#### 432 Class 02 Slides

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### Today's Agenda

- Linear Regression with Categorical Predictors
- Building Two-Way ANOVA models with interaction
- Building Analysis of Covariance Models

We'll use some BRFSS/SMART data about Ohio residents to address the following questions.

- What is the association of diabetes diagnosis and smoking status on BMI?
- ② Does adjusting for subject's age affect our assessments?
- How does the subject's level of physical activity fit into this?

**Chapter 2** of the Course Notes builds the BRFSS/SMART data.

## Setup

```
knitr::opts_chunk$set(comment = NA)
options(width = 60) ## for the slides
library(here)
                        ## project management
library(knitr)
                        ## mostly for kable
library(mosaic)
                        ## mostly for favstats
library(patchwork)
                        ## combine plots
library(janitor)
                        ## mostly for tabyl
library(naniar)
                        ## missing data tools
library(simputation)
                        ## for single imputation
library(broom)
                        ## for tidying model output
                        ## as always (dplyr, ggplot2, etc.)
library(tidyverse)
theme_set(theme_bw())
                        ## my personal preference
```

• I used message = FALSE in the code chunk setup.

## Codebook of variables we'll select from smart\_ohio

```
smart_ohio <- read_csv(here("data/smart_ohio.csv"))
dim(smart_ohio)</pre>
```

[1] 7412 99

We'll sample 432 observations from smart\_ohio on these six variables. . .

Variable	Туре	Description
SEQNO bmi smoker	ID code Quantity 4 levels	we'll represent as a character body-mass index (in kg/m²) smoking status (we'll collapse to 3 levels)
dm_status activity age_imp	4 levels 4 levels Quantity	diabetes status (we'll collapse to 2) physical activity level (we'll re-level) age (imputed from groups, see Chapter 2)

# Create our working data set (day2)

- set seed for random sampling
- modify smart\_ohio data and place in new day2 tibble
- select our six variables
- take a random sample of 432 rows from the data
- onvert all character variables into factors
- o set ID code as a character variable

#### The day2 tibble, printed

#### day2

```
A tibble: 432 \times 6
   SEQNO
               bmi smoker dm status
                                        activity
                                                     age_imp
   <chr>
             <dbl> <fct>
                           <fct>
                                        <fct>
                                                       <int>
  2017000934
               NA
                    Never
                           No-Diabetes
                                        Inactive
                                                          71
 2 2017000411
              27.3 Never
                           No-Diabetes
                                        <NA>
                                                          69
3 2017001084
               19.2 Former No-Diabetes
                                        Inactive
                                                          NA
  2017000065
               20.5 Never
                           No-Diabetes
                                        Highly_Acti~
                                                          22
  2017000240
               25.1 Never
                           No-Diabetes
                                        Insufficien~
                                                          61
  2017000300
               24.8 Never
                           Pregnancy-I~
                                        Inactive
                                                          40
  2017001444
                           No-Diabetes
                                        Highly_Acti~
                                                          57
               31.2 Former
  2017000258
               29.4 Never
                           No-Diabetes
                                        Active
                                                          27
  2017000769
                                        Highly Acti~
                                                          71
               26.7 Former No-Diabetes
10 2017000247
               27.4 Former No-Diabetes
                                        Inactive
                                                          91
# ... with 422 more rows
```

# **Initial Data Checking (Quantities)**

min	Q1	median	Q3	max	mean	sd	n	missing
16.9	23.9	27.4	31.2	56.6	28.4	6.3	400	32

min	Q1	median	Q3	max	mean	sd	n	missing
 18	43	58.5	70	96	56.9	19	424	8

- Do the observed ranges of values look plausible in context?
- ② Are there missing values we need to deal with?

# Checking the Categorical Data (activity)

#### day2 %>% count(activity)

• What does <NA> mean? What should we do with this variable?

# Checking the Categorical Data (activity)

#### day2 %>% count(activity)

```
# A tibble: 5 x 2
activity n
<fct> <int>
1 Active 60
2 Highly_Active 114
3 Inactive 124
4 Insufficiently_Active 87
5 <NA> 47
```

- What does <NA> mean? What should we do with this variable?
- Shortly, we'll **reorder** these levels in a more sensible way (suggestions?) and then we'll have to deal with the missing values, somehow.

# Checking the Categorical Data (smoker)

```
day2 %>% count(smoker)
```

• OK. Some missing values to deal with. What else might we do here?

