Designing and Analyzing Statistical Models

EES 5891-03
Bayesian Statistical Methods
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Class #8: Tuesday, September 20 2022

More Categories of Confounding

General Principle: Identifiability

- Identifiable Models: Each set of model parameters makes different predictions
- Non-Identifiable Models: For any set of parameters, there are many other sets of parameters that make the same prediction
- Example: Categorical variables
 - x has three possible values: Architect, Baker, or Carpenter, and your regression will connect profession to income.
 - Represent x with two variables I_A and I_B , which are 1 if x has that value, and 0 otherwise. \[\begin{align} \text{Income} & \sim \text{Normal}(\mu, \sigma) \\ \mu &= \alpha + \beta_A I_A + \beta_B I_B \end{align} \]
 - Why don't we have I_c ?

Non-Identifiability

Now pick any number \(\delta\) and let \[\begin{align} \alpha' &= \alpha - \delta \\ \beta_A' &= \beta_A + \delta \\ \beta_B' &= \beta_B + \delta \\ \beta_C' &= \beta_C + \delta \end{align} \] And \[\mu' = \alpha' + \beta_A' I_A + \beta_B' I_B + \beta_C' I_C \]

Non-Identifiability (cont.)

```
\[\begin{align} \require{cancel} \mu &= \alpha + \beta_A I_A + \beta_B I_B + \beta_C I_C \\ &= (\alpha + \beta_C) + (\beta_A - \beta_C) I_A + (\beta_B - \beta_C) I_B \\ \mu' &= \alpha' + \beta_A' I_A + \beta_B' I_B + \beta_C' I_C \\ &= (\alpha' + \beta_C) + (\beta_A' - \beta_C') I_A + (\beta_B' - \beta_C') I_B \\ &= [(\alpha - \delta) + (\beta_C + \delta)] + [(\beta_A + \delta) - (\beta_C + \delta)] I_A + [(\beta_B + \delta) - (\beta_C + \delta)] I_B \\ &= [(\alpha - \cancel{\delta})) + (\beta_C + \cancel{\delta})] I_A + [(\beta_B + \cancel{\delta})] I_B \\ &= (\alpha + \beta_C) + (\beta_A - \beta_C) I_A + (\beta_B - \beta_C) I_B \\ &= \mu \end{align} \]
```

- So for any \(\delta\), \(\mu' = \mu\).
 - This means that there isn't a **best** set of values for \(\alpha\), \(\beta_A\), \(\beta_B\), and \(\beta_C\).
 - The problem is if you know I_A and I_B , then you also know I_C .
 - If you don't have an I_c variable, then this problem doesn't come up.
- There should be one fewer indicator variables than there are levels of the category variable.

Worked Example

- Pick values: \(\alpha = 1\), \(\beta_A = 2\), \(\beta_B = 3\), \(\beta_C = 4\)
- \(\delta = 0.5\)
- Alternate values: \(\alpha' = 0.5\), \(\beta_A = 2.5\), \(\beta_B = 3.5\), \(\beta_C = 4.5\) \[
 \begin{align} \require{cancel} \mu &= 1 + 2 I_A + 3 I_B + 4 I_C \\ &= (1 + 4) + (2 4) I_A + (3 4) I_B \\ &= 5 2 I_A 1 I_B \\ \mu' &= 0.5 + 2.5 I_A + 3.5 I_B + 4.5 I_C \\ &= (0.5 + 4.5) + (2.5 4.5) I_A + (3.5 4.5) I_B \\ &= 5 2 I_A 1 I_B \\ &= \mu \end{align} \]

