

Working with logistic regression models

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Further remarks on logistic regression

Test of fit: The Hosmer-Lemeshow test

Conditional logistic regression

Missing data

A small example - non completely random sample

Complete data analysis - bias

Missing at random vs missing completely at random

Introduction to techniques

Sampling weights

Imputation

Full modelling

Sensitivity analyses

Data with several random components: Binary outcome

Clustered binary data with one random component

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Logistic regression models: Test of fit

A common, and to some extend informative, test of fit is the Hosmer-Lemeshow test.

Consider the model for obesity from Day 4

$$\text{logit}(\text{Pr}(\text{obese})) = \beta_0 + \beta_1 \cdot \text{woman} + \beta_2 \cdot (\text{age} - 45)$$

Logit estimates

Number of obs = 4690

LR chi2(2) = 55.68

Prob > chi2 = 0.0000

Pseudo R2 = 0.0155

Log likelihood = -1767.7019

obese	Coef.	Std. Err.	z	P> z	[95% Conf. Interval]
sex					
1	(base)				
2	.2743976	.0903385	3.04	0.002	.0973374 .4514579
age45	.0344723	.0051354	6.71	0.000	.0244072 .0445374
_cons	-2.147056	.0721981	-29.74	0.000	-2.288561 -2.00555

Significantly better than nothing - but is it good?

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Logistic regression models: Test of fit

What about comparing the estimated prevalence with the observed prevalence?

In the Hosmer-Lemeshow test the data is divided into groups (traditionally 10) according to the estimated probabilities

and the observed and expected counts are compared in these groups by a chi-square test.

Most programs, that can fit a logistic regression model, can calculate this test.

In Stata it is done by (after fitting the model):

estat gof, group(10) table

The data is divided into deciles after the estimated probabilities.

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Logistic regression models: Test of fit

OUTPUT

Logistic model for obese, goodness-of-fit test  
(Table collapsed on quantiles of estimated probabilities)

Group	Prob	Obs_1	Exp_1	Obs_0	Exp_0	Total
1	0.0841	64	40.9	462	485.1	526
2	0.0953	43	45.5	453	450.5	496
3	0.1045	44	44.6	398	397.4	442
4	0.1112	42	50.3	422	413.7	464
5	0.1217	44	51.4	394	386.6	438
6	0.1332	52	63.0	441	430.0	493
7	0.1456	53	61.7	389	380.3	442
8	0.1592	62	69.8	392	384.2	454
9	0.1834	98	89.9	424	432.1	522
10	0.2407	99	83.8	314	329.2	413

number of observations = 4690

number of groups = 10

Hosmer-Lemeshow chi2(8) = 26.01

Prob > chi2 = 0.0010

One problem: Too many in the tails

Significant difference between observed and expected!

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Logistic regression models: Test of fit

```
logit obese i.sex##age45
estat gof, group(10) table
Logistic model for obese, goodness-of-fit test
(Table collapsed on quantiles of estimated probabilities)
```

Group	Prob	Obs_1	Exp_1	Obs_0	Exp_0	Total
1	0.0796	36	35.9	466	466.1	502
2	0.1011	42	41.1	406	406.9	448
3	0.1053	49	49.6	429	428.4	478
4	0.1096	50	54.8	458	453.2	508
5	0.1124	52	54.2	436	433.8	488
6	0.1153	51	46.4	355	359.6	406
7	0.1182	52	53.9	410	408.1	462
8	0.1590	76	70.3	428	433.7	504
9	0.2133	96	91.8	391	395.2	487
10	0.3310	97	103.0	310	304.0	407

number of observations = 4690

number of groups = 10

Hosmer-Lemeshow chi2(8) = 2.43

Prob > chi2 = 0.9650

The model 'fits' - when we look at it this way !!!!!

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Conditional logistic regression

When

Used in two situations:

1. Matched studies (binary response).

2. Unmatched studies with a confounder with many distinct values.

In 1. the models correspond to the way data was collected.

