

## exercise5

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### Exercise 1

```
data_df = data.frame(X = c(0, 0, 0, 0, 1, 1, 1, 1), C = c(0, 0, 1, 1, 0, 0, 1, 1),  
  Y = c(0, 1, 0, 1, 0, 1, 0, 1), n = c(80, 20, 20, 10, 80, 20, 80, 40))
```

a

```
PX1cC0 = sum(data_df[(data_df$X == 1) & (data_df$C == 0), "n"])/sum(data_df[data_df$C ==  
  0, "n"])  
print(PX1cC0)
```

```
## [1] 0.5
```

```
PX1cC1 = sum(data_df[(data_df$X == 1) & (data_df$C == 1), "n"])/sum(data_df[data_df$C ==  
  1, "n"])  
print(PX1cC1)
```

```
## [1] 0.8
```

These probabilities are propensity scores of  $X=1$ .

b

```
PX0cC0 = 1 - PX1cC0  
PX0cC1 = 1 - PX1cC1  
WX1cC0 = 1/PX1cC0  
print(WX1cC0)
```

```
## [1] 2
```

```
WX1cC1 = 1/PX1cC1  
print(WX1cC1)
```

```
## [1] 1.25
```

```
WX0cC0 = 1/PX0cC0
print(WX0cC0)
```

```
## [1] 2
```

```
WX0cC1 = 1/PX0cC1
print(WX0cC1)
```

```
## [1] 5
```

c

```
data_df$W = 0
data_df[(data_df$X == 1) & (data_df$C == 0), "W"] = WX1cC0
data_df[(data_df$X == 1) & (data_df$C == 1), "W"] = WX1cC1
data_df[(data_df$X == 0) & (data_df$C == 0), "W"] = WX0cC0
data_df[(data_df$X == 0) & (data_df$C == 1), "W"] = WX0cC1
data_df$YxnW = data_df$Y * data_df$n * data_df$W
data_df$nxW = data_df$n * data_df$W
EY1 = sum(data_df[data_df$X == 1, "YxnW"])/sum(data_df[data_df$X == 1, "nxW"])
print(EY1)
```

```
## [1] 0.2571429
```

```
EY0 = sum(data_df[data_df$X == 0, "YxnW"])/sum(data_df[data_df$X == 0, "nxW"])
print(EY0)
```

```
## [1] 0.2571429
```

```
ATE = EY1 - EY0
print(ATE)
```

```
## [1] 0
```

## Exercise 2

```
library(cobalt)
```

```
## Warning: package 'cobalt' was built under R version 4.3.2
```

```
## cobalt (Version 4.5.4, Build Date: 2024-02-26)
```

```
library(survey)
```

```
## Warning: package 'survey' was built under R version 4.3.2
```

```
## Loading required package: grid
```

```
## Loading required package: Matrix
```

```
## Loading required package: survival
```

```
##
```

```
## Attaching package: 'survey'
```

```
## The following object is masked from 'package:graphics':
```

```
##
```

```
##      dotchart
```

```
library(ggplot2)
```

```
library(gridExtra)
```

```
library(survey)
```

```
load("rhc_exercise.RData")
```

a

```
ps_model = glm(treatment ~ transhx + age + scoma1 + hrt1 + bili1 + wtkilo1 + cat1,  
               data = rhc, family = "binomial")  
print(summary(ps_model))
```

```
##
```

```
## Call:
```

```
## glm(formula = treatment ~ transhx + age + scoma1 + hrt1 + bili1 +
```

```
##      wtkilo1 + cat1, family = "binomial", data = rhc)
```

```
##
```

```
## Coefficients:
```

```
##
```