# Checking the Categorical Data (smoker)

```
day2 %>% count(smoker)
```

- OK. Some missing values to deal with. What else might we do here?
- Shortly, we'll **collapse** this from 4 to 3 levels (how?)

# Checking the Categorical Data (smoker)

#### day2 %>% count(smoker)

- OK. Some missing values to deal with. What else might we do here?
- Shortly, we'll collapse this from 4 to 3 levels (how?)
- I think we'll go with Current, Former and Never

# Checking the Categorical Data (dm\_status)

```
day2 %>% count(dm_status)
```

Next Steps?

# Checking the Categorical Data (dm\_status)

```
day2 %>% count(dm_status)
```

- Next Steps?
- Shortly, we'll collapse this to two levels (how might we do that?) and then we'll deal with the missing information.

## Re-ordering and collapsing in day2

```
day2 <- day2 %>%
    mutate(activity =
               fct_relevel(activity, "Highly_Active",
                            "Active", "Insufficiently_Active",
                            "Inactive")) %>%
    mutate(smoker =
               fct_collapse(smoker,
                    Current = c("Current_not_daily",
                                 "Current_daily"))) %>%
    mutate(dm_status =
               fct_collapse(dm_status,
                            No = c("No-Diabetes".
                                    "Pre-Diabetes".
                                    "Pregnancy-Induced"),
                            Yes = "Diabetes"))
```

## Sanity Checks

```
day2 %>% tabyl(activity) %>% adorn_pct_formatting()
```

 Still need to deal with the missing values, but now the order makes sense.

## **Sanity Checks**

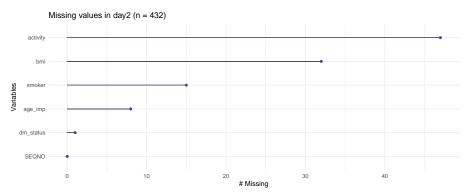
day2 %>% tabyl(dm\_status, smoker)

```
dm_status Current Former Never NA_
Yes 10 23 33 1
No 67 100 183 14
<NA> 0 1 0 0
```

- OK, now we have two dm\_status levels and three smoker levels, although we don't have a lot of currently smoking people with diabetes.
- Once we deal with the missing values, we should be all set.

# How many missing values in each variable?

```
gg_miss_var(day2) +
   labs(title = "Missing values in day2 (n = 432)")
```



• I used warning = FALSE in the code chunk setup.

### Table of missing values in each variable?

We can also just get a count of missing values for each variable with miss\_var\_summary(), which is also part of naniar.

```
miss_var_summary(day2)
```

```
# A tibble: 6 x 3
 variable n_miss pct_miss
 <chr> <int> <dbl>
              47 10.9
1 activity
2 bmi
            32 7.41
           15
                   3.47
3 smoker
                   1.85
4 age imp
                   0.231
 dm status
 SEQNO
                   0
```

# How many missing values per row (subject)?

- How many observations would we lose in a complete case analysis?
- Can we make the necessary assumption for a complete case analysis?

## What do we lose in a complete case analysis?

```
day2_cc <- day2 %>%
    filter(complete.cases(.))
dim(day2_cc)
```

[1] 362 6

This seems clean in some ways (and is the default approach in software), but actually it hides a very important assumption, that the data are **missing completely at random**.

```
prop_miss_case(day2); prop_miss_case(day2_cc)
```

```
[1] 0.162037
```

[1] 0

# Missing Data Mechanisms (Notes, Chapter 3)

- Missing Completely at Random (MCAR)
  - The probability of missing data is the same for every subject, so that throwing out cases with missing data does not bias inferences.
- Missing at Random (MAR)
  - Here, the probability that a variable is missing depends only on available information in your data (the other variables we have). If this is so, then imputation is the most appropriate option.
- Missing Not at Random (MNAR)
  - Whether data are missing is dependent on either unobserved predictors, or on the actual true (but unavailable) value of the observation itself or both. Even imputation cannot solve the problem.

What should we assume in our day2 scenario?

## Formulating a single imputation plan

miss\_var\_summary(day2) %>% kable()

variable	n_miss	pct_miss		
activity	47	10.8796296		
bmi	32	7.4074074		
smoker	15	3.4722222		
age_imp	8	1.8518519		
dm_status	1	0.2314815		
SEQNO	0	0.0000000		

Today, we'll use a *naive* approach to generating a single imputation.

