IDA Assignment 3

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Q1.

Consider the nhanes dataset in mice. For more information please type help(nhanes) in the R console.

a)

(2 marks) What percentage of the cases is incomplete?

Answer:

The percentage of incomplete cases is 48

b)

(4 marks) Impute the data with mice using the defaults with seed=1, in step 2 predict bmi from age, hyp, and chl by the normal linear regression model, and then pool the results. What are the proportions of variance due to the missing data for each parameter? Which parameters appear to be most affected by the nonresponse?

```
pool1 <- pool(with(mice(nhanes, printFlag = F, seed = 1), lm(bmi ~ as.factor(age) + hyp + chl)))
kable(pool1$pooled[,c(1,3,7,8,10)], caption = "Imputation with seed=1") %>%
kable_styling(latex_options = "hold_position")
```

Table 1: Imputation with seed=1

term	estimate	dfcom	df	lambda
(Intercept)	16.0426979	20	9.743911	0.3071661
as.factor(age)2	-5.2508182	20	5.496624	0.5204870
as.factor(age)3	-6.8779315	20	3.466705	0.6819661
hyp	2.4662556	20	8.463472	0.3606031
chl	0.0544513	20	8.530893	0.3576185

c)

(4 marks) Repeat the analysis for $seed \in \{2, 3, 4, 5, 6\}$. Do the conclusions remain the same?

Table 2: Imputation with seed=2,3,4,5,6

parameters	seed=2	seed=3	seed=4	seed=5	seed=6
(Intercept)	0.4144454	0.2772900	0.1315114	0.4855733	0.4168136
age	0.4033924	0.5895051	0.2189333	0.4511896	0.6549523
hyp	0.1430995	0.4101152	0.1961083	0.5942866	0.2960364
chl	0.2959966	0.5621346	0.3305334	0.2346065	0.5196295

d)

(4 marks) Repeat the analysis with M = 100 with the same seeds. Would you prefer these analyses over those with M = 5? Explain why.

Answer:

```
pool1 <- pool(with(mice(nhanes, m = 100, printFlag = F, seed = 1),</pre>
                    lm(bmi ~ age + hyp + chl)))
pool2 <- pool(with(mice(nhanes, m = 100, printFlag = F, seed = 2),</pre>
                    lm(bmi ~ age + hyp + chl)))
pool3 <- pool(with(mice(nhanes, m = 100, printFlag = F, seed = 3),</pre>
                    lm(bmi ~ age + hyp + chl)))
pool4 <- pool(with(mice(nhanes, m = 100, printFlag = F, seed = 4),</pre>
                    lm(bmi ~ age + hyp + chl)))
pool5 <- pool(with(mice(nhanes, m = 100, printFlag = F, seed = 5),</pre>
                    lm(bmi ~ age + hyp + chl)))
pool6 <- pool(with(mice(nhanes, m = 100, printFlag = F, seed = 6),</pre>
                    lm(bmi ~ age + hyp + chl)))
parameters <- c("(Intercept)", "age", "hyp", "chl")</pre>
df <- data.frame(parameters, pool1$pooled[,10], pool2$pooled[,10],</pre>
                  pool3$pooled[,10], pool4$pooled[,10],
                  pool5$pooled[,10], pool6$pooled[,10])
colnames(df) <- c("parameters", "seed=1", "seed=2", "seed=3", "seed=4",</pre>
                   "seed=5", "seed=6")
kable(df, caption="Imputation with seed=1,2,3,4,5,6 and M=100") %>%
 kable_styling(latex_options = "hold_position")
```

Table 3: Imputation with seed=1,2,3,4,5,6 and M=100

parameters	seed=1	seed=2	seed=3	seed=4	seed=5	seed=6
(Intercept)	0.2290445	0.1882474	0.2199607	0.2144722	0.2294356	0.2472607
age	0.4324680	0.4031077	0.3093072	0.3943223	0.3322570	0.4430300
hyp	0.2915346	0.2825108	0.2425105	0.2565132	0.2893046	0.2860700
chl	0.3217837	0.2939693	0.3281911	0.2835232	0.2461956	0.3113085

The pooled estimates is more stable when the value of M is higher. Thus, we would prefer M=100 than M=5

(15 marks) Each of the 100 datasets contained in the file dataex2.Rdata was generated in the following way

$$y_i|x_i \stackrel{\text{ind.}}{\sim} N(\beta_0 + \beta_1 x_i, 1), \quad x_i \stackrel{\text{ind.}}{\sim} Unif(-1, 1), \quad \beta_0 = 1, \quad \beta_1 = 3$$

for $i=1,\ldots,100$. Additionally, some of the responses were set to be missing using a MAR mechanism. The goal of this exercise is to study the effect that acknowledging/not acknowledging parameter uncertainty when performing step 1 of multiple imputation might have on the coverage of the corresponding confidence intervals. Further suppose that the analysis of interest in step 2 is to fit the regression model that was used to generate the data, i.e., a normal linear regression model where the response is y and the covariate is x. With the aid of the mice package, calculate the empirical coverage probability of the 95% confidence intervals for β_1 under the following two approaches: stochastic regression imputation and the corresponding bootstrap based version. Comment. For both approaches, please consider m=20 and seed=1. NOTE 1: In order to calculate the empirical coverage probability, you only need to compute the proportion of the time (over the 100 intervals) that the interval contains the true value of the parameter. NOTE 2: The data are stored in an array structure such that, for instance, dataex2[, , 1], corresponds to the first dataset (which has 100 rows and 2 columns, with the first column containing the values of x and the second the values of y).

