BIOST 515-Project

Michael Cork, Claire Rothschild, Baohai WattShao, Aji John Winter Quarter 2018

We are interested in changes in plasma lipid biomarkers for coronary heart disease (CHD) after hormone replacementtherapy in a sample of 2,763 womenfrom the Heart and Estrogen/progestin Replacement Study (HERS). HERS was randomized, double-blind, placebo-controlled trial designed to test the efficacy and safety of estrogen plus progestin therapy for prevention of recurrent coronary heart disease (CHD) events in women.

Questions of interest to be answered

The data tobe analyzed for this projectis a subset of the data collected from the 2,763 women in the Heart and Estrogen/progestin Replacement Study (HERS) clinical trial of hormone therapy. The questions to be addressed are:

1.

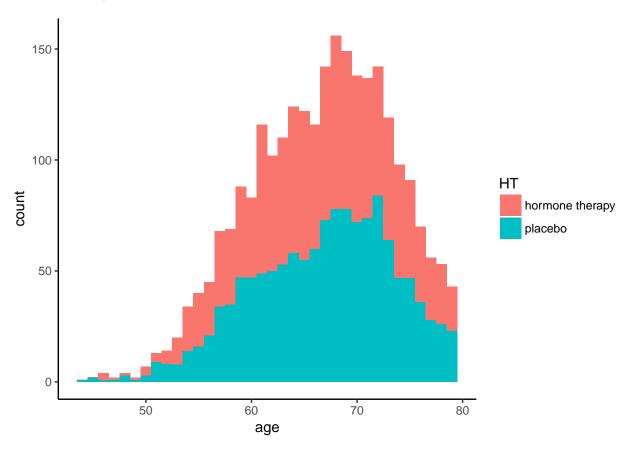
What associations exist between the plasma lipid biomarkers for CHD at baseline (i.e., prior to randomized treatment assignment) and the available data on participant demographics (age, race, BMI), behavior (smoking, alcohol consumption, physical activity), and available clinical and laboratory measures of organ system functioning (e.g., glucose, blood pressure)?

univariate

2.

Is there any evidence of hormone therapy treatment effects on the plasma lipid biomarkers after one year of treatment?

Profile of subjects studied



BMI Profile of subjects studied by Control/Treatment

НТ	n	${\rm meanBMI}$	sdBMI	meanBMI.1	sdBMI.1
hormone therapy	1274	28.59735	5.434025		5.509812
placebo	1306	28.52488	5.495672		5.605677

HT	n	meanLDL	sdLDL	${\rm meanLDL.1}$	sdLDL.1
hormone therapy placebo	1274 1306		38.26348 37.39339		36.95654 39.67142

3.

Is there any attenuation of the hormone therapy treatment effects after adjustment for known risk factors for CHD?

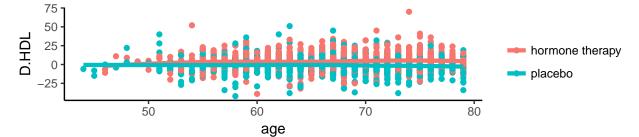
4.

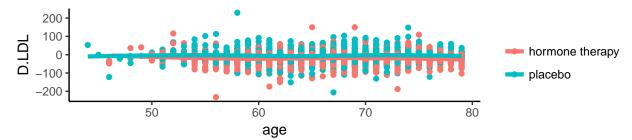
Do any of the hormone therapy treatment effects identified above for the plasma lipid biomarkers differ according to race/ethnicity, statin medication use, smoking behavior, or alcohol consumption.

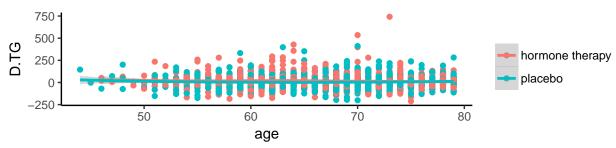
Show the distribution by biomarkers between Placebo and Treatment

Warning: Removed 5 rows containing missing values (geom_smooth).