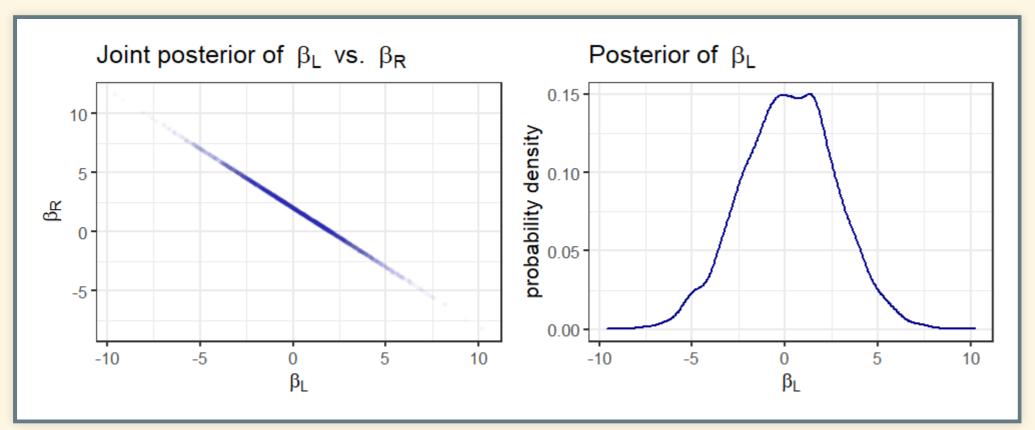
Multicollinearity

Multicollinearity

- Height versus length of legs: \[\begin{align} H &\sim \text{Normal}(\mu, \sigma) \\ \mu &= \alpha + \beta_R R + \beta_L L, \end{align} \] where
 - H is the person's height,
 - R is the length of the right leg,
 - *L* is the length of the left leg.
- The legs don't have identical length, but they are highly correlated.
- This creates a problem of identifiability:
 - Start with \(\beta_L\) and \(\beta_R\),
 - then for some number \(\\delta\\), consider
 - \(\beta_L' = \beta_L + \delta\)
 - \(\beta_R' = \beta_R \delta\)
 - On average (L = R), so $(\mu) = \mu$.
 - \(\beta_L\) and \(\beta_R\) are not identifiable.

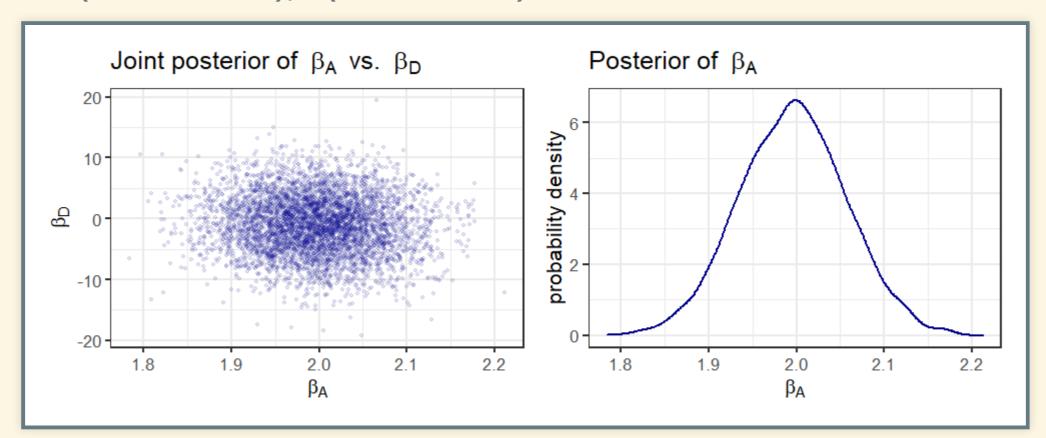
Does Multicollinearity Matter?

- McElreath says it doesn't matter for model predictions
 - Only matters for interpreting model.
 - Large uncertainty in posteriors for parameters when considered,
 - Because many values of \(\beta_L\) and \(\beta_R\) are just as probable.
 - The *joint posterior* for \(\beta_L\) and \(\beta_R\) is very narrow.



