.14831438	40
Intercept	-0.04796015	40
Slope	0.96803833	40

```
plot(summary(m_04_lrm))
```

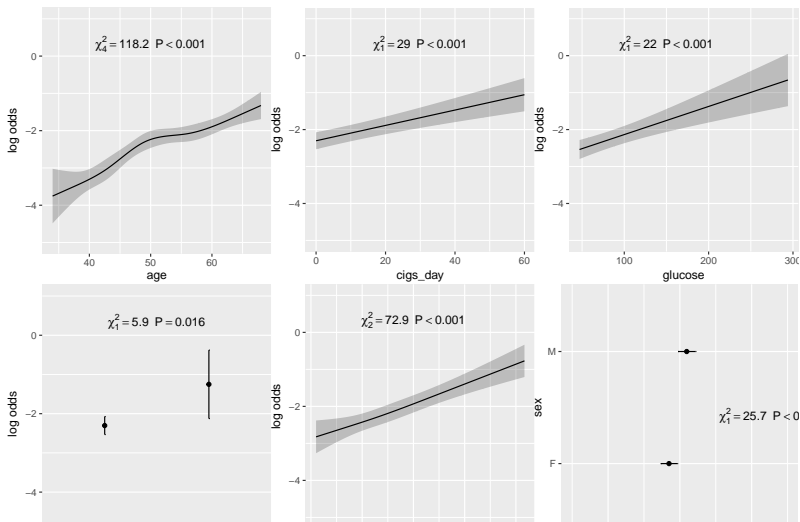


```
plot(anova(m_04_lrm))
```



# Can we see the prediction results?

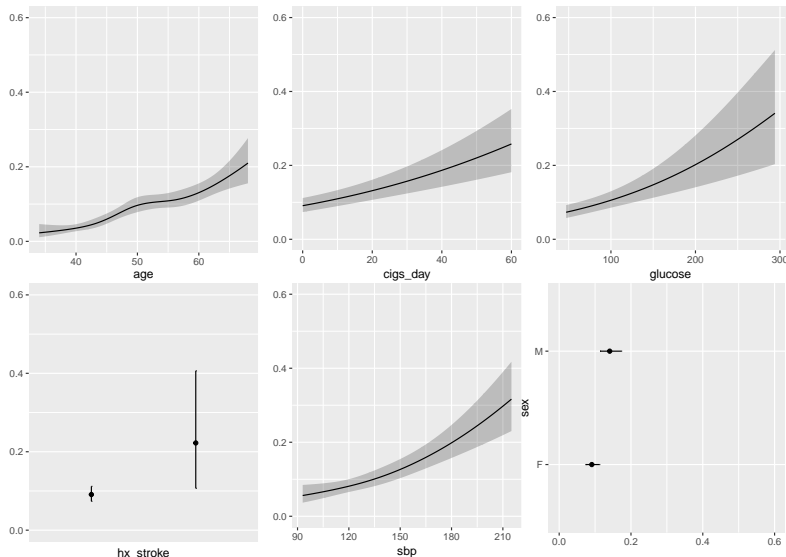
```
ggplot(Predict(m_04_lrm),  
       anova = anova(m_04_lrm), pval = TRUE)
```





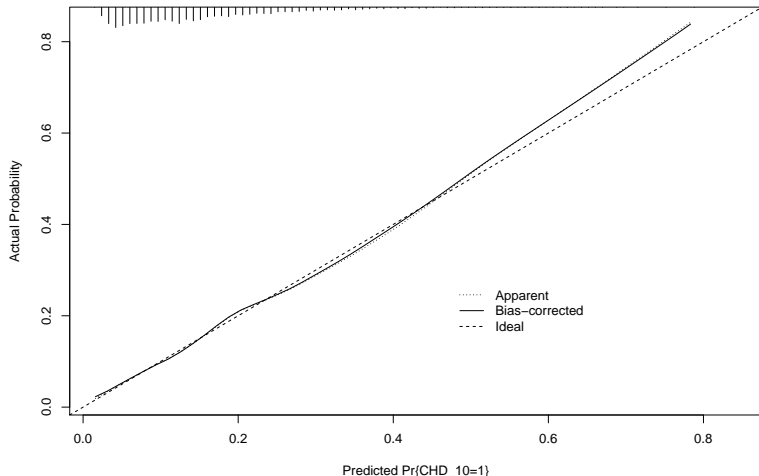
# What about on a better scale?

```
ggplot(Predict(m_04_lrm, fun = plogis))
```



# Calibration of mod\_04\_lrm

```
set.seed(432612); plot(calibrate(m_04_lrm))
```



B= 40 repetitions, boot

Mean absolute error=0.004 n=4240

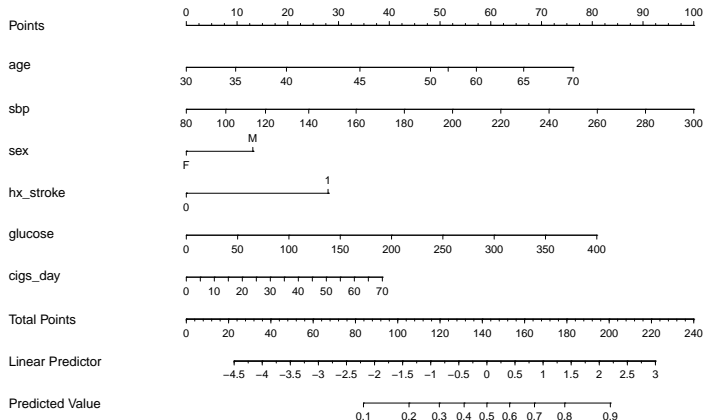
# Goodness of fit test?

```
round(residuals(m_04_lrm, type = "gof"),3)
```

