

LMTLE: A Brief Introduction

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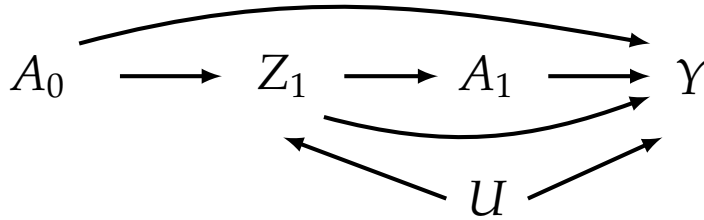
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1 Introduction to Complex Longitudinal Data

Complex longitudinal data is becoming more common in a number of sector. These data are different from more traditional “longitudinal data” that one encounters in classical statistics courses.¹ Complex longitudinal data requires the presence of two features: first, repeated exposure, confounder, and (potentially) outcome measures; second, there has to be time-dependent feedback between these exposure, confounder, and (potentially) outcome variables.

This Figure demonstrates these basic conditions:



¹ For example, courses where you would learn how to estimate models with more sophisticated correlation structures, such as generalized estimating equations or mixed effects models

Figure 1: Causal diagram representing the structure from which the simple simulated data were generated.

This Figure is a simplified version, and several details can be added. For example, baseline covariates will always be present; there may be a Z_0 variable measured; there may be more than one time-dependent confounder; and the outcome may be measured at multiple time points, and may also serve as a time-dependent confounder.

Generally, the presence of a causal relation between Z_1 and A_1 suggests that Z_1 is a confounding variable. However, we cannot simply adjust for Z_1 in a standard regression model, since this would (i) block part of the effect of interest from $A_0 \rightarrow Z_1 \rightarrow Y$, and (ii) induce collider stratification bias through $A_0 \rightarrow Z_1 \leftarrow U \rightarrow Y$.

Because of this, we have to use specialized methods to estimate average treatment effects with complex longitudinal data.

2 IPW and g Computation for CLD

Perhaps the two most common techniques are inverse probability weighting or g computation (aka the parametric g formula) in such settings.

For example, if we're interested in the following ATE:

$$\psi = E(Y^{\bar{a}=1} - Y^{\bar{a}=0})$$

where \bar{a} denotes the entire history of the exposure measurement from the start to the end of follow-up for each person. To make our illustrations concrete, let's use a simple simulated dataset with longitudinal information on an exposure, several time-dependent confounders, and a time-to-event outcome.² Here's what the data look like:

² These data were simulated from Jessica Young's algorithm, modified by Erica Moodie. Details can be found in [Young et al. \(2010\)](#) and [Moodie et al. \(2014\)](#)

```
a <- read_csv(here("data", "2023_04_21-time-dependent.csv")) %>%
  group_by(ID) %>%
  mutate(exposure_lag = lag(exposure, n = 1L,
    default = 0), c1_lag = lag(c1, n = 1L,
    default = 0), c2_lag = lag(c2, n = 1L,
    default = 0)) %>%
  ungroup()

a
```

```
## # A tibble: 3,435 x 9
##       ID   Int exposure    c1    c2    Y
##   <dbl> <dbl>   <dbl> <dbl> <dbl> <dbl>
## 1     1     1     0     0     1     0
## 2     1     2     1     1     1     0
## 3     1     3     1     1     0     0
## 4     1     4     1     1     1     1
## 5     2     1     0     0     1     0
## 6     2     2     1     0     1     0
## 7     2     3     0     1     1     0
## 8     2     4     1     0     1     0
## 9     3     1     1     1     1     0
```

```
## 10      3      2      1      1      1      0
## # i 3,425 more rows
## # i 3 more variables: exposure_lag <dbl>,
## #   c1_lag <dbl>, c2_lag <dbl>
```

```
## look at proportion of outcome at
## each time point
a %>%
  group_by(Int) %>%
  summarise(mY = mean(Y))
```

```
## # A tibble: 4 x 2
##   Int    mY
##   <dbl> <dbl>
## 1     1 0.220
## 2     2 0.254
## 3     3 0.253
## 4     4 0.227
```

Let's talk a little about this data structure.