- Impute dm\_status with a random draw from its distribution.
- Use CART to impute smoker and activity from dm\_status.
- Impute quantities via robust linear models using the factors.

# Single imputation in day2 to yield day2\_im

- impute\_rhd (random hot deck) for dm\_status
- impute\_cart (classification and regression trees) for other factors
- impute\_rlm (robust linear model) for age\_imp and bmi

```
prop_miss_case(day2_im)
```

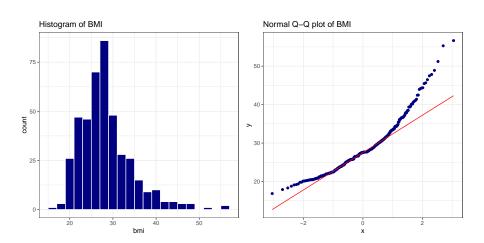
Γ17 0

#### Draw the outcome?

- We're interested in diabetes and smoking's association with BMI
  - What do the BMI data look like? (plots shown on next slide)

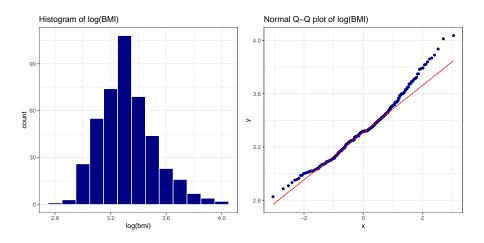
```
p1 <- ggplot(day2 im, aes(x = bmi)) +
    geom_histogram(fill = "navy", col = "white",
                   binwidth = 2) +
    labs(title = "Histogram of BMI")
p2 <- ggplot(day2_im, aes(sample = bmi)) +
    geom qq(col = "navy") + geom qq line(col = "red") +
    labs(title = "Normal Q-Q plot of BMI")
p1 + p2
```

## BMI data in day2\_im

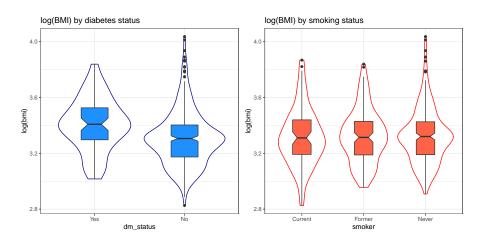


• These data are a little right-skewed. Transform?

## Consider a logarithmic transformation of BMI?

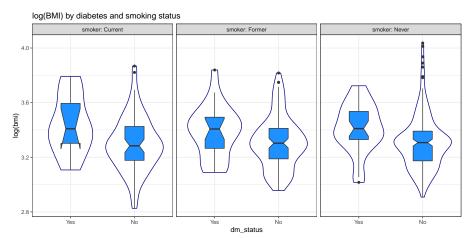


# Compare log(BMI) by diabetes and by smoking



# log(BMI) by diabetes and smoking together

notch went outside hinges. Try setting notch=FALSE.



## Finding the Means of Each Group

We'll plot the mean of log(bmi) in six combinations:

- two levels of dm\_status combined with
- three levels of smoker

`summarise()` has grouped output by 'dm\_status'. You can over

We can suppress this message with message = FALSE in the code chunk label.

## Here are the means of log(BMI) in each group

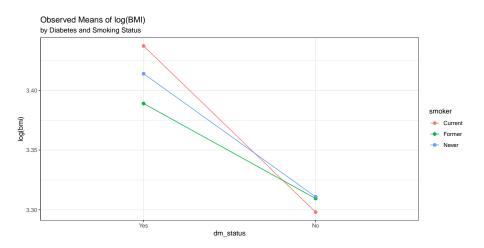
summaries\_1 %>% kable(digits = 2)

dm_status	smoker	n	mean	stdev
Yes	Current	10	3.44	0.22
Yes	Former	23	3.39	0.19
Yes	Never	34	3.41	0.18
No	Current	67	3.30	0.20
No	Former	101	3.31	0.19
No	Never	197	3.31	0.20

• Can we plot this information?