In 2. the method adjust for a 'mathematical' flaw in the unconditional method.

An example of situation 2:

The confounder is "kommune" having 275 distinct values.

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Conditional logistic regression

What

The logistic regression model (outcome disease yes/no):

$$\ln(odds) = \alpha + \sum_{i=1}^k (\beta_i \cdot x_i)$$

ln(odds) in reference

ln(odds ratios)

Suppose the model above hold in each strata:

$$\ln(odds) = \alpha_s + \sum_{i=1}^k (\beta_i \cdot x_i)$$

ln(odds) in reference

ln(odds ratios)

different in each strata the same in each strata

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Conditional logistic regression

What

$$\ln(odds) = \alpha_s + \sum_{i=1}^k (\beta_i \cdot x_i)$$

ln(odds) different in each strata

We are not interested in these !

In a matched study these are 'controlled'.

In a conditional logistic regression one 'condition on the odds in each strata' , i.e. the case/control ratio.

In the conditional model the  $\alpha$ 's disappear !

The  $\beta$ 's , the log OR's, are still in and can be estimated.

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Conditional logistic regression  
How

A study of cancer in the oral cavity

Matched on **gender** and **10-year age groups**

Ten strata (*genage*)

Here we focus on

*textile-worker* and

*life time consumption of alcohol* (three groups)

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Conditional logistic regression  
How

logistic regression in *Stata*

`logit cancer textile i.alkcon i.genage, or`

`binreg cancer textile i.alkcon i.genage, or`

cancer	Odds Ratio	Std. Err.	z	P> z	[95% Conf. Interval]
textile	1.652484	.6843458	1.21	0.225	.7338846 3.720889
alkcon					
0	1 (base)				
1	1.588614	.4485983	1.64	0.101	.9133833 2.763017
2	15.12845	4.890496	8.40	0.000	8.028433 28.50742
genage					
1	1 (base)				
2	1.277731	.598937	0.20	0.845	.1099855 14.84645
3	.6101724	.3357944	-0.90	0.369	.2074977 1.794287
4	1.196961	.7668028	0.28	0.779	.3410196 4.201272
5	.7482746	.4102097	-0.53	0.597	.2555206 2.191271
6	1.237034	.7746878	0.34	0.734	.3625102 4.221272
7	.7940664	.4252551	-0.43	0.667	.2779736 2.26835
8	1.734638	.9130996	1.05	0.295	.6182202 4.867148
9	1.032018	.6072521	0.05	0.957	.3257093 3.269977
10	1.745782	.9768952	1.00	0.319	.5830142 5.227581
_cons	.2301051	.1095992	-3.08	0.002	.0904687 .5852672 10

Conditional logistic regression in *Stata*

The syntax:

`clogit cancer textile i.alkcon, group(genage) or`

Part of the output:

cancer	Odds Ratio	Std. Err.	z	P> z	[95% Conf. Interval]
textile	1.63708	.6717022	1.20	0.230	.732517 3.658661
alkcon					
0	(base)				
1	1.572508	.4390957	1.62	0.105	.909724 2.718168
2	14.30908	4.569879	8.33	0.000	7.651811 26.75835

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Missing data - example 1

Consider the Frammingham study and imagine, that (due to a limited budget) only 500 measurements of SBP were allowed.

It was decided to take SBP measurements on **100** random participants in each of the age groups -40 and 60+ and **150** in each of the age groups 40-50 and 50-60.

That is we have missing SBP on 4190 of the 4,690 participants!

A short description of the design and the data:

agegrp	Freq.	N(sbp)	mean(sbp)	sd(sbp)
0-	1,325	100	122.18	15.4327
40-	1,684	150	130.85	22.2366
50-	1,346	150	140.93	22.4819
60-	335	100	149.51	26.9251
Total	4,690	500	135.87	24.0783

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Missing data - example 1

agegrp	Freq.	N(sbp)	mean(sbp)	sd(sbp)
0-	1,325	100	122.18	15.4327
40-	1,684	150	130.85	22.2366
50-	1,346	150	140.93	22.4819
60-	335	100	149.51	26.9251
Total	4,690	500	135.87	24.0783

We note:

- This is not a **completely** random sample
- the chance of being sample depends on age group!