2.1 Inverse Probability Weighting

Let's start by constructing stabilized IP weights to estimate the ATE in these data:

```
# numerator
num <- glm(exposure ~ factor(Int), data = a,
  family = binomial("logit"))$fitted.values

# denominator
den <- glm(exposure ~ factor(Int) + exposure_lag +
  c1 + c2 + c1_lag + c2_lag, data = a,
  family = binomial("logit"))$fitted.values

# a_last <- a %>% mutate(last_id =
```

```

# !duplicated(ID, fromLast = T)) %>%
# filter(last_id==1) %>% select(ID,
# Int) %>% rename(last_Int = Int)
# a_last a <- left_join(a, a_last, by =
# 'ID') , cum_exp =
# cumsum(exposure)/last_Int a %>%
# select(ID, Int, exposure, Y,
# last_Int)

a <- a %>%
  mutate(sw_ = (num/den) * exposure + ((1 -
    num)/(1 - den)) * (1 - exposure)) %>%
  group_by(ID) %>%
  mutate(sw = cumprod(sw_)) %>%
  ungroup() %>%
  select(-sw_)

a %>%
  group_by(Int) %>%
  summarise(meanSW = mean(sw), maxSW = max(sw))

```

```

## # A tibble: 4 x 3
##   Int meanSW maxSW
##   <dbl> <dbl> <dbl>
## 1     1  0.997  1.60
## 2     2  0.994  2.65
## 3     3  0.988  3.87
## 4     4  0.993  5.79

```

We can then use these weights to fit an IP weighted MSM:

```

library(lmtest)
library(sandwich)

modMSM <- lm(Y ~ factor(Int) + I(exposure/4),

```

```

data = a, weights = sw)

coeftest(modMSM, vcov. = vcovCL(modMSM, cluster = a$ID,
  type = "HC3"))

##
## t test of coefficients:
##
##              Estimate Std. Error t value
## (Intercept)   0.1920508  0.0160912 11.9351
## factor(Int)2   0.0198237  0.0192788  1.0283
## factor(Int)3   0.0303484  0.0222573  1.3635
## factor(Int)4  -0.0092636  0.0241587 -0.3834
## I(exposure/4)  0.1840595  0.0721127  2.5524
##              Pr(>|t|)
## (Intercept)    < 2e-16 ***
## factor(Int)2    0.30390
## factor(Int)3    0.17281
## factor(Int)4    0.70141
## I(exposure/4)  0.01074 *
## ---
## Signif. codes:
## 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

install.packages("ltml", repos = "http://lib.stat.cmu.edu/R/CRAN",
  dependencies = T)

##
## The downloaded binary packages are in
## /var/folders/zm/rqfq5xs0fs86qs2mcxk6q0r0000gr/T/Rtmp22ZxER/downloaded_packages

library(ltml)

# read in data again, to simplify
a <- read_csv(here("data", "2023_04_21-time-dependent.csv"))

```

```
## Rows: 3435 Columns: 6
## -- Column specification -----
## Delimiter: ","
## dbl (6): ID, Int, exposure, c1, c2, Y
##
## i Use `spec()` to retrieve the full column specification for this data.
## i Specify the column types or set `show_col_types = FALSE` to quiet this message.
```

```
# convert data from long to wide
a %>%
  print(n = 3)
```

```
## # A tibble: 3,435 x 6
##       ID   Int exposure    c1    c2    Y
##   <dbl> <dbl>   <dbl> <dbl> <dbl> <dbl>
## 1     1     1       0     0     1     0
## 2     1     2       1     1     1     0
## 3     1     3       1     1     0     0
## # i 3,432 more rows
```

```
# TO DEAL WITH BASELINE CONFOUNDERS,
# KEEP THEM IN DATA
b <- a %>%
  pivot_wider(names_from = Int, values_from = c(exposure,
    c1, c2, Y)) %>%
  mutate(Y_1 = if_else(is.na(Y_1), 1, Y_1),
    Y_2 = if_else(is.na(Y_2), 1, Y_2),
    Y_3 = if_else(is.na(Y_3), 1, Y_3),
    Y_4 = if_else(is.na(Y_4), 1, Y_4)) %>%
  select(exposure_1, c1_1, c2_1, Y_1, exposure_2,
    c1_2, c2_2, Y_2, exposure_3, c1_3,
    c2_3, Y_3, exposure_4, c1_4, c2_4,
    Y_4)
```