Answer:

```
# initialize a counter
count <- 0
for (i in 1:nrow(dataex2)) {
    #impute values for the ith dataset using M=20
    impute.sri <- mice(dataex2[, , i], m = 20, method = "norm.nob", printFlag = F, seed = 1)
    fit.sri <- with(impute.sri, lm(Y ~ X)) #step 2
    pool.sri <- pool(fit.sri) # step 3
    summary.sri <- summary(pool.sri, conf.int = TRUE)
    if (summary.sri[2, 7] <= 3 & summary.sri[2, 8] >= 3) {
        count <- count + 1 #add to the counter if the the value of beta1 is contained in the
        #confidence interval
    }
}
ecp.sri <- count/nrow(dataex2)
cat("the proportion of the time for Stochastic Imputation is", ecp.sri)</pre>
```

the proportion of the time for Stochastic Imputation is 0.88

the proportion of the time for Bootstrap is 0.95

Q3.

(9 marks) Show that for a linear (in the coefficients) regression model, the following two strategies coincide:

- (i) Computing the predicted values (point estimates) from each fitted model in step 2 and then pooling them according to Rubin's rule for point estimates (i.e., averaging the predicted values across the imputed datasets).
- (ii) Pooling the regression coefficients from each fitted model in step 2 using Rubin's rule for point estimates and then computing the predicted values afterwards

Answer:

We consider a linear regression model given a dataset as $\{y_i, x_{1i}, \dots, x_{ni}\}$

$$y_i = \beta_0 + \beta_1 x_{1i} + \dots + \beta_n x_{ni} + \varepsilon_i, \quad \varepsilon_i \sim N(0, \sigma^2)$$

Now we look into Case (i), we compute the predicted values for each fitted model from step 2. Then we obtain as below,

$$\hat{y}_i^{(m)} = \hat{\beta}_0^{(m)} + \hat{\beta}_1^{(m)} x_{1i} + \dots + \hat{\beta}_n^{(m)} x_{ni}$$

Then we pool them according to Rubin's rule for point estimates.

$$\bar{y}_{i} = \frac{1}{M} \sum_{i=1}^{M} \hat{y}^{(m)}
= \frac{1}{M} \sum_{i=1}^{M} \left(\hat{\beta}_{0}^{(m)} + \hat{\beta}_{1}^{(m)} x_{1i} + \dots + \hat{\beta}_{n}^{(m)} x_{ni} \right)
= \frac{1}{M} \sum_{i=1}^{M} \hat{\beta}_{0}^{(m)} + \frac{1}{M} \sum_{i=1}^{M} \hat{\beta}_{1}^{(m)} x_{1i} + \dots + \frac{1}{M} \sum_{i=1}^{M} \hat{\beta}_{n}^{(m)} x_{ni}
= \bar{\beta}_{0} + \bar{\beta}_{1} x_{1i} + \dots + \bar{\beta}_{n} x_{ni}$$

Now, let us consider Case (ii) to validate if they coincide. We pool the regression coefficients from each fitted model in step 2 using Rubin's rule for point estimates.

$$\bar{\beta}_0 = \frac{1}{M} \sum_{i=1}^M \hat{\beta}_0^{(m)}$$
:

$$\bar{\beta}_n = \frac{1}{M} \sum_{i=1}^M \hat{\beta}_n^{(m)}$$

Then we can compute the predicted values as follow

$$\bar{y}_i = \bar{\beta_0} + \bar{\beta_1} x_{1i} + \dots + \bar{\beta_n} x_{ni}$$

As shown above, the order of the computation of predicted values for each fitted model in step 2 and pooling according to Rubin's rule for point estimates do not matter mathematically. Therefore, we conclude here by saying that both cases coincide.

Q4.