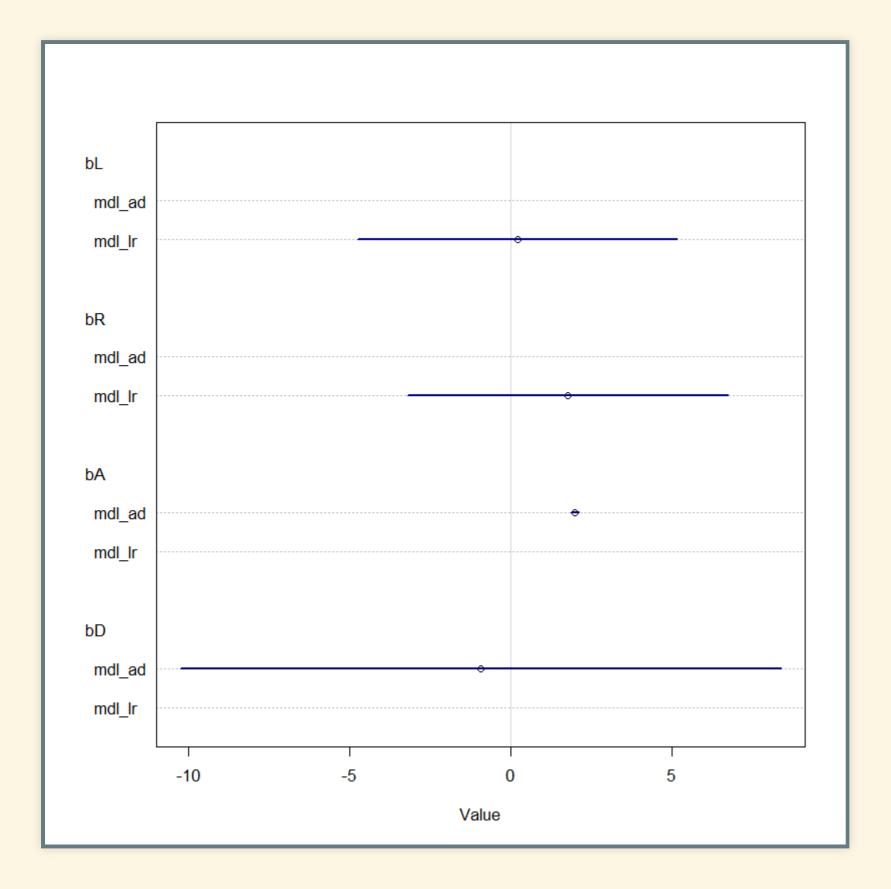
Another perspective

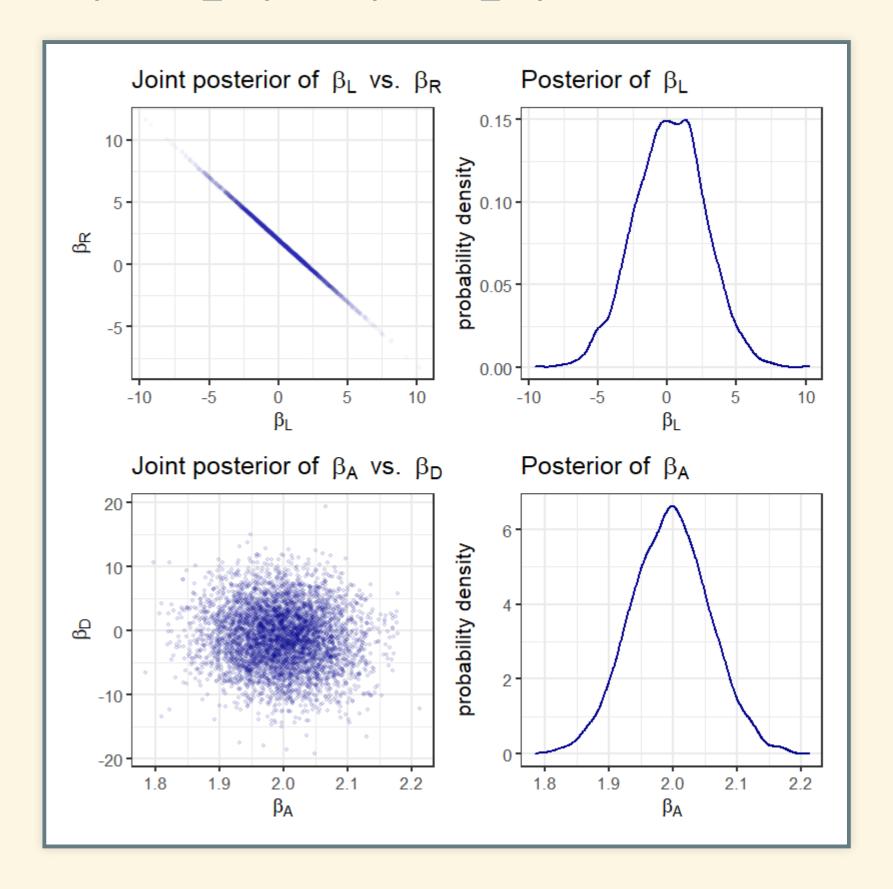
- Multicollinearity can make computational analysis difficult
- One response:
 - Define new variables:
 - $\circ \A = \text{L} + R) / 2$
 - $\circ \(D = \text{difference} = (L R) / 2)$



Summary

• Note how the different scales for \(\beta_A\) vs. \(\beta_L\).