b

```
## # A tibble: 1,228 x 16
##   exposure_1 c1_1 c2_1 Y_1 exposure_2
##         <dbl> <dbl> <dbl> <dbl>      <dbl>
## 1           0     0     1     0          1
## 2           0     0     1     0          1
## 3           1     1     1     0          1
## 4           0     1     1     0          0
## 5           1     1     1     0          1
## 6           1     1     1     0          1
## 7           1     0     1     0          0
## 8           1     1     0     0          1
## 9           0     0     0     0          1
## 10          1     1     1     1          NA
## # i 1,218 more rows
## # i 11 more variables: c1_2 <dbl>, c2_2 <dbl>,
## #   Y_2 <dbl>, exposure_3 <dbl>, c1_3 <dbl>,
## #   c2_3 <dbl>, Y_3 <dbl>, exposure_4 <dbl>,
## #   c1_4 <dbl>, c2_4 <dbl>, Y_4 <dbl>
```

```
# ltmle
```

```
# super learner library
```

```
sl.lib <- c("SL.mean",
           "SL.glm",
           "SL.ranger")
```

```
#ltmle
```

```
result <- ltmle(b,
               Anodes=c(paste0("exposure_",1:4)),
               Lnodes=c("c1_1", "c2_1", "c1_2", "c2_2",
                       "c1_3", "c2_3", "c1_4", "c2_4"),
               Ynodes=c("Y_1","Y_2","Y_3","Y_4"),
```



```

survivalOutcome = TRUE,
SL.library = list(Q = sl.lib, g = sl.lib),
abar=list(treatment = c(1, 1, 1, 1),
          control = c(0, 0, 0, 0)),
# estimate.time = T, need to comment out to run
stratify = T)

```

```
## Qform not specified, using defaults:
```

```
## formula for c1_1:
```

```
## Q.kplus1 ~ 1
```

```
## formula for c1_2:
```

```
## Q.kplus1 ~ c1_1 + c2_1
```

```
## formula for c1_3:
```

```
## Q.kplus1 ~ c1_1 + c2_1 + c1_2 + c2_2
```

```
## formula for c1_4:
```

```
## Q.kplus1 ~ c1_1 + c2_1 + c1_2 + c2_2 + c1_3 + c2_3
```

```
##
```

```
## gform not specified, using defaults:
```

```
## formula for exposure_1:
```

```
## exposure_1 ~ 1
```

```
## formula for exposure_2:
```

```
## exposure_2 ~ c1_1 + c2_1
```

```
## formula for exposure_3:
```

```
## exposure_3 ~ c1_1 + c2_1 + c1_2 + c2_2
```

```
## formula for exposure_4:
```

```

## exposure_4 ~ c1_1 + c2_1 + c1_2 + c2_2 + c1_3 + c2_3

##

## Timing estimate unavailable

## Warning in predict.lm(object, newdata, se.fit,
## scale = 1, type = if (type == : prediction
## from rank-deficient fit; attr(*, "non-estim")
## has doubtful cases

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## Warning: glm.fit: fitted probabilities
## numerically 0 or 1 occurred

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## has doubtful cases

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## numerically 0 or 1 occurred

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## scale = 1, type = if (type == : prediction
## from rank-deficient fit; attr(*, "non-estim")
## has doubtful cases

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## scale = 1, type = if (type == : prediction
## from rank-deficient fit; attr(*, "non-estim")
## has doubtful cases

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## scale = 1, type = if (type == : prediction
## from rank-deficient fit; attr(*, "non-estim")
## has doubtful cases