The overall (total) average SBP is a biased estimate of the mean SBP among participants in the Frammingham study!

I.e. an analysis of the 500 participants (a complete data analysis) will be biased.

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Missing data - example 1

agegrp	Freq.	N(sbp)	mean(sbp)	sd(sbp)
0-	1,325	100	122.18	15.4327
40-	1,684	150	130.85	22.2366
50-	1,346	150	140.93	22.4819
60-	335	100	149.51	26.9251
Total	4,690	500	135.87	24.0783

We also note:

- Within each age group** the sample is **completely** random.
- Within each age group** the average SBP is an **unbiased** estimate of the mean SBP in the age group.
- We know the size of each age group.
- We can **calculate an unbiased** estimate of the total mean by weighing the group averages.

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Missing data - example 1

agegrp	Freq.	N(sbp)	mean(sbp)	sd(sbp)
0-	1,325	100	122.18	15.4327
40-	1,684	150	130.85	22.2366
50-	1,346	150	140.93	22.4819
60-	335	100	149.51	26.9251
Total	4,690	500	135.87	24.0783

An unbiased estimate can be found as the **weighted average** of the group averages using the group sizes as weights:

$$\frac{122.18 \cdot 1325 + 130.85 \cdot 1684 + 140.93 \cdot 1346 + 149.51 \cdot 335}{4690} = 132.62$$

**Conclusion:** Although this is not a completely random sample, we have enough information in the data to find an unbiased estimate!!!!  
(Assuming completely random sample **within** age group!)

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Assuming that SBP is related to age:

Being missing is **not independent** of the **unobserved** SBP.

but

Being missing is **independent** of the unobserved SBP, **when we know the age group of the individual.**

The first statement means that the data is not **missing completely at random (MCAR).**

The second statement corresponds to **missing at random (MAR)**, i.e. that given **all what we have observed** (including age group), then the missingness is (completely) random, i.e. independent of the unobserved data.

Mathematically Missing At Random implies that one (in theory) has enough information in the **observed data** to correct for the missing data - in principle.

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**Missing data: Standard terminology**

**Missing completely at random (MCAR).**  
The observed data is a (completely) random sample:  
**A complete data analysis will be unbiased**

**Missing at random (MAR)**  
Given **all what we have observed**, then the missingness is (completely) random (independent of the unobserved data):  
The biased sampling **might be adjusted for**.

**Missing not at random (MNAR)**  
Non of the two above apply:  
We will need further assumptions in order to analyse the data.

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**Missing at random**

When the data is **missing at random**, then one can, in theory, make unbiased inference based on the observed data.

In the SBP example such an analysis could be to use the **weighted average** SBP instead of the biased unweighted average.

**In general**  
If the sampled persons are not a completely random sample, but the  $i$ th person is sampled with a **known** probability,  $p_i$ , then we can obtain unbiased estimates by weighing the  $i$ th person with  $1/p_i$ .  
The method is called Inverse Probability Weighing.

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**Inverse probability weighting**

The SBP data:  
Four different sampling probabilities and weights:  
 $p_0 = 100/1325 = 0.0755$      $w_0 = 1/p_0 = 13.25$   
 $p_1 = 150/1684 = 0.0891$      $w_1 = 1/p_1 = 11.23$   
 $p_2 = 150/1346 = 0.1114$      $w_2 = 1/p_2 = 8.97$   
 $p_3 = 100/335 = 0.2985$      $w_3 = 1/p_3 = 3.35$