Multicollinearity with Milk Data

Multicollinearity with Milk Data

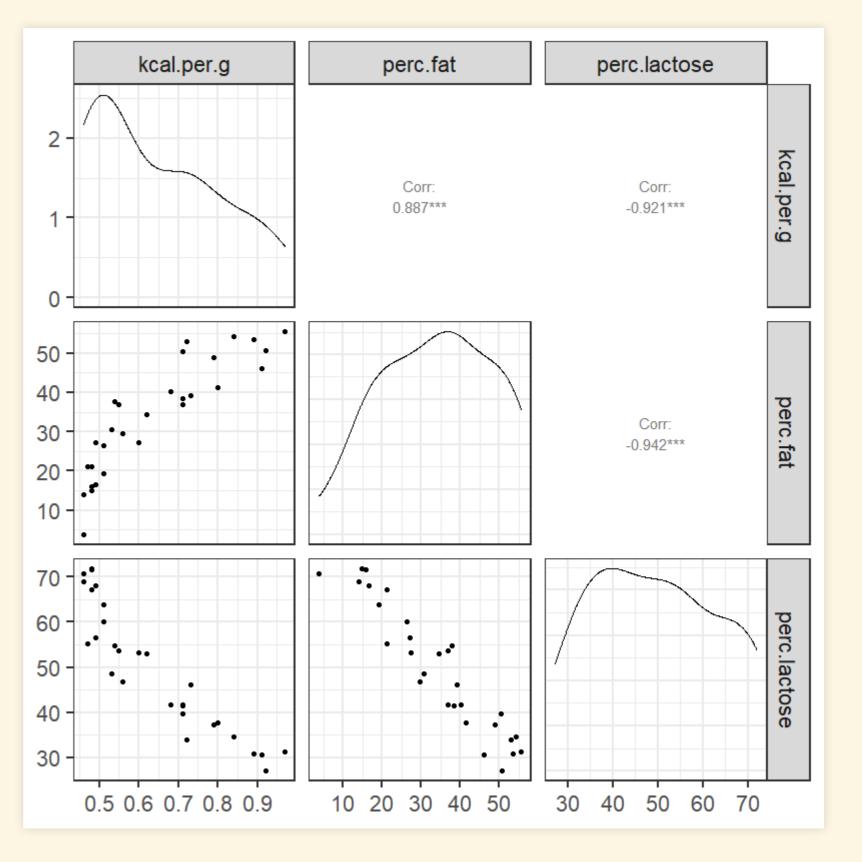
• Calories come from fat and sugar (lactose):

```
data(milk)
d <- milk
d$K <- standardize( d$kcal.per.g )
d$F <- standardize( d$perc.fat )
d$L <- standardize( d$perc.lactose )</pre>
```

• Make a pairwise correlation plot

```
library(tidyverse)
library(GGally)

d %>% select(kcal.per.g, perc.fat, perc.lactose) %>%
    ggpairs()
```



Making a model

```
mdl_milk <- quap(
    alist(
        K ~ dnorm(mu, sigma),
        mu <- a + bF * F + bL * L,
        a ~ dnorm(0, 0.2),
        bF ~ dnorm(0, 0.5),
        bL ~ dnorm(0, 0.5),
        sigma ~ dexp(1)
    ), data=d )

precis_show(precis(mdl_milk, digits = 2))</pre>
```