Sum of squared errors	Expected value H0
488.554	489.423
SD	Z
1.183	-0.735
P	
0.462	

# Nomogram of mod\_04\_lrm

```
plot(nomogram(m_04_lrm, fun = plogis))
```



## Comparing Models 3 and 4 (which aren't nested)

```
glance(m_03) # kitchen sink but no non-linear terms
```

	null.deviance	df.null	logLik	AIC	BIC
1	3612.209	4239	-1603.407	3242.813	3357.155

	deviance	df.residual
1	3206.813	4222

```
glance(m_04) # six predictors but with non-linear terms
```

	null.deviance	df.null	logLik	AIC	BIC	deviance
1	3612.209	4239	-1605.382	3232.764	3302.64	3210.764

	df.residual
1	4229

# Checking Residuals?

- Yes/No outcomes contain less information than quantitative outcomes
- Residuals cannot be observed - predicted
  - There are several different types of residuals defined
- Assumptions of logistic regression are different
  - Model is deliberately non-linear
  - Error variance is a function of the mean, so it isn't constant
  - Errors aren't assumed to follow a Normal distribution
  - Only thing that's the same: leverage and influence

So, plot 5 (residuals/leverage/influence) can be a little useful, but that's it.

- We'll need better diagnostic tools for generalized linear models.

# Any observations particularly influential on Model 4?

```
which.influence(m_04_lrm, cutoff = 0.3)
```

```
$Intercept
```

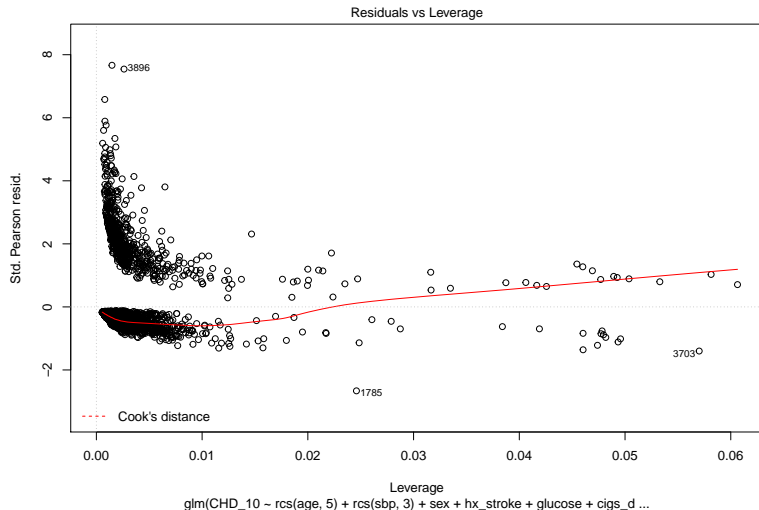
```
[1] "3896"
```

```
$glucose
```

```
[1] "1785" "3703"
```

# Influence and Model 4?

```
plot(m_04, which = 5)
```





# The HERS Trial

# Hormone Therapy and Baseline LDL in the HERS Trial

HERS clinical trial of hormone therapy (HT). We're excluding the women with diabetes here.

```
hers <- read.csv("data/hersdata.csv") %>% tbl_df

hers1 <- hers %>%
  filter(diabetes == "no") %>%
  select(subject, LDL, HT, age, smoking, drinkany, SBP,
         physact, BMI, diabetes)
```

# The Data

```
head(hers1)
```

```
# A tibble: 6 x 10
```

	subject	LDL	HT	age	smoking	drinkany	SBP	physact
	<int>	<dbl>	<fct>	<int>	<fct>	<fct>	<int>	<fct>
1	1	122	place~	70	no	no	138	much m~
2	2	242	place~	62	no	no	118	much l~
3	4	116	place~	64	yes	yes	152	much l~
4	5	151	place~	65	no	no	175	somewh~
5	6	138	hormo~	68	no	yes	174	about ~
6	8	121	hormo~	69	no	no	178	much m~

```
# ... with 2 more variables: BMI <dbl>, diabetes <fct>
```

# The Codebook (n = 2,032 women without diabetes)

Variable	Description	Missing?
subject	subject code	0
LDL	LDL cholesterol in mg/dl	7
HT	factor: hormone therapy or placebo	0
age	age in years	0
smoking	yes or no	0
drinkany	yes or no	2
SBP	systolic BP in mm Hg	0
physact	5-level factor	0
BMI	body-mass index in kg/m <sup>2</sup>	2
diabetes	yes or no (all of these are no)	0

# Our Modeling Goal

Predict LDL using

- age
- smoking
- drinkany
- SBP
- physact
- BMI
- the interaction of smoking and BMI

# Details on physact variable

```
hers1 %>% count(physact)
```

```
# A tibble: 5 x 2
```

	physact	n
	<fct>	<int>
1	about as active	674
2	much less active	107
3	much more active	252
4	somewhat less active	322
5	somewhat more active	677