The goal of this exercise is to study different ways of using mice when the analysis model of interest/substantive model involves an interaction term between incomplete variables. The model used to generate the data (available in dataex4.Rdata), which corresponds to our model of interest in step 2, was the following one:

$$y_i = \beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + \beta_3 x_{1i} x_{2i} + \varepsilon_i, x_{1i} \stackrel{\text{iid}}{\sim} \mathrm{N}(0,1), \quad x_{2i} \stackrel{\text{iid}}{\sim} \mathrm{N}(1.5,1), \quad \varepsilon_i \stackrel{\text{iid}}{\sim} \mathrm{N}(0,1)$$

for $i=1,\ldots,1000$, $\beta_0=1.5$, $\beta_1=1$, $\beta_2=2$ and $\beta_3=1$. Additionally, missingness was imposed on y and x_1 and so the interaction variable x_1x_2 also has missing values, although the missingness in this interaction variable is induced by the missing in the covariate x_1 . In the following, please use M=50 and seed=1.

```
kable(head(dataex4), caption="dataex4 observation") %>%
kable_styling(latex_options = "hold_position")
```

у	x1	x2
NA	NA	1.0983213
4.609476	0.0265944	1.1290673
NA	-1.5165531	1.0748538
1.718231	-1.3626533	1.9411515
NA	1.1784892	0.8272496
1.820674	-0.9341513	1 9166941

Table 4: dataex4 observation

a)

(6 marks) By only imputing the y and x_1 variables in step 1, provide the estimates of β_1, β_2 , and β_3 along with 95% confidence intervals. Comment. Note that this approach where the interaction variable is left outside the imputation process and calculated afterwards in the analysis model, is known as Impute, then transform

Table 5: Summary Statistics of Imputation of y and x_1

term	estimate	std.error	2.5 %	97.5 %
(Intercept)	1.5929831	0.0954133	1.404501	1.7814655
x1	1.4112333	0.0973291	1.219397	1.6030697
x2	1.9658191	0.0532322	1.860657	2.0709812
x1:x2	0.7550367	0.0570146	0.642302	0.8677715

b)

(10 marks) Now, start by calculating the interaction variable in the incomplete data and append it as a variable to your dataset. Then, use *passive imputation* to impute the interaction variable. Provide the estimates of β_1, β_2 , and β_3 along with 95% confidence intervals. Comment.

```
x1 <- dataex4$x1; x2 <- dataex4$x2; dataex4$x1x2 <- x1*x2
impute.null <- mice(dataex4, maxit = 0)</pre>
method <- impute.null$method</pre>
method["x1x2"] \leftarrow "\sim I(x1*x2)"
pred <- impute.null$predictorMatrix</pre>
pred[c("x1", "x2"), "x1x2"] <- 0</pre>
visit.seq <- impute.null$visitSequence</pre>
visit.seq
## [1] "y"
                      "x2"
               "x1"
                              "x1x2"
impute.passive <- mice(dataex4, method = method, predictorMatrix = pred,</pre>
                 visitSequence = visit.seq, m = 50, seed = 1, printFlag = FALSE)
pool.passive <- pool(with(impute.passive, lm(y \sim x1 + x2 + x1*x2)))
kable(summary(pool.passive, conf.int=TRUE)[,c(1,2,3,7,8)],
      caption = "Summary Statistics of Imputation of $y$ and $x_1$") %>%
  kable_styling(latex_options = "hold_position")
```

Table 6: Summary Statistics of Imputation of y and x_1

term	estimate	std.error	2.5 %	97.5 %
(Intercept)	1.5534782	0.0884221	1.3788626	1.7280939
x1	1.1926170	0.0958435	1.0034980	1.3817360
x2	1.9964402	0.0493658	1.8989468	2.0939336
x1:x2	0.8740573	0.0567852	0.7615712	0.9865434

c)

(10 marks) Now that you have already appended the interaction variable to the dataset, impute it as it was just another variable (or like any other variable) in the dataset and use this variable for the interaction term in step 2. Provide the estimates of β_1, β_2 and β_3 along with 95% confidence intervals. Comment.

Table 7: Summary Statistics of Imputation of y and x_1

term	estimate	std.error	2.5 %	97.5 %
(Intercept)	1.499714	0.0782144	1.3452011	1.654227
x1	1.003930	0.0822837	0.8414967	1.166363
x2	2.026180	0.0437161	1.9398113	2.112548
x1x2	1.017793	0.0442807	0.9303479	1.105238

d)

(6 marks) What is the obvious conceptual drawback of the $just\ another\ variable$ approach for imputing interactions?

 $\mathbf{Answer:}$

Q_5

(30 marks) The file NHANES2.Rdata contains a subset of data from the *National Health and Nutrition Examination Survey* (NHANES), whose goal is to assess the health and nutritional status of adults and children in the United States. The variables in the dataset are the following:

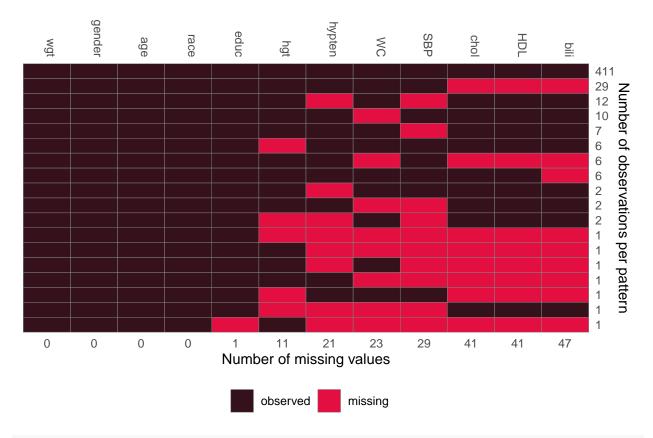
- wgt: weight in kg,
- gender: male vs female,
- bili: bilirubin concentration in mg/dL,
- age: in years,
- chol: total serum cholestrol in mg/dL,
- HDL: High-density lipoprotein cholestrol in mg/dL,
- hgt: height in metres,
- educ: educational status; 5 ordered categories,
- race: 5 unordered categories,
- SBP: systolic blood pressure in mmHg,
- hypten: hyptertensive status; binary,
- WC: waist circumference in cm.