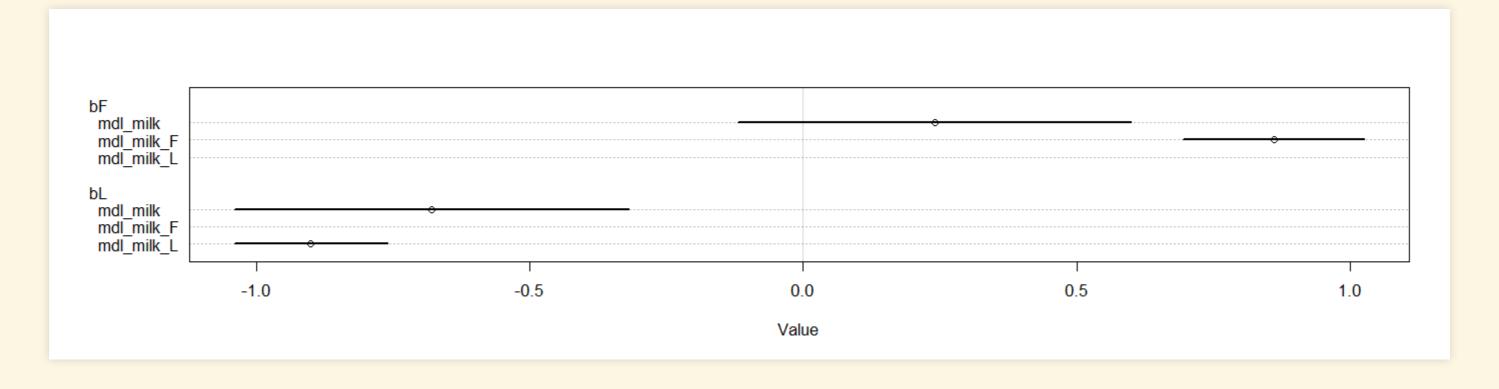
```
## mean sd 5.5% 94.5%

## a 0.00 0.07 -0.11 0.11

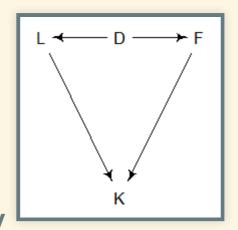
## bF 0.24 0.18 -0.05 0.54

## bL -0.68 0.18 -0.97 -0.38

## sigma 0.38 0.05 0.30 0.46
```



Explaining the multicollinearity



- Knowledge of biology
- Density D is important
 - Frequent nursing: watery, low-energy milk, high in sugar (lactose)
 - Infrequent nursing: rich, dense, high-energy milk, high in fat

Post-Treatment Bias

Anti-Fungal Treatment Experiment

- You do an experiment
 - Divide plants in 2 groups
 - Apply anti-fungal treatment to one group (\(T = 1\))
 - \circ The other is a control (\(T = 0\))
 - Observe whether there is fungus after treatment (\(F\))
 - Compare height before treatment (\
 (H_0\)) to height some time after treatment (\(H_1\)).
 - Growth rate \(p \ge 0\) unless fungus is very bad.

```
mdl_fungus <- quap(
    alist(
        H1 ~ dnorm(mu, sigma),
        mu <- H0 * p,
        # p is growth rate
        p <- a + bT * T + bF * F,
        a ~ dlnorm(0, 0.2),
        bT ~ dnorm(0, 0.5),
        bF ~ dnorm(0, 0.5),
        sigma ~ dexp(1)
    ), data=d)
precis_show(precis(mdl_fungus, digits = 2))</pre>
```

```
## mean sd 5.5% 94.5%

## a 1.48 0.02 1.44 1.52

## bT 0.00 0.03 -0.05 0.05

## bF -0.27 0.04 -0.33 -0.21

## sigma 1.41 0.10 1.25 1.57
```

- Why doesn't the treatment have an effect?
 - \blacksquare mean(bT) = 0.

Understanding the problem

- Fungus is the big thing that affects the plants' growth
- Treatment affects fungus.
 - Doesn't affect plants directly
 - Doesn't always eliminate all fungus
- Fungus is a better predictor
 - But we don't know how bad fungus will be until after we treat.
- DAG

```
H_0 \longrightarrow H_1 \longleftarrow F \longleftarrow T
```

```
## Implied Conditional Independencies

## F _ | | _ H _ 0

## H _ 0 _ | | _ T

## H _ 1 _ | | _ T | F
```

```
mdl fungus <- quap (
    alist(
         H1 ~ dnorm(mu, sigma),
         mu \leftarrow H0 * p
         # p is growth rate
         p <- a + bT * T + bF *
         F,
         a \sim dlnorm(0, 0.2),
         bT \sim dnorm(0, 0.5),
         bF \sim dnorm(0, 0.5),
         sigma \sim dexp(1)
    ), data=d)
```