That is information from each of the youngest should weight by 13.25 and information from the each of the oldest should weight by 3.35.  
Sampling weights can be used in many Stata commands:

```
mean sbp [pw= sampw]
Mean estimation      Number of obs   =      500
-----+-----
      |      Mean   Std. Err.   [95% Conf. Interval]
-----+-----
sbp |  132.6242   1.032943    130.5947    134.6536
-----+-----
```

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**Missing values - not by design**

Most often the missing is **not per design** and both in the **outcome** and in the **covariates**:

id	y	x <sub>1</sub>	x <sub>2</sub>	x <sub>3</sub>
1	o	o	o	o
2	o	m	o	o
3	m	o	o	o
4	m	m	o	o
5	o	o	o	o
6	o	m	m	o

o observed  
m observed

Here we have only **complete data** on 2 persons, but partial information on 4 additoinal persons.

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Missing values – not by design

If the missing is **completely at random**, then the analysis of the complete cases will be unbiased.

If this is not the case, then complete data analysis can give biased estimates.

If the data is **missing at random**, then it is **in theory** possible to make an unbiased analysis of all the data.

id	y	x <sub>1</sub>	x <sub>2</sub>	x <sub>3</sub>
1	o	o	o	o
2	o	m	o	o
3	m	o	o	o
4	m	m	o	o
5	o	o	o	o
6	o	m	m	o

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Imputation

One way to try solve the problem with missing is to **fill in** the data for the missing values and then make the analysis on the whole data set with the **'imputed'** values.

The imputation can be done in many ways.

One way is to fill in an "average" value.

This could be the total average of the observed values for the specific variable or the average in a **relevant subgroup**.

This method will not in general solve the bias problem.

And of course the **standard error** stated in the output, when you analyse the imputed data set, is **wrong**.

id	y	x <sub>1</sub>	x <sub>2</sub>	x <sub>3</sub>
1	o	o	o	o
2	o	a <sub>1</sub>	o	o
3	a <sub>y</sub>	o	o	o
4	a <sub>y</sub>	a <sub>1</sub>	o	o
5	o	o	o	o
6	o	a <sub>1</sub>	a <sub>2</sub>	o

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The missing SBP example

Imputation by **observed mean** in age group:

```
bysort agegrp: egen msbp=mean(sbp)
generate isbp=msbp
replace isbp=msbp if missing(sbp)
```

```
mean isbp
Mean estimation      Number of obs   =    4690
-----+-----
            |      Mean   Std. Err.   [95% Conf. Interval]
-----+-----
isbp |    132.6242   .1627486   132.3051   132.9432
-----+-----
```

Correct mean, but a much too small standard error – incorrectly **assuming 4690 independent observations**.

Correct analysis using sampling weights:

```
mean sbp [pw=sampw]
Mean estimation      Number of obs   =    500
-----+-----
            |      Mean   Std. Err.   [95% Conf. Interval]
-----+-----
sbp |    132.6242   1.032943   130.5947   134.6536
-----+-----
```

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Imputation – random multiple

A fixed imputation will not take into account the random variation of the unobserved observation or the uncertainty of the parameters.

Imputation methods should add some random variation to the imputed data.

For that we need a **statistical model** for the missing data.

In **multiple imputations** one generates **several imputed** data sets.

For each imputed data set one fit the model of interest.

The point estimate, then the average across the imputed data sets.

One tricky thing is **calculation of the standard errors**.

id	y	x <sub>1</sub>	x <sub>2</sub>	x <sub>3</sub>
1	o	o	o	o
2	o	m	o	o
3	m	o	o	o
4	m	m	o	o
5	o	o	o	o
6	o	m	m	o

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Multiple imputations

Questions:

How to find the models from which to generate the missing data?

How should you handle missing data in this process?

How to find the uncertainty (standard errors) of the estimates?

Bookkeeping.