# Skim?

```
hers1 %>% select(-subject) %>% skim()
```

```
> hers1 %>% select(-subject) %>% skim()
```

Skim summary statistics

n obs: 2032

n variables: 9

Variable type: factor

variable	missing	complete	n	n_unique	top_counts	ordered
diabetes	0	2032	2032	1	no: 2032, yes: 0, NA: 0	FALSE
drinkany	2	2030	2032	2	no: 1135, yes: 895, NA: 2	FALSE
HT	0	2032	2032	2	pla: 1031, hor: 1001, NA: 0	FALSE
physact	0	2032	2032	5	som: 677, abo: 674, som: 322, muc: 252	FALSE
smoking	0	2032	2032	2	no: 1733, yes: 299, NA: 0	FALSE

Variable type: integer

variable	missing	complete	n	mean	sd	p0	p25	median	p75	p100	hist
age	0	2032	2032	66.89	6.75	44	62	67	72	79	
SBP	0	2032	2032	133.38	18.47	83	120	132	145	197	

Variable type: numeric

variable	missing	complete	n	mean	sd	p0	p25	median	p75	p100	hist
BMI	2	2030	2032	27.67	5.14	15.21	24.2	26.89	30.27	54.13	
LDL	7	2025	2032	145.65	37.07	36.8	120.6	141.4	166	351.2	

# Missingness pattern?

```
na.pattern(hers1) # from Hmisc
```

```
pattern
```

```
0000000000 0000000010 0000010000 0100000000  
          2021          2          2          7
```

```
names(hers1)
```

```
[1] "subject" "LDL"      "HT"       "age"      "smoking"  
[6] "drinkany" "SBP"     "physact"  "BMI"      "diabetes"
```

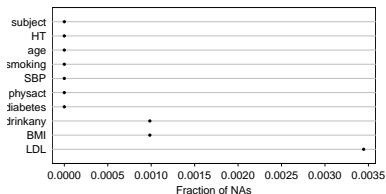
## Next slide

```
par(mfrow = c(2,2))  
naplot(naclus(hers1))  
par(mfrow = c(1,1))
```

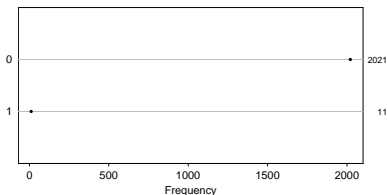


# naplot(naclus(hers1))

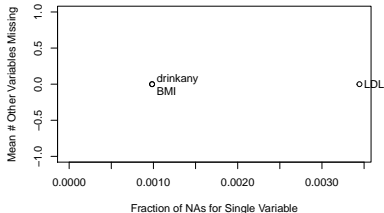
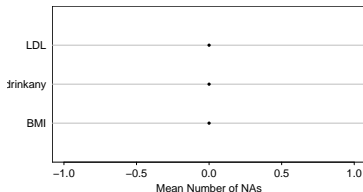
Fraction of NAs in each Variable



Number of Missing Variables Per Observation



Mean Number of Other Variables Missing for Observations where Indicated Variable is NA



## Simple Imputation into `hers2`

# Simple Imputation for drinkany, BMI and LDL

Since drinkany is a factor, we have to do some extra work to impute.

```
set.seed(432092)

hers2 <- hers1 %>%
  mutate(drinkany_n =
    ifelse(drinkany == "yes", 1, 0)) %>%
  impute_pmm(drinkany_n ~ age + smoking) %>%
  mutate(drinkany =
    ifelse(drinkany_n == 1, "yes", "no")) %>%
  impute_rlm(BMI ~ age + smoking + SBP) %>%
  impute_rlm(LDL ~ age + smoking + SBP + BMI)
```

## Now, check missingness...

```
na.pattern(hers2)
```

```
pattern
000000000000
      2032
```

```
names(hers2)
```

```
[1] "subject"      "LDL"          "HT"           "age"
[5] "smoking"      "drinkany"     "SBP"          "physact"
[9] "BMI"          "diabetes"     "drinkany_n"
```

# Multiple Imputation with aregImpute

# Multiple Imputation using aregImpute from Hmisc

Model to predict all missing values of any variables, using additive regression bootstrapping and predictive mean matching.

Steps are:

- 1 aregImpute draws a sample with replacement from the observations where the target variable is observed, not missing.
- 2 It then fits a flexible additive model to predict this target variable while finding the optimum transformation of it.
- 3 It then uses this fitted flexible model to predict the target variable in all of the original observations.
- 4 Finally, it imputes each missing value of the target variable with the observed value whose predicted transformed value is closest to the predicted transformed value of the missing value.