	Estimate	Std. Error	z value	Pr(> z )	
## (Intercept)	-1.9073406	0.1967016	-9.697	< 2e-16	***
## transhx	0.5426888	0.0875449	6.199	5.68e-10	***
## age	0.0032199	0.0017884	1.800	0.07179	.
## scoma1	-0.0019473	0.0011566	-1.684	0.09224	.
## hrt1	0.0028841	0.0007398	3.899	9.67e-05	***
## bili1	0.0267466	0.0061889	4.322	1.55e-05	***
## wtkilo1	0.0098138	0.0013141	7.468	8.13e-14	***
## cat1CHF	0.3973320	0.1064659	3.732	0.00019	***
## cat1COPD	-1.3804068	0.1485325	-9.294	< 2e-16	***
## cat1Cirrhosis	-0.9184740	0.1743862	-5.267	1.39e-07	***
## cat1Colon Cancer	-1.1519303	1.0871074	-1.060	0.28931	
## cat1Coma	-0.5785267	0.1429332	-4.048	5.18e-05	***

```
## cat1Lung Cancer      -1.2690193  0.4826942  -2.629  0.00856 **
## cat1MOSF w/Malignancy 0.0984301  0.1127698   0.873  0.38275
## cat1MOSF w/Sepsis    0.7993355  0.0727869  10.982 < 2e-16 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for binomial family taken to be 1)
##
##      Null deviance: 7621.4  on 5734  degrees of freedom
## Residual deviance: 7049.6  on 5720  degrees of freedom
## AIC: 7079.6
##
## Number of Fisher Scoring iterations: 4
```

Except age and scoma1, other variables are strongly associated with treatment.

**b**

```
rhc$ps1 = predict(ps_model, newdata = rhc, type = "response")
rhc$ipw1 = 1/rhc$ps1
rhc$ipw1[rhc$treatment == "no RHC"] = 1/(1 - rhc$ps1[rhc$treatment == "no RHC"])
```

**c**

```
vars1 = c("transhx", "age", "scoma1", "hrt1", "bili1", "wtkilo1", "cat1", "surv2md1",
          "aps1")
covariates = rhc[, vars1]
table1 = bal.tab(covariates, treat = rhc$treatment, weights = rhc$ipw1, method = "weighting",
                un = TRUE)
```

## Note: 's.d.denom' not specified; assuming "pooled".

```
print(table1)
```

```
## Balance Measures
##
##      Type Diff.Un Diff.Adj
## transhx      Binary  0.0554  0.0007
## age          Contin. -0.0614  0.0005
## scoma1        Contin. -0.1098  0.0227
## hrt1          Contin.  0.1469 -0.0164
## bili1         Contin.  0.1446 -0.0075
## wtkilo1       Contin.  0.2256 -0.0012
## cat1_ARF      Binary -0.0290 -0.0040
## cat1_CHF      Binary  0.0261  0.0000
## cat1_COPD     Binary -0.0858  0.0002
## cat1_Cirrhosis Binary -0.0268  0.0001
## cat1_Colon Cancer Binary -0.0012  0.0000
## cat1_Coma     Binary -0.0525  0.0057
```

```
## cat1_Lung Cancer      Binary -0.0073 -0.0003
## cat1_MOSF w/Malignancy Binary  0.0045 -0.0008
## cat1_MOSF w/Sepsis    Binary  0.1721 -0.0010
## surv2md1             Contin. -0.1985 -0.2997
## aps1                 Contin.  0.5014  0.4165
##
## Effective sample sizes
##           no RHC      RHC
## Unadjusted 3551.    2184.
## Adjusted   3297.72 1738.58
```

```
# love.plot(covariates, treat = rhc$treatment, weights = rhc$ipw1, method =
# 'weighting', binary = 'std', threshold = .1)
```

After propensity score weight adjustment by transhx, age, scoma1, hrt1, bili1, wtkilo1, cat1, the treatment is still imbalanced on surv2md1 and aps1.