# Plotting the Means (code)

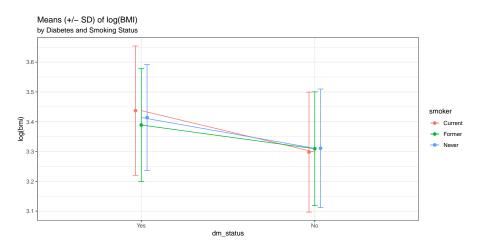
## Plotting the Means (results)



# Adding in standard deviations (code)

```
pd <- position dodge(0.1)
ggplot(summaries_1, aes(x = dm_status, y = mean,
                        col = smoker)) +
    geom_errorbar(aes(ymin = mean - stdev,
                      vmax = mean + stdev),
                      width = 0.1, position = pd) +
    geom_point(size = 2, position = pd) +
    geom_line(aes(group = smoker)) +
    labs(y = "log(bmi)",
         title = "Means (+/- SD) of log(BMI)",
         subtitle = "by Diabetes and Smoking Status")
```

## Adding in standard deviations (code)



## Review: One-Factor Analysis of Variance

```
m1 <- lm(log(bmi) ~ dm status, data = day2 im)
anova(m1)
Analysis of Variance Table
Response: log(bmi)
          Df Sum Sq Mean Sq F value Pr(>F)
dm status 1 0.5748 0.57482 15.113 0.0001172 ***
Residuals 430 16.3546 0.03803
Signif. codes:
0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

### Tidied m1 output

term	estimate	low90	high90	se	t	р
(Intercept)	3.41	3.37	3.45	0.02	143.07	0.00000
dm_statusNo	-0.10	-0.14	-0.06	0.03	-3.89	0.00012

# Revising the order?

term	estimate	low90	high90	se	t	р
(Intercept)	3.31	3.29	3.32	0.01	324.07	0.00000
dm_statusYes	0.10	0.06	0.14	0.03	3.89	0.00012

# **Glancing** at m1

```
glance(m1) %>%
    select(r.squared, adj.r.squared, sigma, AIC, BIC) %>%
    kable(digits = c(3, 3, 2, 1, 1))
```

r.squared	adj.r.squared	sigma	AIC	BIC
0.034	0.032	0.2	-182.4	-170.2

### **Developing a Two-Factor Model**

We want to describe the mean of log(BMI) as a function of both

- the two-level factor dm\_status, and
- the three-level factor smoker

One decision is whether we'll consider an **interaction** term between these two factors.

- A model with the interaction will fit the data a bit better, by some measures.
- A model with the interaction is most appropriate if we believe the dm\_status relationship with log(BMI) changes depending on the level of smoker.
  - or at least if we are unwilling to assume the smoker effect is the same regardless of dm\_status

# What is an interaction term (a product term)?

When we build our two-way model with interaction, we'll include a product term

```
m2 <- lm(log(bmi) ~ dm_status*smoker, data = day2_im)</pre>
```

as compared to a model without interaction, which we'd fit with:

Our main tool in thinking about these will be a means plot.

### Two-Way ANOVA with Interaction

```
m2 <- lm(log(bmi) ~ dm_status*smoker, data = day2_im)</pre>
anova(m2)
Analysis of Variance Table
Response: log(bmi)
                 Df Sum Sq Mean Sq F value Pr(>F)
                  1 0.5748 0.57482 14.9970 0.0001246 ***
dm status
         2 0.0051 0.00253 0.0659 0.9362544
smoker
dm_status:smoker 2 0.0214 0.01070 0.2791 0.7566000
Residuals 426 16.3282 0.03833
Signif. codes:
0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

#### Tidied m2 coefficients

term	estimate	low90	high90	se	р
(Intercept)	3.298	3.26	3.34	0.02	0.000
dm_statusYes	0.139	0.03	0.25	0.07	0.037
smokerFormer	0.011	-0.04	0.06	0.03	0.713
smokerNever	0.013	-0.03	0.06	0.03	0.644
dm_statusYes:smokerFormer	-0.060	-0.19	0.07	0.08	0.459
dm_statusYes:smokerNever	-0.036	-0.16	0.09	0.08	0.634

m2 estimates derived from the indicator (1/0) variables

• Estimated mean for a current smoker with no diabetes diagnosis?

- Estimated mean for a current smoker with no diabetes diagnosis?
- log(BMI) = 3.298, so estimated BMI = exp(3.298) = 27.06

- Estimated mean for a current smoker with no diabetes diagnosis?
- log(BMI) = 3.298, so estimated BMI = exp(3.298) = 27.06
- Estimated mean for a never smoker with no diabetes diagnosis?