# Fitting a Multiple Imputation Model

```
set.seed(4320132)
dd <- datadist(hers1)
options(datadist = "dd")
fit3 <- aregImpute(~ LDL + age + smoking + drinkany +
                  SBP + physact + BMI,
                  nk = c(0, 3:5), tlinear = FALSE,
                  data = hers1, B = 10, n.impute = 20)
```

Iteration 1

Iteration 2

Iteration 3

Iteration 4

Iteration 5

Iteration 6

Iteration 7

Iteration 8

Iteration 9

# Multiple Imputation using `aregImpute` from `Hmisc`

`aregImpute` requires specifications of all variables, and several other details:

- `n.impute` = number of imputations, we'll run 20
- `nk` = number of knots to describe level of complexity, with our choice `nk = c(0, 3:5)` we'll fit both linear models and models with restricted cubic splines with 3, 4, and 5 knots
- `tlinear = FALSE` allows the target variable to have a non-linear transformation when `nk` is 3 or more
- `B = 10` specifies 10 bootstrap samples will be used
- `data` specifies the source of the variables



# aregImpute Imputation Results (1 of 3)

```
fit3
```

```
> fit3
```

Multiple Imputation using Bootstrap and PMM

```
aregImpute(formula = ~LDL + age + smoking + drinkany + SBP +  
  physact + BMI, data = hers1, n.impute = 5, nk = c(0, 3:5),  
  tlinear = FALSE, B = 10)
```

```
n: 2032      p: 7      Imputations: 5      nk: 0
```

Number of NAs:

LDL	age	smoking	drinkany	SBP	physact	BMI
7	0	0	2	0	0	2

	type	d.f.
--	------	------

LDL	s	1
age	s	1
smoking	c	1
drinkany	c	1
SBP	s	1
physact	c	4
BMI	s	1

## aregImpute Imputation Results (2 of 3)

R-squares for Predicting Non-Missing Values for Each Variable  
Using Last Imputations of Predictors

	LDL	drinkany	BMI
	0.019	0.029	0.093

Resampling results for determining the complexity of imputation models

Variable being imputed: LDL

			nk=0	nk=3	nk=4	nk=5
Bootstrap bias-corrected	R <sup>2</sup>		0.0151	0.0149	0.0101	0.00976
10-fold cross-validated	R <sup>2</sup>		0.0154	0.0224	0.0129	0.01913
Bootstrap bias-corrected	mean	error	28.2956	42.5462	43.9458	39.56345
10-fold cross-validated	mean	error	145.8339	43.2577	44.8477	45.07517
Bootstrap bias-corrected	median	error	22.7110	35.0460	38.9420	32.95988
10-fold cross-validated	median	error	142.1896	35.3638	38.3446	38.08496

## aregImpute Imputation Results (3 of 3)

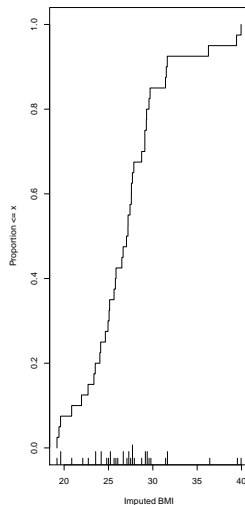
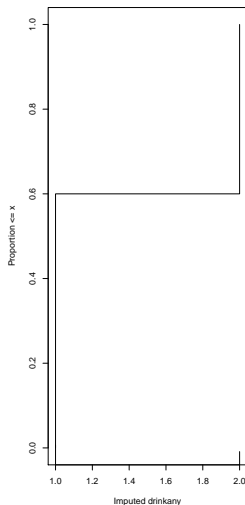
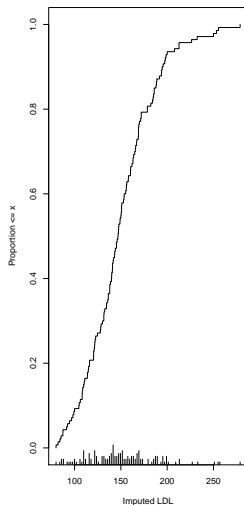
Variable being imputed: drinkany

		nk=0	nk=3	nk=4	nk=5
Bootstrap bias-corrected	R <sup>2</sup>	0.0138	0.0110	0.0131	0.0104
10-fold cross-validated	R <sup>2</sup>	0.0191	0.0184	0.0182	0.0122
Bootstrap bias-corrected	mean  error	0.4535	0.4529	0.4551	0.4567
10-fold cross-validated	mean  error	0.4482	0.4499	0.4382	0.4644
Bootstrap bias-corrected	median  error	0.0000	0.0000	0.0000	0.0000
10-fold cross-validated	median  error	0.2000	0.3000	0.0000	0.2000

Variable being imputed: BMI

		nk=0	nk=3	nk=4	nk=5
Bootstrap bias-corrected	R <sup>2</sup>	0.0851	0.0878	0.0858	0.0892
10-fold cross-validated	R <sup>2</sup>	0.0880	0.0940	0.0930	0.0923
Bootstrap bias-corrected	mean  error	3.8316	4.8352	4.9466	5.1806
10-fold cross-validated	mean  error	27.6718	4.8115	4.9504	4.9765
Bootstrap bias-corrected	median  error	2.9944	4.0176	4.0021	4.2517
10-fold cross-validated	median  error	27.0170	4.0261	3.9930	4.0836