A Better Model

- DAG
- Conditioning on *F* induces a *D-separation* (*directional* separation) between *T* and *H1*.
- Remove fungus data from the model.

```
mdl_fungus_2 <- quap(
    alist(
        h1 ~ dnorm( mu , sigma ),
        mu <- h0 * p,
        p <- a + bt*treatment,
        a ~ dlnorm( 0 , 0.2 ),
        bt ~ dnorm( 0 , 0.5 ),
        sigma ~ dexp( 1 )
    ), data=d )
precis_show(precis(mdl_fungus_2, digits = 2))</pre>
```

```
## mean sd 5.5% 94.5%

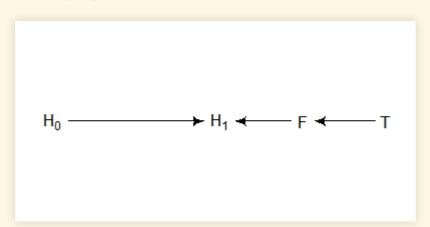
## a 1.38 0.03 1.34 1.42

## bt 0.08 0.03 0.03 0.14

## sigma 1.75 0.12 1.55 1.94
```

Other Post-Treatment Bias Problems

Suppose we have this DAG:

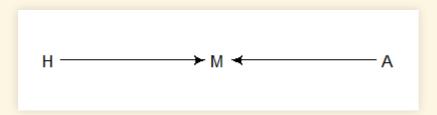


- Fungus does not influence plant growth.
- Moisture influences both plant growth and fungus
- Fitting our original model falsely implies that treatment benefits plants.
- This is a kind of *collider* effect.

Collider Bias

Happiness and Age

- Do people get happier as they get older?
- Suppose:
 - Everyone's happiness is something they are born with and it doesn't change.
 - Happier people are more likely to get married
 - Older people are more likely to be married.
 - DAG:



This diagram is a **collider**: Causal paths from *H* and *A collide* at *M*

Analyze Happiness Data

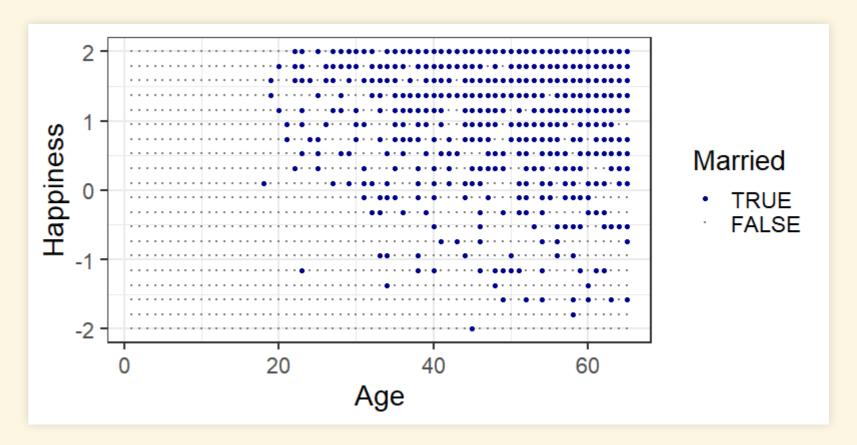
Load data

```
d <- sim_happiness( seed=1977 , N_years=1000 )</pre>
```

- Look for an association between age and happiness.
 - We suspect that the relationship between age and happiness may be different for married people, so we include marriage as a variable.
- Clean the data: Select adults and convert age to a variable that goes from 0 to 1, and create a marriage index:

```
d2 <- d[ d$age>17 , ] # only adults
d2$A <- ( d2$age - 18 ) / ( 65 - 18 )
d2$mid <- d2$married + 1</pre>
```