- Estimated mean for a current smoker with no diabetes diagnosis?
- log(BMI) = 3.298, so estimated BMI = exp(3.298) = 27.06
- Estimated mean for a never smoker with no diabetes diagnosis?

m2 estimates derived from the indicator (1/0) variables

- Estimated mean for a current smoker with no diabetes diagnosis?
- log(BMI) = 3.298, so estimated BMI = exp(3.298) = 27.06

-0.036 (dm = Yes)(smoker = Never)

- Estimated mean for a never smoker with no diabetes diagnosis?
- Estimated mean for a never smoker with a diabetes diagnosis?

m2 estimates derived from the indicator (1/0) variables

- Estimated mean for a current smoker with no diabetes diagnosis?
- log(BMI) = 3.298, so estimated BMI = exp(3.298) = 27.06

-0.036 (dm = Yes)(smoker = Never)

- Estimated mean for a never smoker with no diabetes diagnosis?
- Estimated mean for a never smoker with a diabetes diagnosis?

#### What if we assume there's no interaction?

```
m2_no <- lm(log(bmi) ~ dm_status + smoker, data = day2_im)</pre>
anova(m2 no)
Analysis of Variance Table
Response: log(bmi)
           Df Sum Sq Mean Sq F value Pr(>F)
dm status 1 0.5748 0.57482 15.0477 0.0001213 ***
smoker 2 0.0051 0.00253 0.0661 0.9360460
Residuals 428 16.3496 0.03820
Signif. codes:
0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

### Tidied m2\_no coefficients

term	estimate	low90	high90	se	р
(Intercept)	3.303	3.27	3.34	0.02	0.000
dm_statusYes	0.101	0.06	0.14	0.03	0.000
smokerFormer	0.002	-0.04	0.05	0.03	0.932
smokerNever	0.008	-0.03	0.05	0.03	0.751

### Interpreting m2\_no (no interaction model)

```
log(BMI) = 3.303 + 0.101 (dm_status = Yes)
+ 0.002 (smoker = Former)
+ 0.008 (smoker = Never)
```

- Estimated mean for a current smoker with no diabetes diagnosis?
  - log(BMI) = 3.303, so estimated BMI = exp(3.303) = 27.19
- Estimated mean for a never smoker with no diabetes diagnosis?
  - log(BMI) = 3.303 + 0.008 = 3.311, so BMI = exp(3.311) = 27.41
- Estimated mean for a never smoker with a diabetes diagnosis?
  - log(BMI) = 3.303 + 0.008 + 0.101 = 3.412 so BMI = 30.33

#### What did we see in the data?

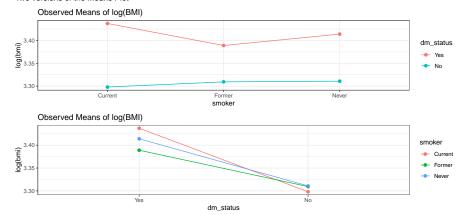
Estimates of mean (log(BMI)) from the two models, vs. the actual data.

dm_status	smoker	n	actual	m2 est.	m2_no est.
No	Current	67	3.298	3.298	3.303
No	Never	197	3.311	3.311	3.311
Yes	Never	34	3.414	3.414	3.412
No	Former	101	3.309	3.309	3.305
Yes	Current	10	3.437	3.437	3.404
Yes	Former	23	3.389	3.389	3.406

- The two-way ANOVA model with interaction simply reproduces the observed means.
- Not clear we want to assume the interaction effect is actually zero.

### Interaction Plot shows non-parallel lines?

Two versions of the Means Plot



# How about % of variation explained measures?

```
tidy(anova(m2)) %>% select(term, df, sumsq) %>%
   kable(dig = c(0, 0, 4))
```

term	df	sumsq
dm_status	1	0.5748
smoker	2	0.0051
dm_status:smoker	2	0.0214
Residuals	426	16.3282

- $R^2$  associated with the interaction term?
  - SS(interaction) is 0.0214
  - SS(m2) = 0.5748 + 0.0051 + 0.0214 = 0.6013
  - Interaction accounts for 3.6% of the variation explained by m2
  - Interaction accounts for 0.0214 / (0.6013+16.3282)=0.0013, or about 0.13% of the variation in log(BMI)

### Comparison of Fit across the models?