# A plot of the imputed values... (results)



## A plot of the imputed values... (code)

```
par(mfrow = c(1,3))  
plot(fit3)  
par(mfrow = c(1,1))
```

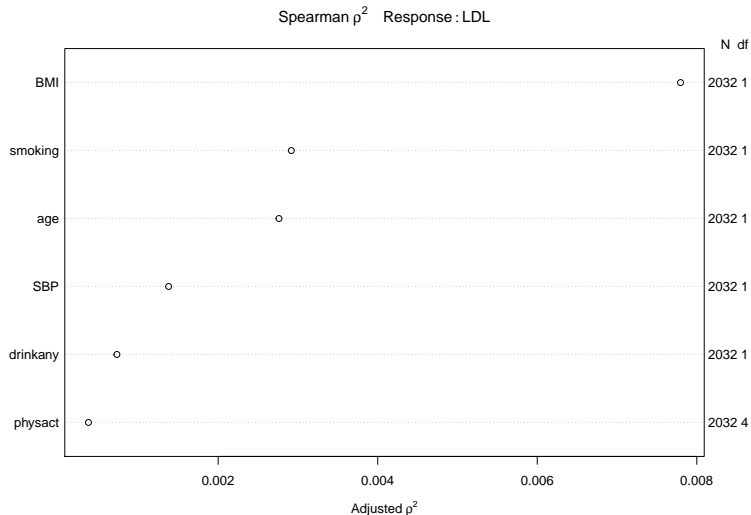
- For LDL, we imputed most of the 7 missing subjects in most of the 20 imputation runs to values within a range of around 120 through 200, but occasionally, we imputed values that were substantially lower than 100.
- For `drinkany` we imputed about 70% no and 30% yes.
- For BMI, we imputed values ranging from about 23 to 27 in many cases, and up near 40 in other cases.
- This method never imputes a value for a variable that doesn't already exist in the data.

# Spearman $\rho^2$ Plot

We've already decided to include a BMI\*smoking product term, but how should we prioritize the degrees of freedom we spend on non-linearity otherwise?

```
plot(spearman2(LDL ~ age + smoking + drinkany + SBP +  
              physact + BMI, data = hers2))
```

# Spearman $\rho^2$ Plot Result



## Fitting a Linear Regression with `ols`



# Model we'll fit

Fitting a model to predict LDL using

- BMI with a restricted cubic spline, 5 knots
- age with a quadratic polynomial
- SBP as a linear term
- drinkany indicator
- physact factor
- smoking indicator and its interaction with BMI

We could fit this to the data

- restricted to complete cases (hers1, effectively)
- after simple imputation (hers2)
- after our multiple imputation (fit3)

# Fitting the model after simple imputation

```
dd <- datadist(hers2)
options(datadist = "dd")

m2 <- ols(LDL ~ rcs(BMI, 5) + pol(age, 2) + SBP +
          drinkany + physact + smoking +
          smoking %ia% BMI, data = hers2,
          x = TRUE, y = TRUE)
```

where %ia% identifies the linear interaction alone.

## m2 results (slide 1 of 2)

```
> m2
Linear Regression Model

ols(formula = LDL ~ rcs(BMI, 5) + pol(age, 2) + SBP + drinkany +
     physact + smoking + smoking %ia% BMI, data = hers2, x = TRUE,
     y = TRUE)
```

		Model Likelihood		Discrimination	
		Ratio Test		Indexes	
Obs	2032	LR chi2	53.14	R2	0.026
sigma	36.6503	d.f.	14	R2 adj	0.019
d.f.	2017	Pr(> chi2)	0.0000	g	6.631

### Residuals

	Min	1Q	Median	3Q	Max
	-113.379	-24.326	-3.835	20.832	197.097

## m2 results (slide 2 of 2)

	Coef	S.E.	t	Pr(> t )
Intercept	120.2662	67.6113	1.78	0.0754
BMI	1.5508	1.0071	1.54	0.1237
BMI'	-8.4486	9.0978	-0.93	0.3532
BMI''	39.6413	37.1378	1.07	0.2859
BMI'''	-54.8924	44.2677	-1.24	0.2151
age	-0.5249	1.9490	-0.27	0.7877
age^2	0.0014	0.0148	0.10	0.9233
SBP	0.1209	0.0451	2.68	0.0074
drinkany=yes	-3.7023	1.6544	-2.24	0.0253
physact=much less active	-4.7408	3.8621	-1.23	0.2198
physact=much more active	-0.2635	2.7391	-0.10	0.9234
physact=somewhat less active	0.0130	2.5101	0.01	0.9959
physact=somewhat more active	3.8031	2.0193	1.88	0.0598
smoking=yes	-6.8961	12.0196	-0.57	0.5662
smoking=yes * BMI	0.4892	0.4375	1.12	0.2636

# Validation of summary statistics

```
validate(m2)
```

	index.orig	training	test	optimism
R-square	0.0258	0.0307	0.0188	0.0119
MSE	1333.3300	1320.0677	1342.9027	-22.8350
g	6.6306	7.1548	5.8726	1.2821
Intercept	0.0000	0.0000	26.2153	-26.2153
Slope	1.0000	1.0000	0.8208	0.1792

	index.corrected	n
R-square	0.0139	40
MSE	1356.1650	40
g	5.3485	40
Intercept	26.2153	40
Slope	0.8208	40