The analysis of interest is the following:

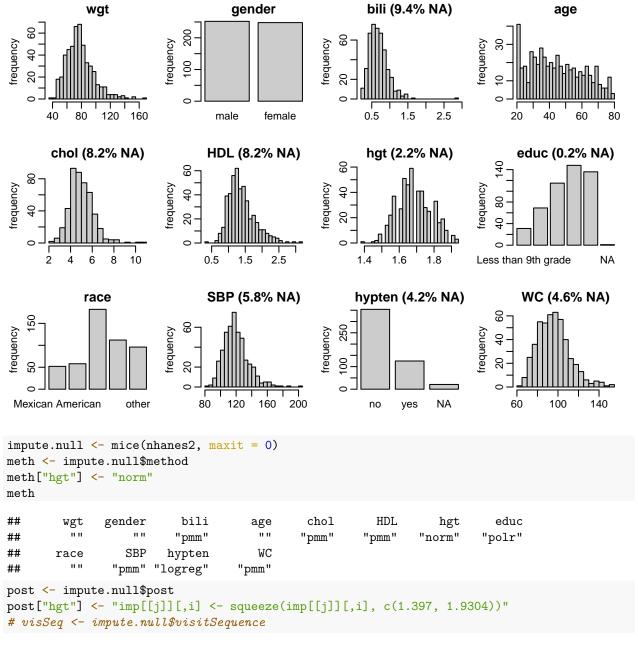
wgt =
$$\beta_0 + \beta_1$$
gender + β_2 age + β_3 hgt + β_4 WC + ε , $\varepsilon \sim N(0, \sigma^2)$.

Using multiple imputation and conducting all necessary checks, report your findings.

```
nhanes2 <- NHANES2
md_pattern(nhanes2, pattern = FALSE, color = c('#34111b', '#e30f41'))</pre>
```



```
par(mar = c(3,3,2,1), mgp = c(2,0.6,0))
plot_all(nhanes2, breaks = 30, ncol = 4)
```



While inspecting the missing data pattern, we found 411 observations with observed values on all 12 variables. Also, 10 observations for which on WC is missing, 6 observations for which on hgt is missing Interestingly, we can observe that there are 29 observations for which on chol, HDL and bili are missing.

We can visualise the variables' distributions using plot_all(). In the plot above, depicting the distribution of the observed data for the different variables, we could appreciate that hgt following a normal distribution is possibly not a completely unreasonable idea. Let us then change the default from predictive mean matching method (pmm) to norm for the variable hgt.

However, we need to be careful, because we do not want to risk imputing a negative value for the height. With the below syntax all imputed values of hgt that are outside the interval (min(nhanes2\$hgt), max(nhanes2\$hgt)) will be set to those limiting values.

Since our model is limited to this, wgt = $\beta_0 + \beta_1$ gender + β_2 age + β_3 hgt + β_4 WC + ε , $\varepsilon \sim N(0, \sigma^2)$.. We will end our data preprocessing and proceed to the next them in tuning the hyperparameters for imputation

```
seed1.5 <- pool(with(mice(nhanes2, methods = meth, post = post,</pre>
                           m = 5, seed = 1, printFlag = FALSE),
                     lm(wgt ~ gender + age + hgt + WC)))
seed1.10 <- pool(with(mice(nhanes2, methods = meth, post = post,</pre>
                           m = 10, seed = 1, printFlag = FALSE),
                      lm(wgt ~ gender + age + hgt + WC)))
seed1.20 <- pool(with(mice(nhanes2, methods = meth, post = post,</pre>
                           m = 20, seed = 1, printFlag = FALSE),
                     lm(wgt ~ gender + age + hgt + WC)))
seed1.25 <- pool(with(mice(nhanes2, methods = meth, post = post,</pre>
                           m = 25, seed = 1, printFlag = FALSE),
                     lm(wgt ~ gender + age + hgt + WC)))
seed2.5 <- pool(with(mice(nhanes2, methods = meth, post = post, maxit = 20,</pre>
                           m = 5, seed = 2, printFlag = FALSE),
                     lm(wgt ~ gender + age + hgt + WC)))
seed2.10 <- pool(with(mice(nhanes2, methods = meth, post = post, maxit = 20,</pre>
                           m = 10, seed = 2, printFlag = FALSE),
                      lm(wgt ~ gender + age + hgt + WC)))
seed2.20 <- pool(with(mice(nhanes2, methods = meth, post = post, maxit = 20,</pre>
                           m = 20, seed = 2, printFlag = FALSE),
                      lm(wgt ~ gender + age + hgt + WC)))
seed2.25 <- pool(with(mice(nhanes2, methods = meth, post = post, maxit = 20,</pre>
                           m = 25, seed = 2, printFlag = FALSE),
                     lm(wgt ~ gender + age + hgt + WC)))
# parameters <- c("(Intercept)", "gender", "age", "hgt", "WC")</pre>
# df \leftarrow data.frame(parameters, seed1.5$pooled[,10], seed1.10$pooled[,10],
                   seed1.20$pooled[,10], seed1.25$pooled[,10],
                   seed2.5$pooled[,10], seed2.10$pooled[,10],
#
                   seed2.20$pooled[,10], seed2.25$pooled[,10])
#
\# colnames(df) <- c("parameters", "seed1.5", "seed1.10", "seed1.20", "seed1.25",
                     "seed2.5", "seed2.10", "seed2.20", "seed2.25")
# kable(df, caption="Imputation with various seed and m values") %>%
   kable_styling(latex_options = "hold_position")
kable(data.frame(summary(seed1.5, conf.int = TRUE)[, c(1, 2, 3, 7, 8)],
                  lambda = seed1.5$pooled[,10]), caption = "Seed=1 and m=5") %>%
 kable_styling(latex_options = "hold_position")
```