```
comp_table <- bind_rows( glance(m2), glance(m2_no) ) %>%
    mutate(mod = c("m2", "m2_no"))

comp_table %>%
    select(mod, r.squared, adj.r.squared, sigma, AIC, BIC) %>%
    kable(dig = c(0, 3, 3, 3, 1, 1))
```

mod	r.squared	adj.r.squared	sigma	AIC	BIC
m2	0.036	0.024	0.196	-175.1	-146.6
m2_no	0.034	0.027	0.195	-178.5	-158.2

• Is there much to choose from in comparing the in-sample performance?

#### How else can we assess the fit of these models?

We're not keen on making model decisions based on significance tests. Model selection doesn't actually work well, in practice.

```
anova(m2, m2_no)
```

Analysis of Variance Table

```
Model 1: log(bmi) ~ dm_status * smoker

Model 2: log(bmi) ~ dm_status + smoker

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

1 426 16.328

2 428 16.350 -2 -0.021396 0.2791 0.7566
```

• We'd rather think about how the two models reflect the data we have and predict on new data.

# OK, what if we add age as a covariate?

ANCOVA model m3 takes our model m2 (with interaction) and adds in age\_imp (centered by subtracting off its mean) as a predictor.

```
day2_im <- day2_im %>%
    mutate(age_c = age_imp - mean(age_imp))

day2_im %>% select(age_imp, age_c) %>% summary()
```

```
age_imp age_c

Min. :18.00 Min. :-38.942

1st Qu.:44.00 1st Qu.:-12.942

Median :59.00 Median : 2.058

Mean :56.94 Mean : 0.000

3rd Qu.:70.00 3rd Qu.: 13.058

Max. :96.00 Max. : 39.058
```

# OK, what if we add age as a covariate?

Here's an analysis of **covariance** model m3 with a factor-factor interaction, plus a centered quantitative covariate.

```
m3 <- lm(log(bmi) ~ dm_status * smoker + age_c,
data = day2_im)
```

Does this change the nature of the relationship we see between dm\_status, smoker and bmi?

#### Model m3 coefficients

term	estimate	low90	high90
(Intercept)	3.288	3.249	3.328
dm_statusYes	0.148	0.039	0.257
smokerFormer	0.026	-0.025	0.077
smokerNever	0.019	-0.026	0.064
age_c	-0.001	-0.002	-0.001
dm_statusYes:smokerFormer	-0.056	-0.188	0.075
dm_statusYes:smokerNever	-0.025	-0.149	0.099

#### ANOVA results for m2 and m3

tidy(anova(m3)) %>% select(term, df, sumsq) %>% kable(dig = 3)

term	df	sumsq
dm_status	1	0.575
smoker	2	0.005
age_c	1	0.307
dm_status:smoker	2	0.021
Residuals	425	16.021

tidy(anova(m2)) %>% select(term, df, sumsq) %>% kable(dig = 3)

term	df	sumsq
dm_status	1	0.575
smoker	2	0.005
dm_status:smoker	2	0.021
Residuals	426	16.328

# Does age\_imp improve quality of fit?

```
comp_table <- bind_rows( glance(m2), glance(m3) ) %>%
    mutate(mod = c("m2", "m3"))

comp_table %>%
    select(mod, r.squared, adj.r.squared, sigma, AIC, BIC) %>%
    kable(dig = c(0, 3, 3, 3, 1, 1))
```

mod	r.squared	adj.r.squared	sigma	AIC	BIC
m2	0.036	0.024	0.196	-175.1	-146.6
m3	0.054	0.040	0.194	-181.3	-148.7

# How about if we add a third factor (activity)?

term	df	sumsq
dm_status	1	0.575
smoker	2	0.005
age_c	1	0.307
activity	3	0.269
dm_status:smoker	2	0.036
Residuals	422	15.737

# Does activity improve quality of fit?

mod	r.squared	adj.r.squared	sigma	AIC	BIC
m1	0.034	0.032	0.195	-182.4	-170.2
m2	0.036	0.024	0.196	-175.1	-146.6
m3	0.054	0.040	0.194	-181.3	-148.7
m4	0.070	0.051	0.193	-183.0	-138.3

#### What's next?

- Is it feasible to look at the assumptions of these models?
- Could we consider additional interaction terms?
  - factor-factor interactions?
  - factor-covariate interactions?
- Interaction is just one type of non-linearity. Can we include other types?
- Should we think more about back-transformation in this setting?
- Could we split the sample and consider how well we'd predict in new data?
- Is 1m the best way to fit regression models to a quantitative outcome like log(bmi)?

Can we build up our framework for developing regression models?