## anova(m2) results

```
> anova(m2)
```

	Analysis of Variance		Response: LDL		
Factor	d.f.	Partial SS	MS	F	P
BMI (Factor+Higher Order Factors)	5	2.758824e+04	5517.64861	4.11	0.0010
All Interactions	1	1.679813e+03	1679.81344	1.25	0.2636
Nonlinear	3	9.735452e+03	3245.15068	2.42	0.0647
age	2	9.175762e+03	4587.88077	3.42	0.0330
Nonlinear	1	1.244351e+01	12.44351	0.01	0.9233
SBP	1	9.657476e+03	9657.47569	7.19	0.0074
drinkany	1	6.726918e+03	6726.91809	5.01	0.0253
physact	4	9.709992e+03	2427.49791	1.81	0.1247
smoking (Factor+Higher Order Factors)	2	1.085405e+04	5427.02463	4.04	0.0177
All Interactions	1	1.679813e+03	1679.81344	1.25	0.2636
smoking * BMI (Factor+Higher Order Factors)	1	1.679813e+03	1679.81344	1.25	0.2636
TOTAL NONLINEAR	4	9.738807e+03	2434.70175	1.81	0.1237
TOTAL NONLINEAR + INTERACTION	5	1.171134e+04	2342.26845	1.74	0.1214
REGRESSION	14	7.178905e+04	5127.78931	3.82	<.0001
ERROR	2017	2.709327e+06	1343.24569		

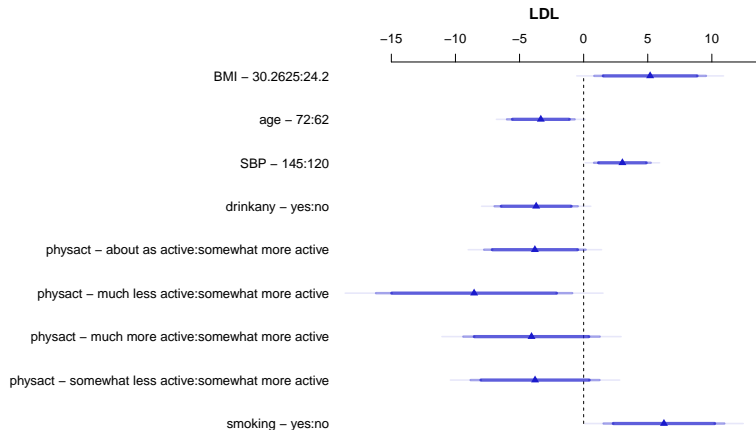
## summary(m2) results

```
> summary(m2)
```

Effects	Response : LDL							
Factor	Low	High	Diff.	Effect	S.E.	Lower 0.95	Upper 0.95	
BMI	24.2	30.263	6.0625	5.1862	2.2217	0.82921	9.54330	
age	62.0	72.000	10.0000	-3.3412	1.3450	-5.97890	-0.70357	
SBP	120.0	145.000	25.0000	3.0218	1.1270	0.81165	5.23190	
drinkany - yes:no	1.0	2.000	NA	-3.7023	1.6544	-6.94690	-0.45779	
physact - about as active:somewhat more active	5.0	1.000	NA	-3.8031	2.0193	-7.76310	0.15695	
physact - much less active:somewhat more active	5.0	2.000	NA	-8.5439	3.9035	-16.19900	-0.88862	
physact - much more active:somewhat more active	5.0	3.000	NA	-4.0666	2.7125	-9.38630	1.25310	
physact - somewhat less active:somewhat more active	5.0	4.000	NA	-3.7901	2.5633	-8.81720	1.23690	
smoking - yes:no	1.0	2.000	NA	6.2635	2.4009	1.55500	10.97200	

Adjusted to: BMI=26.9 smoking=no

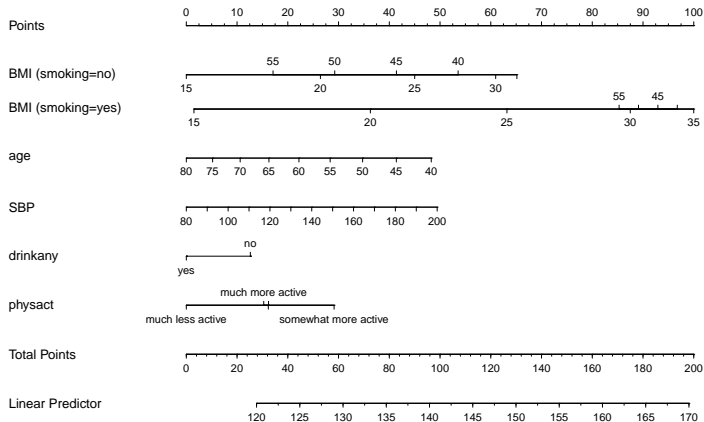
# plot(summary(m2)) results



Adjusted to: BMI=26.9 smoking=no



# plot(nomogram(m2))



# Making Predictions for an Individual

Suppose now that we want to use R to get a prediction for a new individual subject with BMI = 30, age = 50, smoking = yes and physact = about as active, drinkany= yes and SBP of 150.

```
predict(m2, expand.grid(BMI = 30, age = 50, smoking = "yes",  
                        physact = "about as active",  
                        drinkany = "yes", SBP = 150),  
        conf.int = 0.95, conf.type = "individual")
```

\$linear.predictors	\$lower	\$upper
160.9399	88.48615	233.3936

# Making Predictions for a Long-Run Mean

The other kind of prediction we might wish to make is for the mean of a series of subjects whose BMI = 30, age = 50, smoking = yes and physact = about as active, drinkany= yes and SBP of 150.

```
predict(m2, expand.grid(BMI = 30, age = 50, smoking = "yes",  
                        physact = "about as active",  
                        drinkany = "yes", SBP = 150),  
        conf.int = 0.95, conf.type = "mean")
```

\$linear.predictors	\$lower	\$upper
160.9399	151.8119	170.0679

Of course, the confidence interval will always be narrower than the prediction interval given the same predictor values.