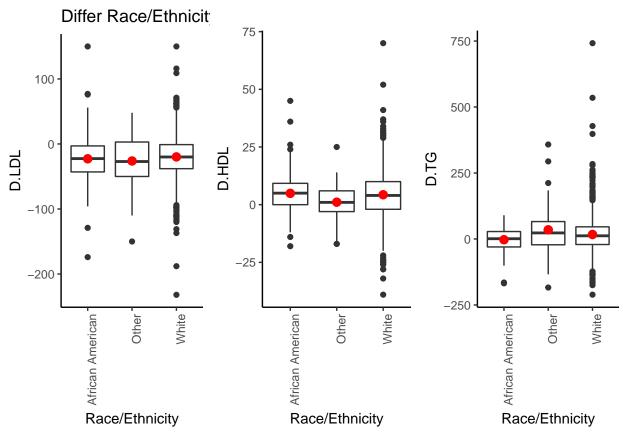
Warning: Removed 5 rows containing missing values (geom_smooth).







Checking by Race

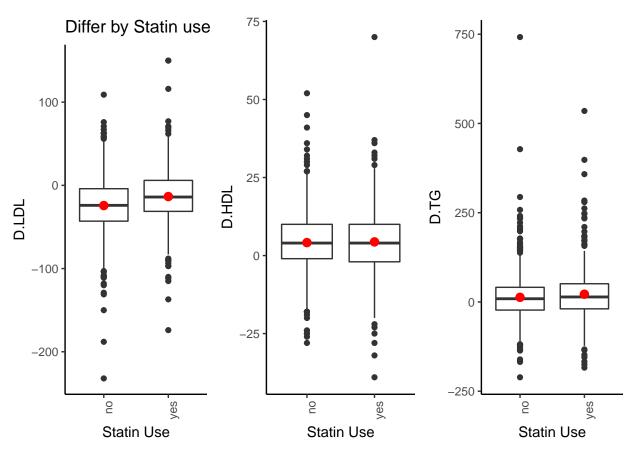


By looking at the boxplots (SE-Max, Mean, SE-Min), we could hypothesize that TG might be affected by race. We could do an ANOVA to confirm that.

For LDL, the effect is found to be not significant (p-value 0.96), but for TG and HDL, the effect do differ by race/ethnicity (p-value 0.01, and p-value < 0.001 respectively). The analysis was facilitated by ANOVA where null model included demograhics (excluding BMI),diabetic indicator and behavior. Race/Ethnicity was omitted in the null model.

```
## Analysis of Deviance Table
##
## Model 1: D.LDL ~ HT * raceth
## Model 2: D.LDL ~ age + smoking + drinkany + exercise + statins + diabetes
## Resid. Df Resid. Dev Df Deviance Pr(>Chi)
## 1 2574 2955598
## 2 2573 3087299 1 -131701
```

Checking by Statin Use

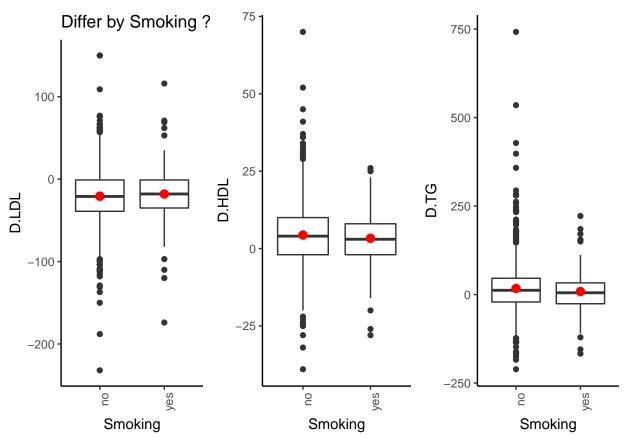


By looking at the boxplots(SE-Max, Mean, SE-Min), we could hypothesize that LDL might be affected by statin use We could do an ANOVA to confirm that.

```
## Analysis of Deviance Table
##
## Model 1: D.LDL ~ raceth + age + smoking + drinkany + exercise + statins +
##
       diabetes
## Model 2: D.LDL ~ raceth + age + smoking + drinkany + exercise + diabetes
##
     Resid. Df Resid. Dev Df Deviance Pr(>Chi)
          2571
                  3087210
## 1
          2572
## 2
                  3112096 -1
                                -24886 5.302e-06 ***
##
                   0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
## Signif. codes:
```

For TG, and HDL the effect is found to be not significant (p-value = 0.21, and p-value = 0.97 respectively), but for LDL, the effect do differ by statin use (p-value < 0.001). The analysis was facilitated by ANOVA where null model included demograhics (excluding BMI), diabetic indicator and behavior. Statin use was omitted in the null model.