d

```
ps_model1 = glm(treatment ~ transhx + age + scoma1 + hrt1 + bili1 + wtkilo1 + cat1 +
  surv2md1 + aps1, data = rhc, family = "binomial")
print(summary(ps_model1))
```

```
##
## Call:
## glm(formula = treatment ~ transhx + age + scoma1 + hrt1 + bili1 +
##      wtkilo1 + cat1 + surv2md1 + aps1, family = "binomial", data = rhc)
##
## Coefficients:
##              Estimate Std. Error z value Pr(>|z|)
## (Intercept)    -1.0372749   0.3362492   -3.085  0.00204 **
## transhx         0.6860904   0.0904192    7.588 3.25e-14 ***
## age            -0.0051040   0.0020279   -2.517  0.01184 *
## scoma1         -0.0068209   0.0013238   -5.153 2.57e-07 ***
## hrt1           0.0014886   0.0007586    1.962  0.04973 *
## bili1         -0.0002230   0.0063827   -0.035  0.97213
## wtkilo1        0.0109611   0.0013536    8.098 5.60e-16 ***
## cat1CHF        0.9062462   0.1146630    7.904 2.71e-15 ***
## cat1COPD       -1.0939852   0.1519484   -7.200 6.03e-13 ***
## cat1Cirrhosis  -0.9407151   0.1779204   -5.287 1.24e-07 ***
## cat1Colon Cancer -1.0907678   1.1049168   -0.987  0.32355
## cat1Coma       -0.7041024   0.1532261   -4.595 4.32e-06 ***
## cat1Lung Cancer -1.1897262   0.4922679   -2.417  0.01566 *
## cat1MOSF w/Malignancy -0.6373686   0.1340213   -4.756 1.98e-06 ***
## cat1MOSF w/Sepsis  0.6492980   0.0754932    8.601 < 2e-16 ***
## surv2md1      -1.8602550   0.2608313   -7.132 9.89e-13 ***
## aps1          0.0177368   0.0019704    9.002 < 2e-16 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for binomial family taken to be 1)
```

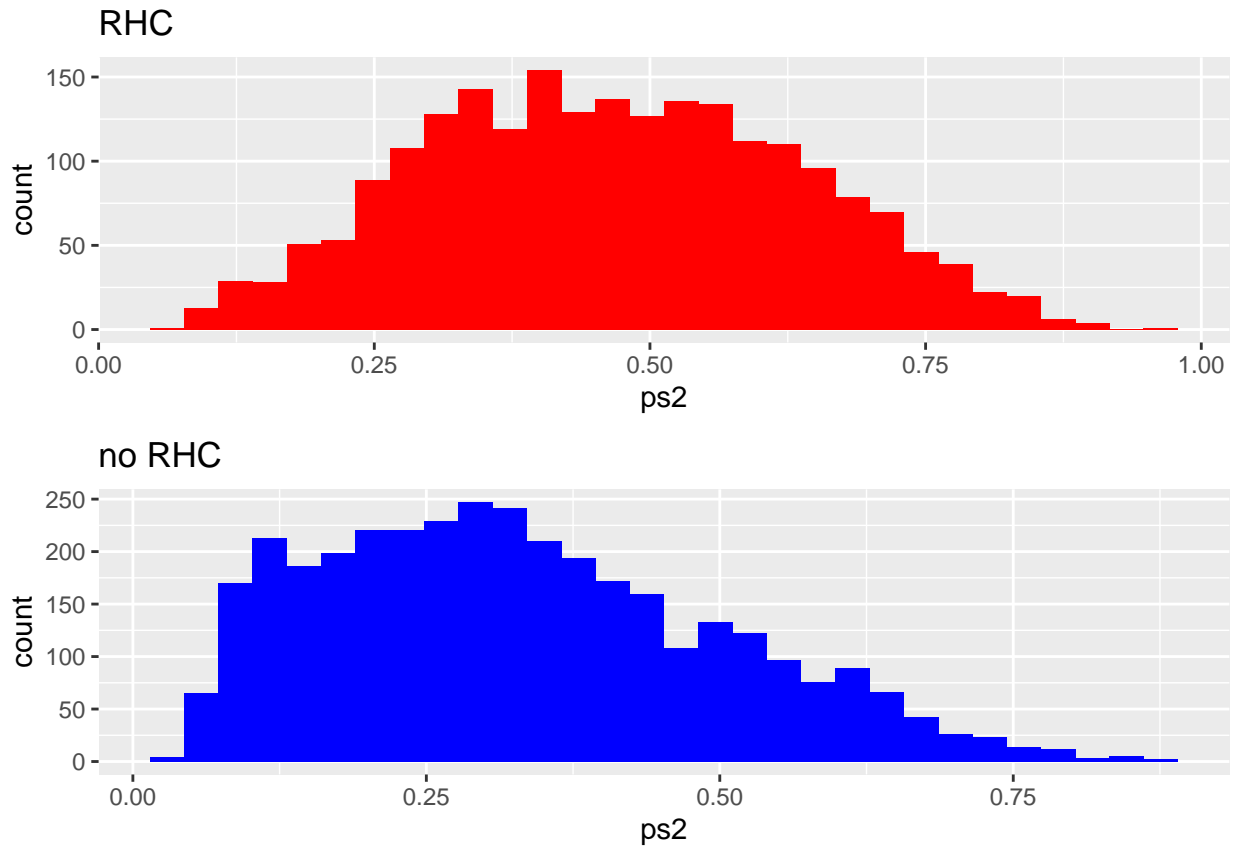
```
##
## Null deviance: 7621.4 on 5734 degrees of freedom
## Residual deviance: 6779.1 on 5718 degrees of freedom
## AIC: 6813.1
##
## Number of Fisher Scoring iterations: 4
```

```
rhc$ps2 = predict(ps_model1, newdata = rhc, type = "response")
print(summary(rhc$ps2))
```

```
## Min. 1st Qu. Median Mean 3rd Qu. Max.
## 0.03975 0.24215 0.36314 0.38082 0.51358 0.96591
```

```
p1 = ggplot(data = rhc[rhc$treatment == "RHC", ], mapping = aes(x = ps2)) + geom_histogram(fill = "red")
labs(title = "RHC")
p2 = ggplot(data = rhc[rhc$treatment == "no RHC", ], mapping = aes(x = ps2)) + geom_histogram(fill = "blue")
labs(title = "no RHC")
grid.arrange(p1, p2, nrow = 2)
```

```
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
```