# Influential Points?

```
which.influence(m2, cutoff = 0.4)
```

```
$Intercept
```

```
[1] 1135
```

```
$age
```

```
[1] 1135
```

```
$smoking
```

```
[1] 132
```

```
$`smoking * BMI`
```

```
[1] 132
```

# Fitting the model to the complete cases

```
d <- datadist(hers1)
options(datadist = "d")

m1 <- ols(LDL ~ rcs(BMI, 5) + pol(age, 2) + SBP +
          drinkany + physact + smoking +
          smoking %ia% BMI, data = hers1,
          x = TRUE, y = TRUE)
```

where %ia% identifies the linear interaction alone.

## Putting it Together

# What have we got?

- An imputation model `fit3`

```
fit3 <- aregImpute(~ LDL + age + smoking + drinkany + SBP +  
  physact + BMI, nk = c(0, 3:5), tlinear = FALSE,  
  data = hers1, B = 10, n.impute = 20, x = TRUE)
```

- A prediction model

```
m1 <- ols(LDL ~ rcs(BMI, 5) + pol(age, 2) + SBP +  
  drinkany + physact + smoking + smoking %ia% BMI,  
  x = TRUE, y = TRUE)
```

Now we put them together

# Linear Regression & Imputation Model

```
m3imp <-  
  fit.mult.impute(LDL ~ rcs(BMI, 5) + pol(age, 2) + SBP +  
    drinkany + physact + smoking +  
    smoking %ia% BMI,  
    fitter = ols, xtrans = fit3,  
    data = hers1)
```

Variance Inflation Factors Due to Imputation:

Intercept	BMI
1.00	1.00
BMI '	BMI ' '
1.00	1.00
BMI ' ' '	age
1.00	1.00
age^2	SBP



## m3imp results (1 of 2)

```
> m3imp
Linear Regression Model

fit.mult.impute(formula = LDL ~ rcs(BMI, 5) + pol(age, 2) + SBP +
  drinkany + physact + smoking + smoking %ia% BMI, fitter = ols,
  xtrans = fit3, data = hers1)
```

		Model Likelihood		Discrimination	
		Ratio Test		Indexes	
Obs	2032	LR chi2	53.30	R2	0.026
sigma	36.7128	d.f.	14	R2 adj	0.019
d.f.	2017	Pr(> chi2)	0.0000	g	6.652

### Residuals

Min	1Q	Median	3Q	Max
-113.10	-24.46	-3.81	20.92	197.42

## m3imp results (2 of 2)

	Coef	S.E.	t	Pr(> t )
Intercept	121.1499	67.7998	1.79	0.0741
BMI	1.5445	1.0097	1.53	0.1263
BMI'	-8.2945	9.1027	-0.91	0.3623
BMI''	39.0890	37.3055	1.05	0.2949
BMI'''	-54.2119	44.4779	-1.22	0.2230
age	-0.5521	1.9547	-0.28	0.7776
age^2	0.0016	0.0148	0.11	0.9119
SBP	0.1216	0.0453	2.69	0.0073
drinkany=yes	-3.7404	1.6625	-2.25	0.0246
physact=much less active	-4.7426	3.8692	-1.23	0.2204
physact=much more active	-0.2665	2.7455	-0.10	0.9227
physact=somewhat less active	0.0313	2.5214	0.01	0.9901
physact=somewhat more active	3.8060	2.0257	1.88	0.0604
smoking=yes	-6.9198	12.0472	-0.57	0.5658
smoking=yes * BMI	0.4917	0.4388	1.12	0.2626

# anova(m3imp)

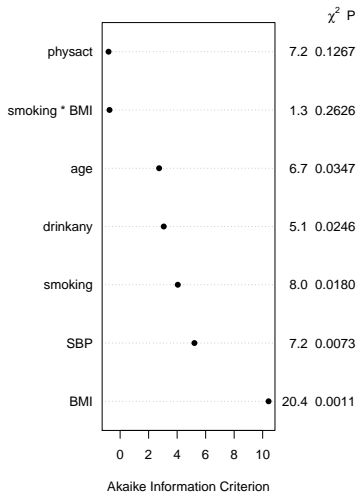
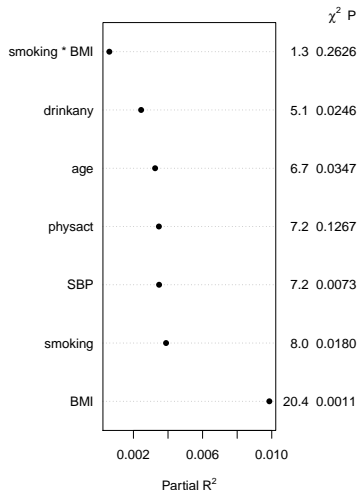
```
> anova(m3imp)
```