Table 8: Seed=1 and m=5

term	estimate	std.error	X2.5	X97.5	lambda
(Intercept)	-101.4588664	7.5938549	-116.3816321	-86.5361006	0.0195179
genderfemale	-1.4195844	0.8425729	-3.0758469	0.2366781	0.0355672
age	-0.1589451	0.0213354	-0.2008787	-0.1170115	0.0293297
hgt	53.0413585	4.3496144	44.4934595	61.5892575	0.0227495
WC	1.0225600	0.0223080	0.9787286	1.0663914	0.0049737

Table 9: Seed=1 and m=10

term	estimate	std.error	X2.5	X97.5	lambda
(Intercept)	-101.6700795	7.8551715	-117.1260787	-86.2140802	0.0937749
genderfemale	-1.3085502	0.8493953	-2.9787291	0.3616288	0.0667687
age	-0.1576161	0.0215356	-0.1999575	-0.1152746	0.0616516
hgt	52.9908125	4.4813711	44.1741136	61.8075113	0.0899366
WC	1.0244107	0.0222726	0.9806462	1.0681752	0.0168680

Table 10: Seed=1 and m=20

term	estimate	std.error	X2.5	X97.5	lambda
(Intercept)	-101.2897304	7.8011807	-116.6273935	-85.9520673	0.0822178
genderfemale	-1.3626948	0.8438271	-3.0211578	0.2957682	0.0527619
age	-0.1602819	0.0216544	-0.2028488	-0.1177150	0.0690126
hgt	52.7103557	4.4457568	43.9703249	61.4503865	0.0766459
WC	1.0265327	0.0222736	0.9827681	1.0702973	0.0122374

Table 11: Seed=1 and m=25

term	estimate	std.error	X2.5	X97.5	lambda
(Intercept)	-101.0547825	7.7120024	-116.2127029	-85.8968621	0.0624659
genderfemale	-1.3203389	0.8335918	-2.9583505	0.3176727	0.0279959
age	-0.1593655	0.0214497	-0.2015208	-0.1172102	0.0512832
hgt	52.5781406	4.4232119	43.8838452	61.2724359	0.0683664
WC	1.0259679	0.0224005	0.9819518	1.0699840	0.0233236

Table 12: Seed=2 and m=5

term	estimate	std.error	X2.5	X97.5	lambda
(Intercept)	-100.9966596	7.7300251	-116.2136672	-85.7796520	0.0752096
genderfemale	-1.3775163	0.8404887	-3.0302117	0.2751791	0.0471154
age	-0.1580193	0.0213131	-0.1999240	-0.1161147	0.0436862
hgt	52.5424942	4.4106181	43.8613258	61.2236627	0.0718380
WC	1.0259230	0.0222557	0.9821911	1.0696550	0.0142010

Table 13: Seed=2 and m=10

term	estimate	std.error	X2.5	X97.5	lambda
(Intercept)	-101.5186858	7.5571364	-116.3699779	-86.6673938	0.0311981
genderfemale	-1.3215513	0.8256067	-2.9438439	0.3007412	0.0185686
age	-0.1592563	0.0213802	-0.2012842	-0.1172285	0.0510297
hgt	52.8833997	4.3363259	44.3609058	61.4058936	0.0384335
WC	1.0255032	0.0222773	0.9817285	1.0692778	0.0197263

Table 14: Seed=2 and m=20

term	estimate	std.error	X2.5	X97.5	lambda
(Intercept)	-101.6993777	7.6823878	-116.7986779	-86.600078	0.0547441
genderfemale	-1.3663151	0.8370945	-3.0113631	0.278733	0.0398821
age	-0.1576975	0.0214614	-0.1998791	-0.115516	0.0557808
hgt	53.0055071	4.4023585	44.3525473	61.658467	0.0590317
WC	1.0246288	0.0223533	0.9807051	1.068553	0.0239163