Most important: Missing at random is required!

id	y	x <sub>1</sub>	x <sub>2</sub>	x <sub>3</sub>
1	o	o	o	o
2	o	m	o	o
3	m	o	o	o
4	m	m	o	o
5	o	o	o	o
6	o	m	m	o

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The missing SBP example

```
use sbpdata,clear
mi set mlong
mi register imputed sbp
(4190 m=0 obs. now marked as incomplete)

mi impute regress sbp i.agegrp, add(20)
```

Univariate imputation                      Imputations =        20  
Linear regression                                added =        20  
Imputed: m=1 through m=20                        updated =        0

variable	complete	incomplete	imputed	total
sbp	500	4190	4190	4690

(complete + incomplete = total; imputed is the minimum across m of the number of filled in observations.)

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The missing SBP example

```
codebook, comp
```

Variable	Obs	Unique	Mean	Min	Max	Label
sbp	84300	83383	132.3204	44.52609	270	Systolic Blood Pressure
id	88490	4690	2352.429	1	4699	
agegrp	88490	4	1.107481	0	3	
_mi_id	88490	4690	2357.795	1	4690	
_mi_miss	4690	2	.8933902	0	1	
_mi_m	88490	21	9.943496	0	20	

```
sum if _mi_m==1
```

Variable	Obs	Mean	Std. Dev.	Min	Max
sbp	4190	131.2507	21.65931	59.92363	209.6556
id	4190	2352.611	1359.59	2	4699
agegrp	4190	1.105251	.8895275	0	3
_mi_id	4190	2358.483	1331.661	101	4690
_mi_miss	0				
_mi_m	4190	1	0	1	1

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The missing SBP example

```
. table agegrp if _mi_m>0, c(count sbp mean sbp sd sbp)
```

agegrp	N(sbp)	mean(sbp)	sd(sbp)	
0-	24,500	121.5843	22.32535	20*1225=24500
40-	30,680	131.1271	22.37045	
50-	23,920	141.2539	22.4434	
60-	4,700	150.2313	22.19089	20*235=4700

```
. table agegrp if _mi_m==0,c(count sbp mean sbp sd sbp)
```

agegrp	N(sbp)	mean(sbp)	sd(sbp)	
0-	100	122.18	15.4327	
40-	150	130.85	22.2366	
50-	150	140.93	22.4819	
60-	100	149.51	26.9251	

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The missing SBP example

mi estimate: mean sbp

Multiple-imputation estimates

Imputations = 20

Mean estimation

Number of obs = 4690

Average RVI = 7.4275

Complete DF = 4689

DF adjustment: Small sample

DF: min = 23.43

Within VCE type: ANALYTIC

avg = 23.43

max = 23.43

	Mean	Coef.	Std. Err.	t	P> t	[95% Conf. Interval]
sbp	132.6799	1.017506	130.40	0.000	130.5772	134.7826

Correct analysis using sampling weights:

mean sbp [pw=sampw]

Mean estimation

Number of obs = 500

	Mean	Std. Err.	[95% Conf. Interval]
sbp	132.6242	1.032943	130.5947 134.6536

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A more complicated example

use sbp2data,clear

codebook,comp

Variable	Obs	Unique	Mean	Min	Max	Label
sex	4188	2	1.566141	1	2	Sex
sbp	4216	112	132.6945	80	270	Systolic Blood Pressure
dbp	4281	67	82.62766	40	148	Diastolic Blood Pressure
sc1	4192	244	228.2011	115	568	Serum Cholesterol
age	4245	37	46.0636	30	66	Age in Years
bmi	4218	245	25.63148	16.2	57.6	Body Mass Index
id	4690	4690	2349.172	1	4699	

regress sbp age i.sex

Source	SS	df	MS	Number of obs	=	3,406
Model	281261.425	2	140630.713	F(2, 3403)	=	320.62
Residual	1492627.36	3,403	438.621029	Prob > F	=	0.0000
Total	1773888.79	3,405	520.96587	R-squared	=	0.1586
				Adj R-squared	=	0.1581
				Root MSE	=	20.943