Analysis of Variance

Response: LDL

Factor	d.f.	Partial SS	MS	F	P
BMI (Factor+Higher Order Factors)	5	27514.6406	5502.9281	4.08	0.0011
All Interactions	1	1692.6044	1692.6044	1.26	0.2626
Nonlinear	3	9741.6194	3247.2065	2.41	0.0653
age	2	9078.9851	4539.4926	3.37	0.0347
Nonlinear	1	16.5032	16.5032	0.01	0.9119
SBP	1	9721.1667	9721.1667	7.21	0.0073
drinkany	1	6822.3861	6822.3861	5.06	0.0246
physact	4	9690.3632	2422.5908	1.80	0.1267
smoking (Factor+Higher Order Factors)	2	10845.6127	5422.8063	4.02	0.0180
All Interactions	1	1692.6044	1692.6044	1.26	0.2626
smoking * BMI (Factor+Higher Order Factors)	1	1692.6044	1692.6044	1.26	0.2626
TOTAL NONLINEAR	4	9747.0966	2436.7741	1.81	0.1246
TOTAL NONLINEAR + INTERACTION	5	11717.3715	2343.4743	1.74	0.1225
REGRESSION	14	71571.1297	5112.2236	3.79	<.0001
ERROR	2017	2718570.0412	1347.8285		

# Evaluation via Partial R<sup>2</sup> and AIC (result)



# Evaluation via Partial $R^2$ and AIC (code)

```
par(mfrow = c(1,2))  
plot(anova(m3imp), what="partial R2")  
plot(anova(m3imp), what="aic")  
par(mfrow = c(1,1))
```

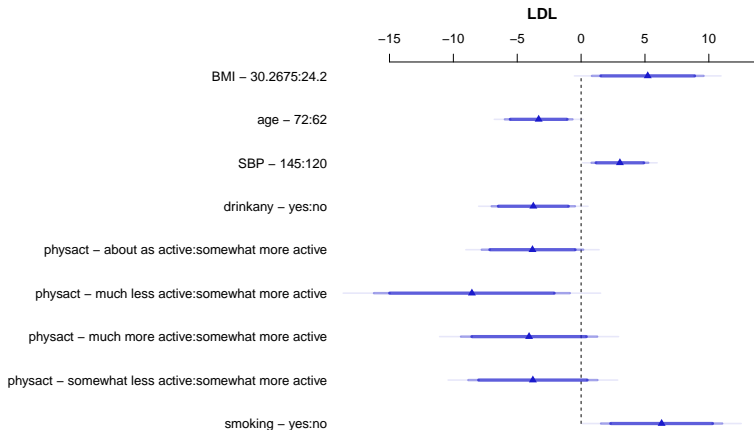
# summary(m3imp)

```
> summary(m3imp)
```

Effects	Response : LDL						
Factor	Low	High	Diff.	Effect	S.E.	Lower 0.95	Upper 0.95
BMI	24.2	30.268	6.0675	5.2165	2.2287	0.84565	9.58730
age	62.0	72.000	10.0000	-3.3219	1.3498	-5.96910	-0.67463
SBP	120.0	145.000	25.0000	3.0394	1.1317	0.81989	5.25880
drinkany - yes:no	1.0	2.000	NA	-3.7404	1.6625	-7.00080	-0.47996
physact - about as active:somewhat more active	5.0	1.000	NA	-3.8060	2.0257	-7.77860	0.16663
physact - much less active:somewhat more active	5.0	2.000	NA	-8.5486	3.9114	-16.21900	-0.87779
physact - much more active:somewhat more active	5.0	3.000	NA	-4.0724	2.7198	-9.40640	1.26160
physact - somewhat less active:somewhat more active	5.0	4.000	NA	-3.7746	2.5773	-8.82900	1.27980
smoking - yes:no	1.0	2.000	NA	6.3043	2.4196	1.55900	11.05000

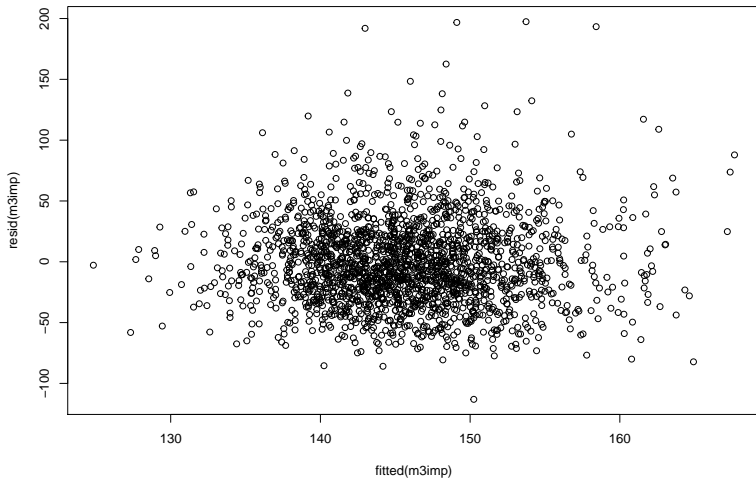
Adjusted to: BMI=26.895 smoking=no

```
plot(summary(m3imp))
```



Adjusted to: BMI=26.895 smoking=no

```
plot(resid(m1imp) ~ fitted(m1imp))
```





# plot(nomogram(m3imp))