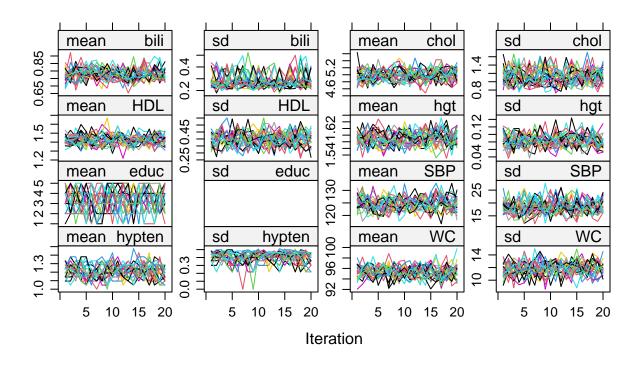
Table 15: Seed=2 and m=25

term	estimate	std.error	X2.5	X97.5	lambda
(Intercept)	-101.4521949	7.6290783	-116.4452338	-86.4591561	0.0472501
genderfemale	-1.3125011	0.8317349	-2.9468852	0.3218831	0.0303398
age	-0.1585538	0.0213320	-0.2004759	-0.1166316	0.0456505
hgt	52.8201428	4.3724292	44.2269381	61.4133475	0.0516546
WC	1.0255459	0.0223605	0.9816083	1.0694836	0.0243072

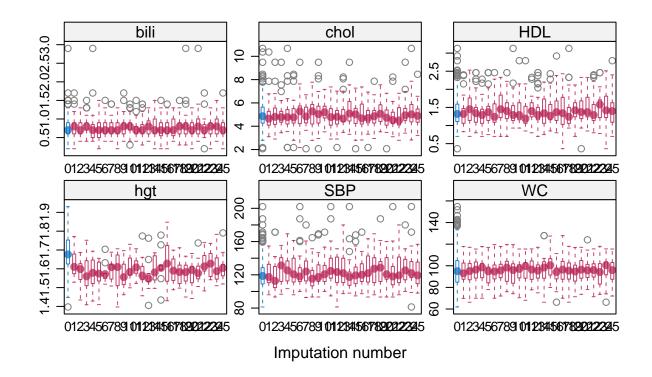
Looking at Tables above, we have the (pooled) estimates, standard errors, the confident intervals and lambda which get more stable as M increases and we can be more confident in any one specific run. Since we run the multiple imputation with a sufficiently large M, the results will with high probability only differ by a small amount. Thus we will choose $\mathtt{seed=1}$ with $\mathtt{M=25}$ and proceed to the multiple imputation.

NULL

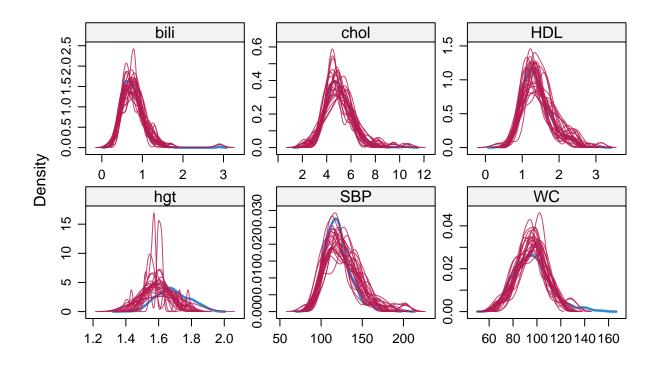
We can also confirm through loggedEvents that no problems occurred during the imputation.



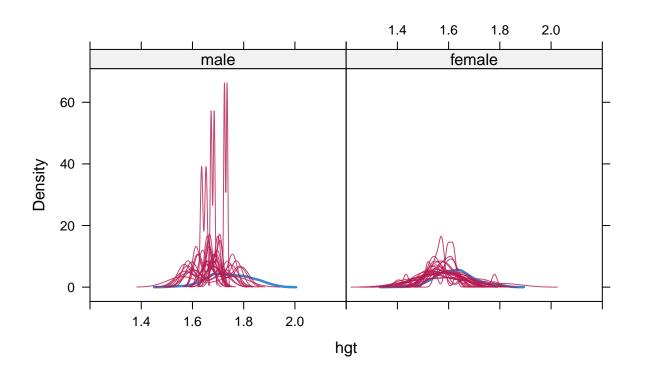
bwplot(imp)[c(2,4,5,6,7,8)]

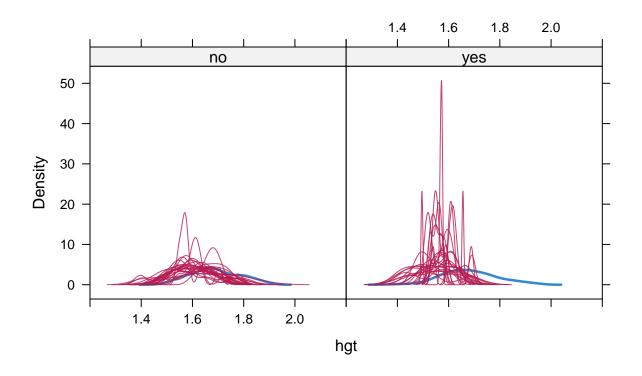


densityplot(imp)