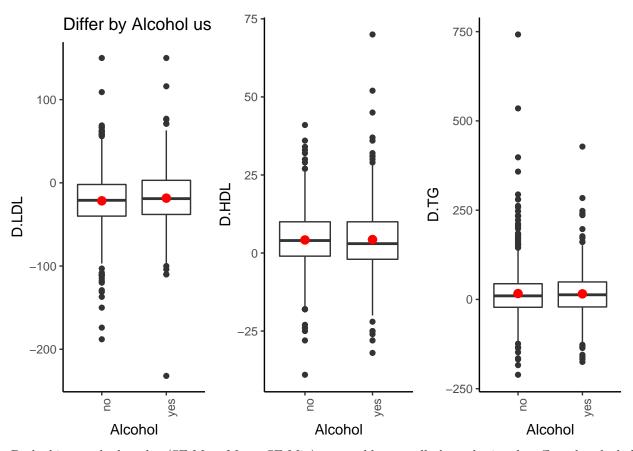
Smoking Behavior



By looking at the boxplots(SE-Max, Mean, SE-Min), we could not really hypothesize the affects by smoking behavior. We could do an ANOVA to confirm which ones are significant

For TG,LDL,HDL the effect is found to be not significant (p-value = 0.29, p-value = 0.42, and p-value = 0.99 respectively). The analysis was facilitated by ANOVA where null model included demograhics (excluding BMI),diabetic indicator and behavior. Smoking indicator was omitted in the null model.

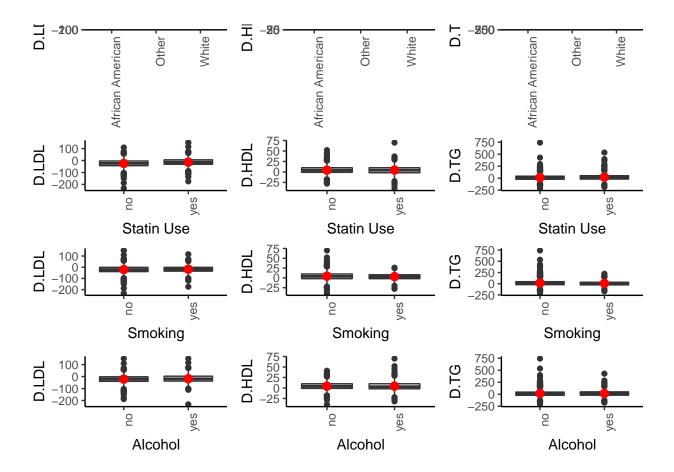
Alcohol use



By looking at the boxplots(SE-Max, Mean, SE-Min), we could not really hypothesize the affects by alcohol use. We could do an ANOVA to confirm which ones are significant

```
## Analysis of Deviance Table
##
## Model 1: D.TG ~ raceth + age + smoking + drinkany + exercise + statins +
## diabetes
## Model 2: D.TG ~ raceth + age + smoking + exercise + statins + diabetes
## Resid. Df Resid. Dev Df Deviance Pr(>Chi)
## 1 2571 11564115
## 2 2572 11564115 -1 -0.01827 0.9984
```

For TG,LDL,HDL the effect is found to be not significant(p-value = 0.99, p-value = 0.13, and p-value = 0.24 respectively). The analysis was facilitated by ANOVA where null model included demograhics(excluding BMI),diabetic indicator and behavior. Drinking indicator was omitted in the null model.