 The model says that people become unhappy as they get older



```
mdl_happy <- quap(
   alist(
     happiness ~ dnorm(mu, sigma),
     mu <- a[mid] + bA * A,
     a[mid] ~ dnorm(0, 1),
     bA ~ dnorm(0, 2),
     sigma ~ dexp(1)
   ), data=d2)
precis_show(precis(mdl_happy, digits = 2))</pre>
```

```
## mean sd 5.5% 94.5%

## bA -0.75 0.11 -0.93 -0.57

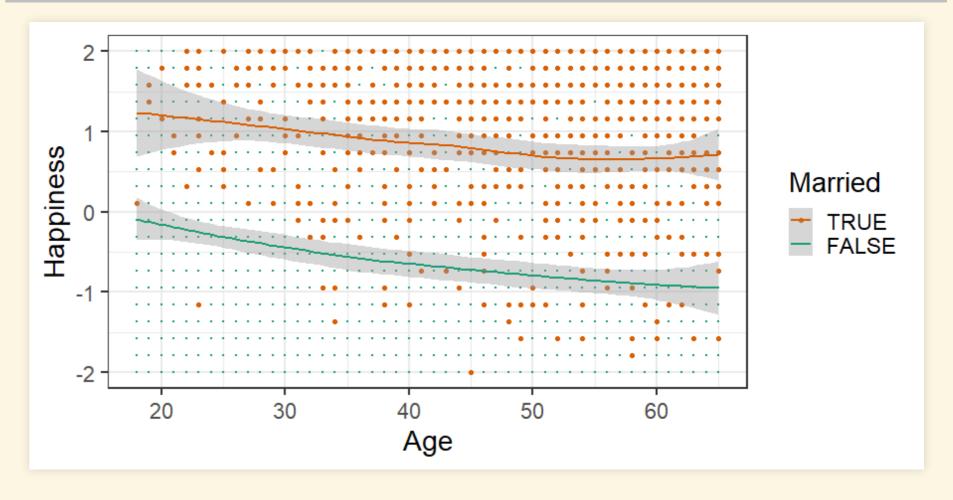
## sigma 0.99 0.02 0.95 1.03
```

A different Model

- Try a different model that does not control for mdl_happy_2 <- quap(marriage.
- This model shows no association between age and happiness.
- What happened?
- Consider married people:
 - Older people are more likely to get married
 - Happier people are more likely to get married
 - Happy people get married younger
 - Unhappy people get married older
 - Thus, among married people, younger people are happier, and older ones are unhappier.
- Consider single people
 - As people age, happier ones marry,
 - So the older someone is, if they are still single, they're more likely to be unhappy.

```
mdl_happy_2 <- quap(
    alist(
        happiness ~ dnorm( mu , sigma ),
        mu <- a + bA*A,
        a ~ dnorm( 0 , 1 ),
        bA ~ dnorm( 0 , 2 ),
        sigma ~ dexp(1)
    ) , data=d2 )
precis_show(precis(mdl_happy_2, digits = 2))</pre>
```

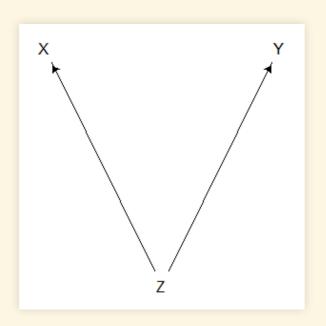
```
## a 0.00 0.08 -0.12 0.12
## bA 0.00 0.13 -0.21 0.21
## sigma 1.21 0.03 1.17 1.26
```



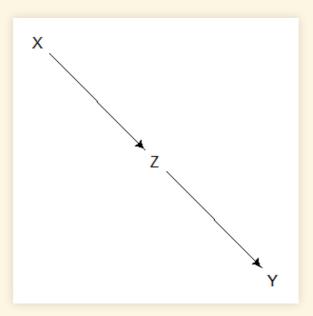
Categories of Confounding Relationships

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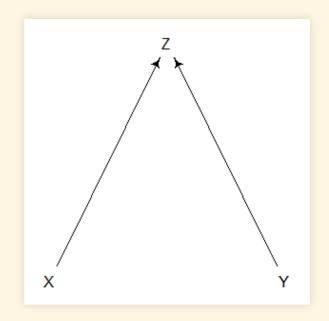
- There are four major categories of confounding relationships:
- 1. Fork



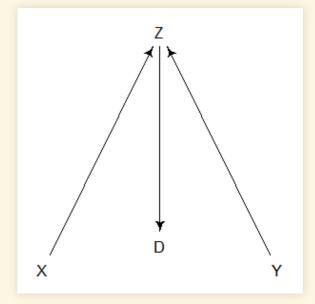
2. Pipe



3. Collider



4. Descendant



• All causal DAGs are build of combinations of these four patterns.