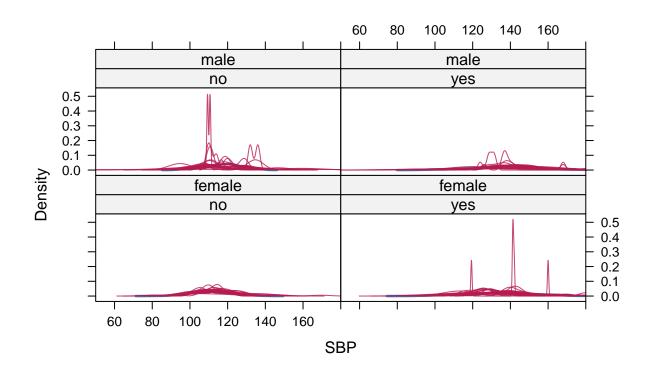


densityplot(imp,~hgt|gender, xlim = c(1.2, 2.2))

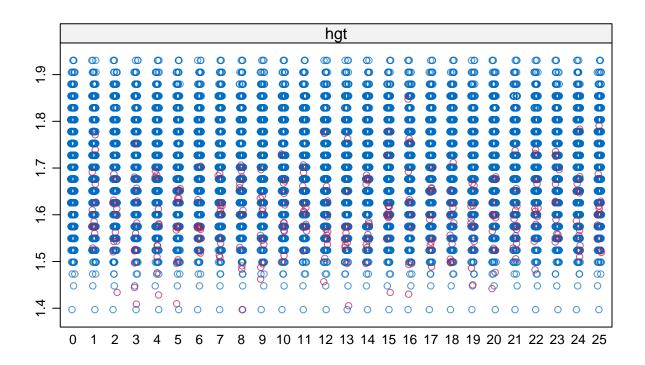


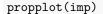


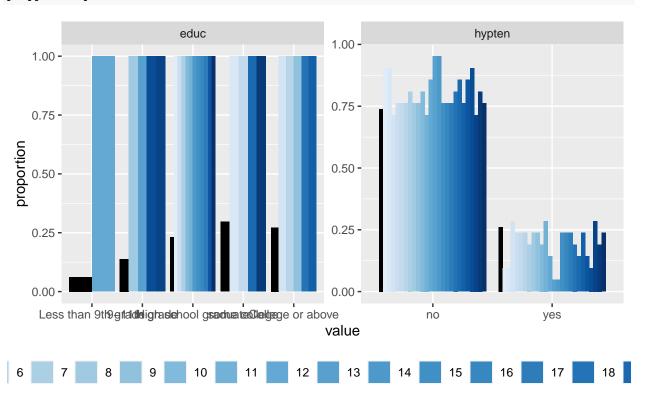
densityplot(imp,~SBP|hypten + gender, xlim=c(50, 180))



stripplot(imp)[6]







Now let us process the plots one by one to check if the mice() converged. Except for educ, rest of the variables are continuous are we proceed to the first plot. The traceplot indicates that the iterative algorithm appears to have converged for all variables that were imputed. The next plots are boxplots and density plots for the observed and imputed ones where we label them as blue and red respectively.

Note that we are using M=25 and because the density of the observed data is possibly plotted first, we can barely see it. However, we can generally confirm that most imputed datasets follow a similar distribution. The most outstanding plots are the ones from SBP and hgt as there is a shift towards lower values.

Specifically, we investigated with hgt conditional on the gender and hgt conditional on hypten. We can see that to a certain extent, both plots explain the differences between the observed and imputed values for hgt. This is because male and positive have much narrower distribution compared to wider distribution for female and negative. For SBP we could observe similarity. Using stripplot() we can double confirm that the imputation seems reasonable.

We observe an abnormal imputation pattern for the educ variable however this is **not** a cause for concern since we are only imputing a single missing value out of 500 cases. Meanwhile, for the hypten variable, we observe a reasonable amount between imputation variance but not enough such that the general distribution of this variable is lost. In summary, all imputations for variables with missing data have been performed successfully.

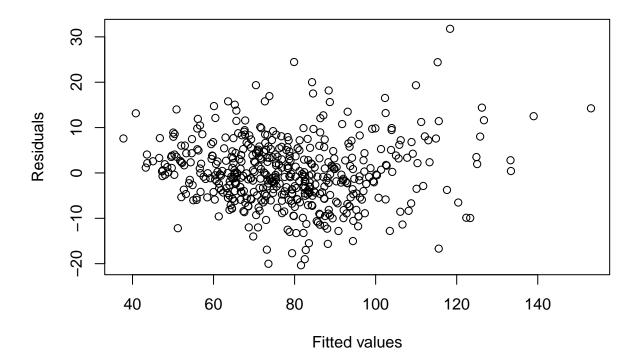
we can compare the distribution of the imputed values against the distribution of the observed categorical variables' values using propplot(). This compares the proportion of values in each category. This shows a large discrepancy between the observed and imputed data distributions for the educ. However educ only contains 1 missing variable and we can conclude that there is insufficient evidence to say that the imputation educ has problem. For the hypten, there is a reasonable proportion between the imputation variances.