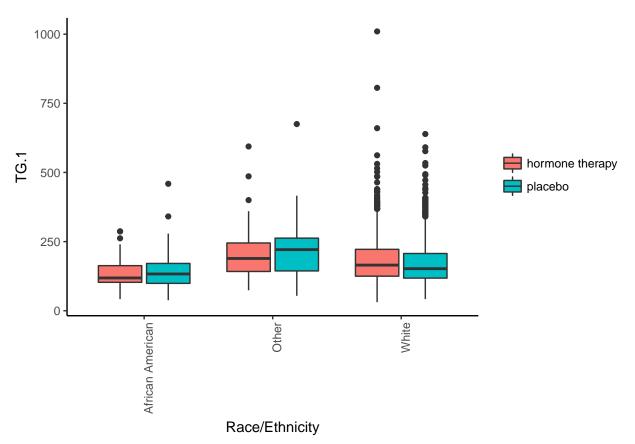


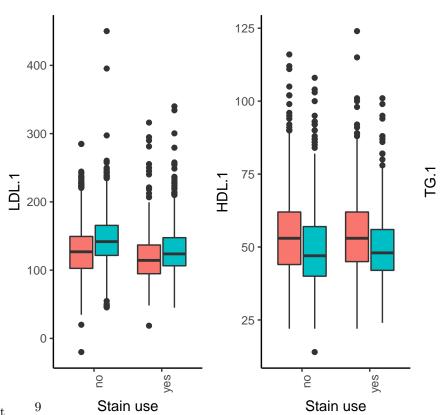
Changing direction

Methods : We tested the

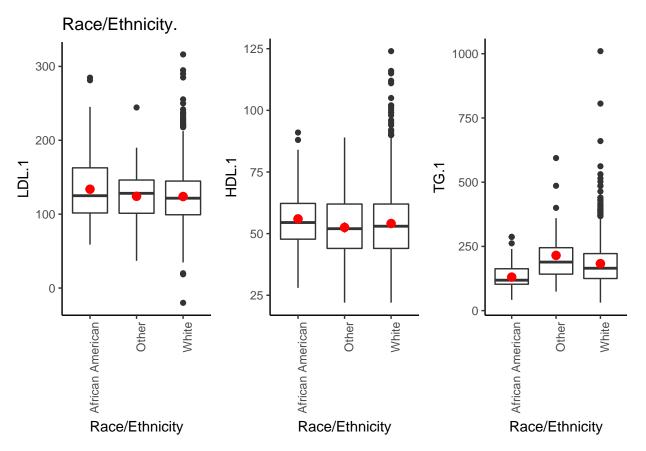
final figures

TG Treatment vs Placebo - Endpoint

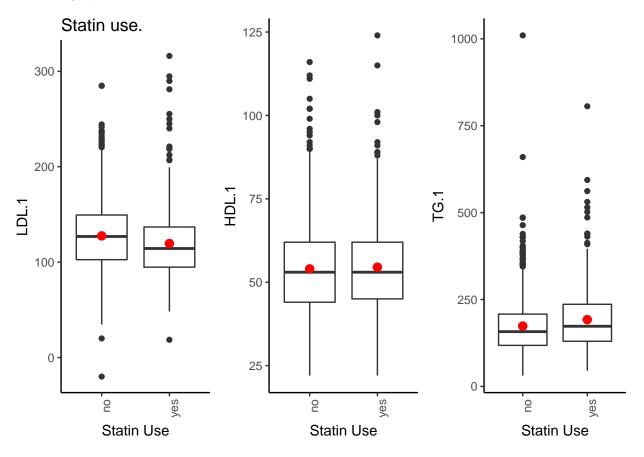




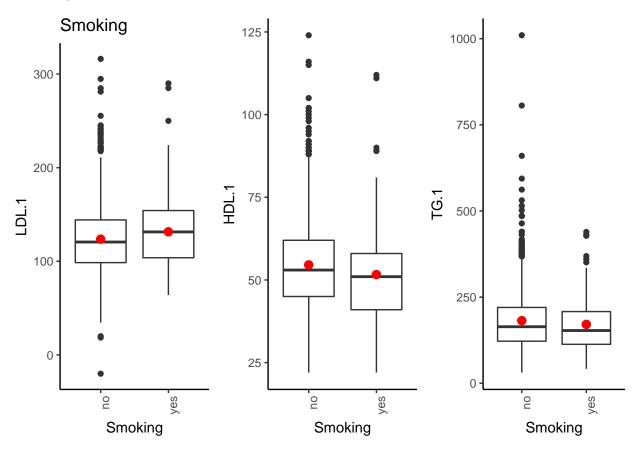
Checking by Race



Checking by Statin Use



Smoking Behavior



Alcohol use