	Coef.	Std. Err.	t	P> t	[95% Conf. Interval]
sbp	1.072026	.0423621	25.31	0.000	.9889686 1.155084
age	0	(base)			
sex					
Male	.2701054	.7247534	0.37	0.709	-1.150891 1.691101
Female	83.39557	2.017962	41.33	0.000	79.43903 87.35211
_cons					

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A more complicated example

mi set mlong

mi register imputed sbp age sex dbp bmi sc1

(2201 m=0 obs. now marked as incomplete)

mi describe

Style: mlong

last mi update 24nov2016 16:43:28, 0 seconds ago

Obs.: complete 2,489

incomplete 2,201 (M = 0 imputations)

total 4,690

Vars.: imputed: 6; sbp(474) age(445) sex(502) dbp(409) bmi(472) sc1(498)

passive: 0

regular: 0

system: 3; \_mi\_m \_mi\_id \_mi\_miss

(there is one unregistered variable; id)

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mi misstable pattern, freq

Missing-value patterns

(1 means complete)

Frequency	Pattern	1	2	3	4	5	6
2,489		1	1	1	1	1	1
314		1	1	1	1	0	1
301		1	1	1	1	1	0
281		1	1	1	0	1	1
278		1	1	0	1	1	1
253		1	0	1	1	1	1
243		0	1	1	1	1	1
42		1	1	1	0	0	1
37		1	0	1	1	1	0
37		1	1	1	0	1	0
36		1	1	0	1	1	0
35		1	1	1	1	0	0
34		0	0	1	1	1	1
33		1	0	0	1	1	1
32		0	1	0	1	1	1
30		1	1	0	0	1	1
28		1	1	0	1	0	1
27		1	0	1	0	1	1
25		0	1	1	0	1	1
25		0	1	1	1	1	0
25		1	0	1	1	0	1

21	0	1	1	1	0	1
7	1	0	0	1	0	1
5	0	0	0	1	1	1
5	1	1	0	1	0	0
4	0	0	1	0	1	1
4	0	1	0	0	1	1
4	0	1	1	1	0	0
4	1	0	1	0	1	0
4	1	1	0	0	1	0
3	1	0	0	0	1	1
3	1	0	1	0	0	1
3	1	0	1	1	0	0
2	0	0	1	1	0	1
2	0	1	1	0	0	1
2	0	1	1	0	1	0
2	1	0	0	1	1	0
2	1	1	1	0	0	0
1	0	0	1	0	1	0
1	0	0	1	1	0	0
1	0	0	1	1	1	0
1	0	1	0	0	0	1
1	0	1	0	1	0	1
1	0	1	0	1	1	0
1	1	1	0	0	0	0
1	1	1	0	0	0	1

4,690

Variables are (1) dbp (2) age (3) bmi (4) sbp (5) sc1 (6) sex

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Linear regression models for continuous and binary data : Note 6

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A more complicated example

```
mi impute chained ///
(regress,include( i.sex age bmi dbp scl))sbp ///
(regress,include( i.sex age bmi sbp scl))dbp ///
(regress,include( i.sex age bmi scl))bmi ///
(regress,include( i.sex age bmi sbp dbp scl))age ///
(regress,include( i.sex age bmi scl))scl ///
(logit, include( age bmi ))sex ///
```

add(100) noimputed

Conditional models:

```
dbp: regress dbp i.sex age bmi sbp scl
age: regress age i.sex bmi sbp dbp scl
bmi: regress bmi i.sex age scl
sbp: regress sbp i.sex age bmi dbp scl
scl: regress scl i.sex age bmi
sex: logit sex age bmi
```

Performing chained iterations ...