All together, we can confirm that the multiple imputation is successful and proceed to analysis of the imputed data.

```
fit <- with(imp, lm(wgt ~ gender + age + hgt + WC))
summary(fit$analyses[[1]])</pre>
```

```
##
## Call:
## lm(formula = wgt ~ gender + age + hgt + WC)
##
## Residuals:
##
       Min
                1Q
                    Median
                                3Q
                                       Max
                                    31.774
##
   -20.349
            -4.568
                    -0.428
                             4.068
##
## Coefficients:
##
                  Estimate Std. Error t value Pr(>|t|)
## (Intercept)
               -100.83781
                              7.51251 -13.423
                                               < 2e-16 ***
  genderfemale
                  -1.36542
                              0.82182
                                       -1.661
##
                                                 0.0973
                  -0.15556
## age
                              0.02085
                                       -7.460 3.91e-13 ***
                                       12.203
                  52.42865
                              4.29633
                                               < 2e-16 ***
## hgt
## WC
                   1.02524
                              0.02210
                                       46.387
                                               < 2e-16 ***
##
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 7.192 on 495 degrees of freedom
## Multiple R-squared: 0.857, Adjusted R-squared: 0.8558
## F-statistic: 741.4 on 4 and 495 DF, p-value: < 2.2e-16
```

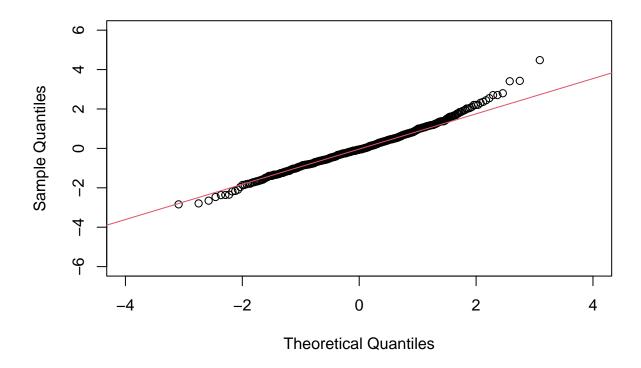
Looking at the summary we can check that only the coefficient for **genderfemale** has a p-value greater than 0.05. To this, we will check using Wald Test later for the importance of it.



In this fitted values versus residuals plot we can observe two clusters. The residuals are symmetric about zero, and we can suspect that maybe the homoscedastic assumption is slightly violated, but nothing that we cannot live with. I was however curious and decided to plot the response variable, SBP, against the other variables. They seem to indicate that the major cause of the two clusters is the hypertensive status.

```
qqnorm(rstandard(fit$analyses[[1]]), xlim = c(-4, 4), ylim = c(-6, 6))
qqline(rstandard(fit$analyses[[1]]), col = 2)
```

Normal Q-Q Plot



By observing the normal Q-Q plot, we do not see the deviance from the red dotted base line. Therefore, we can conclude that the data after imputation follows normal distribution and the normality assumption holds. Lastly, we will proceed to pooling the estimates with our imputed dataset.

Table 16: Summary Statistics

term	estimate	p.value	2.5 %	97.5 %
(Intercept)	-101.1180859	0.0000000	-116.0326895	-86.2034823
genderfemale	-1.3477467	0.1063107	-2.9844502	0.2889568
age	-0.1582103	0.0000000	-0.2004609	-0.1159597
hgt	52.6066777	0.0000000	44.0729374	61.1404180
WC	1.0256687	0.0000000	0.9818758	1.0694617

```
kable(pool.r.squared(pooled_ests, adjusted = TRUE), caption = "") %>%
kable_styling(latex_options = "hold_position")
```

Table 17:

	est	lo 95	hi 95	fmi
adj R^2	0.8563534	0.8308076	0.8783204	0.0261765

Looking at the Table, the adjusted R^2 values indicates good fit.

Now we conduct Wald Test to check the importance of the feature.

Table 18: Wald Test on each features

excluded.features	F.value	df1	df2	PF.	RIV
WC	2117.811020	1	488.6612	0.0000000	0.0174882
hgt	146.751120	1	472.7845	0.0000000	0.0394115
age	54.164973	1	447.6375	0.0000000	0.0621865
gender	2.618292	1	479.2870	0.1062959	0.0319663

We conducted the Wald test by excluding each feature one by one and used D1(). From the Table, we can see that gender is the only feature having p-value greater than 0.05. Although it is above 0.05, we can still consider its important but limiting it into 0.1 significance level. Thus, WC, hgt and age have more influence than gender here.