```

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## has doubtful cases
```

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## has doubtful cases
```

```
## Warning in predict.lm(object, newdata, se.fit,
## scale = 1, type = if (type == : prediction
## from rank-deficient fit; attr(*, "non-estim")
## has doubtful cases
```

```
summary(result)
```

```
## Estimator:  tmle
## Call:
## ltmle(data = b, Anodes = c(paste0("exposure_", 1:4)), Lnodes = c("c1_1",
##      "c2_1", "c1_2", "c2_2", "c1_3", "c2_3", "c1_4", "c2_4"),
##      Ynodes = c("Y_1", "Y_2", "Y_3", "Y_4"), survivalOutcome = TRUE,
##      abar = list(treatment = c(1, 1, 1, 1), control = c(0, 0, 0,
##      0)), stratify = T, SL.library = list(Q = sl.lib, g = sl.lib))
##
## Treatment Estimate:
##      Parameter Estimate:  0.68612
##      Estimated Std Err:  0.019473
##      p-value:  <2e-16
##      95% Conf Interval: (0.64795, 0.72429)
##
## Control Estimate:
##      Parameter Estimate:  0.46255
##      Estimated Std Err:  0.052735
```

```
##                p-value: <2e-16
##      95% Conf Interval: (0.35919, 0.5659)
##
## Additive Treatment Effect:
##      Parameter Estimate: 0.22357
##      Estimated Std Err: 0.056215
##                p-value: 6.977e-05
##      95% Conf Interval: (0.11339, 0.33375)
##
## Relative Risk:
##      Parameter Estimate: 1.4833
##      Est Std Err log(RR): 0.11749
##                p-value: 0.00079055
##      95% Conf Interval: (1.1782, 1.8675)
##
## Odds Ratio:
##      Parameter Estimate: 2.5399
##      Est Std Err log(OR): 0.2306
##                p-value: 5.2941e-05
##      95% Conf Interval: (1.6163, 3.9912)
```

```
result$fit$g
```

```
## [[1]]
## [[1]]$exposure_1
##              Estimate Std. Error t value
## (Intercept) 0.4061438 0.05827759 6.969124
##              Pr(>|t|)
## (Intercept) 5.19302e-12
##
## [[1]]$exposure_2
##              Risk      Coef
## SL.mean_All  0.09853545 0.0000000
## SL.glm_All   0.09763598 0.5063361
## SL.ranger_All 0.09764399 0.4936639
```

```
##
## [[1]]$exposure_3
##           Risk      Coef
## SL.mean_All  0.1341856 0.7965582
## SL.glm_All   0.1361927 0.2034418
## SL.ranger_All 0.1365624 0.0000000
##
## [[1]]$exposure_4
##           Risk      Coef
## SL.mean_All  0.1165104 0.2171115
## SL.glm_All   0.1172869 0.0000000
## SL.ranger_All 0.1143180 0.7828885
##
##
## [[2]]
## [[2]]$exposure_1
##           Estimate Std. Error  t value
## (Intercept) 0.4061438 0.05827759 6.969124
##           Pr(>|t|)
## (Intercept) 5.19302e-12
##
## [[2]]$exposure_2
##           Risk      Coef
## SL.mean_All  0.2351189 0.1902279
## SL.glm_All   0.2306222 0.8097721
## SL.ranger_All 0.2316999 0.0000000
##
## [[2]]$exposure_3
##           Risk      Coef
## SL.mean_All  0.2440609 0.56230791
## SL.glm_All   0.2491922 0.04838326
## SL.ranger_All 0.2467136 0.38930883
##
## [[2]]$exposure_4
##           Risk      Coef
```

```
## SL.mean_All    0.2564660 0.8639936
## SL.glm_All     0.3004954 0.1360064
## SL.ranger_All  0.2836678 0.0000000
```

```
result$fit$Q
```

```
## [[1]]
## [[1]]$c1_1
##           Estimate Std. Error  t value
## (Intercept) 0.7820347 0.03031254 25.79905
##           Pr(>|t|)
## (Intercept) 4.828552e-105
##
## [[1]]$c1_2
##           Risk      Coef
## SL.mean_All    0.06823469 0.1490461
## SL.glm_All     0.06749214 0.8509539
## SL.ranger_All  0.16189686 0.0000000
##
## [[1]]$c1_3
##           Risk      Coef
## SL.mean_All    0.1112692 0.3686925
## SL.glm_All     0.1103943 0.6313075
## SL.ranger_All  0.1431575 0.0000000
##
## [[1]]$c1_4
##           Risk Coef
## SL.mean_All    0.1819373    1
## SL.glm_All     0.1900696    0
## SL.ranger_All  0.1903400    0
##
##
## [[2]]
## [[2]]$c1_1
##           Estimate Std. Error  t value
## (Intercept) -0.1500947 0.04783664 -3.137651
```



```
##                Pr(>|t|)
## (Intercept) 0.001805626
##
## [[2]]$c1_2
##                Risk Coef
## SL.mean_All  0.09688196    1
## SL.glm_All   0.10009314    0
## SL.ranger_All 0.12193717    0
##
## [[2]]$c1_3
##                Risk Coef
## SL.mean_All  0.06207995    1
## SL.glm_All   0.06902620    0
## SL.ranger_All 0.07987094    0
##
## [[2]]$c1_4
##                Risk Coef
## SL.mean_All  0.09408145    1
## SL.glm_All   0.15911422    0
## SL.ranger_All 0.11498982    0
```

References

- Erica E. M. Moodie, David A. Stephens, and Marina B. Klein. A marginal structural model for multiple-outcome survival data: assessing the impact of injection drug use on several causes of death in the canadian co-infection cohort. *Stat Med*, 33(8):1409–1425, 2014.
- J. G. Young, M. A. Hernán, S. Picciotto, and J. M. Robins. Relation between three classes of structural models for the effect of a time-varying exposure on survival. *Lifetime Data Anal*, 16(1):71–84, 2010.