Multivariate imputation	Imputations =	100
Chained equations	added =	100
Imputed: m=1 through m=100	updated =	0
Initialization: monotone	Iterations =	1000
	burn-in =	10

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A more complicated example

Variable	Observations per m			
	Complete	Incomplete	Imputed	Total
sbp	4216	474	474	4690
dbp	4281	409	409	4690
bmi	4218	472	472	4690
age	4245	445	445	4690
scl	4192	498	498	4690
sex	4188	502	502	4690

(complete + incomplete = total; imputed is the minimum across m of the number of filled-in observations.)

codebook, comp

Variable	Obs	Unique	Mean	Min	Max	Label
-						
sbp	224316	47338	132.3185	47.08539	270	Systolic Blood Pressure
dbp	224381	40808	82.44368	40	148	Diastolic Blood Pressure
scl	224292	49896	227.064	34.90916	568	Serum Cholesterol
age	224345	44422	45.95276	12.26457	82.2043	Age in Years
bmi	224318	47293	25.52148	9.895696	57.6	Body Mass Index
id	224790	4690	2348.082	1	4699	
sex	224288	2	.5676273	0	1	RECODE of koen (Sex)
_mi_m	224790	101	49.44637	0	100	
_mi_id	224790	4690	2329.055	1	4690	
_mi_miss	4690	2	.4692964	0	1	

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A more complicated example

```
mi estimate: regress sbp age i.sex
```

Multiple-imputation estimates

Linear regression	Imputations =	100
	Number of obs =	4,690
	Average RVI =	0.1130
	Largest FMI =	0.1394
	Complete DF =	4687
DF adjustment: Small sample	DF: min =	2,256.34
	avg =	2,715.41
	max =	3,031.05
Model F test: Equal FMI	F( 2, 3480.5) =	396.38
Within VCE type: OLS	Prob > F =	0.0000

	sbp	Coef.	Std. Err.	t	P> t	[95% Conf. Interval]
age		1.072957	.0376538	28.50	0.000	.9991277 1.146787
sex						
Male		0 (base)				
Female		.2033005	.6617939	0.31	0.759	-1.094488 1.501089
_cons		83.29757	1.802549	46.21	0.000	79.76314 86.83199

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Clustered data / data with several random components

Dichotomous outcome

A different outcome:

$$H_{fpd} = \begin{cases} 1 & \text{if the person has hayfewer} \\ 0 & \text{else} \end{cases}$$

A statistical model:

Systematic part

$$\text{logit}(H_{fpd} = 1) = \beta_0 + \beta_I \cdot I + \beta_U \cdot U + \beta_A \cdot A + \beta_S \cdot S + \beta_G \cdot G$$

$$+ F_f + P_{fp} + X_{m_i}$$

Random part

This is not needed due to the binomial error

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**Clustered data / data with several random components**  
**Dichotomous outcome**

$$\text{logit}(H_{jpd} = 1) = \beta_0 + \beta_I \cdot I + \beta_U \cdot U + \beta_A \cdot A + \beta_S \cdot S + \beta_G \cdot G + F_f + P_{fp}$$

That is, an ordinary logistic regression + **random components**.

- A **generalized linear mixed model**
- A **multilevel model for dichotomous outcome**

Comments 1:

- It is **important** to include the **relevant random components** in the model.
- 'Multilevel models' is **essential** in medical/epidemiological research.

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**Clustered data / data with several random components**  
**Dichotomous outcome**

Comments 2:

- The theory and insight into the models for non-normal data are **not yet fully developed**.
- The main problem being that it is very difficult to find **valid (unbiased) estimates**.
- Several software programs **falsely claim** to estimate the models.
- Some programs like Stata and NLwin can give you valid estimates if you take care and have **a lot of data**.

**Advice:**

Do not try to estimate this kind of models without consulting a specialist.

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**Clustered data / data with one random components**  
**Dichotomous outcome**

If the models only involve **one random component**, e.g. **variation between families** or between **GP's**, then methods exist which can **adjust the standard errors**.

Remember that if the **data contains clusters**, then the precision of the estimates are overestimated, that is, the reported **standard errors are too small**.

So-called **robust methods** or **sandwich estimates** of the standard errors will (try to) adjust for this problem.

Only a **few** programs have this option - Stata does!

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