The minimal ps2 is 0.04 and the maximal ps2 is 0.97, which means no individual will get a propensity score 1 or 0. The histograms show overlap between two treatment groups. Positivity assumption holds.

e

```
rhc$ipw2 = (rhc$treatment == "RHC") * 1/rhc$ps2 + (rhc$treatment == "no RHC") * 1/(1 -  
rhc$ps2)  
table2 = bal.tab(covariates, treat = rhc$treatment, weights = rhc$ipw2, method = "weighting",  
un = TRUE)
```

```
## Note: 's.d.denom' not specified; assuming "pooled".
```

```
print(table2)
```

```
## Balance Measures  
##  
##          Type Diff.Un Diff.Adj  
## transhx      Binary  0.0554   0.0018  
## age          Contin. -0.0614  -0.0023  
## scoma1       Contin. -0.1098  -0.0034  
## hrt1         Contin.  0.1469   0.0150  
## bili1        Contin.  0.1446   0.0060  
## wtkilo1      Contin.  0.2256   0.0045  
## cat1_ARF     Binary -0.0290   0.0061  
## cat1_CHF     Binary  0.0261   0.0034  
## cat1_COPD    Binary -0.0858  -0.0088  
## cat1_Cirrhosis Binary -0.0268  -0.0022  
## cat1_Colon Cancer Binary -0.0012  0.0001  
## cat1_Coma     Binary -0.0525  -0.0055  
## cat1_Lung Cancer Binary -0.0073  -0.0014  
## cat1_MOSF w/Malignancy Binary  0.0045  0.0036  
## cat1_MOSF w/Sepsis Binary  0.1721  0.0047  
## surv2md1     Contin. -0.1985  -0.0187  
## aps1         Contin.  0.5014   0.0235  
##  
## Effective sample sizes  
##          no RHC    RHC  
## Unadjusted 3551.   2184.  
## Adjusted   3116.93 1680.32
```

The distribution of all covariates become balanced after reweighting by ps2.

f

```
print(table(rhc$death30))
```

```
##  
## alive death  
## 3817 1918
```

```
rhc$death30 = as.numeric(rhc$death30) - 1  
print(table(rhc$death30))
```

```
##  
##    0    1  
## 3817 1918
```

g

```
EY1 = weighted.mean(x = rhc[rhc$treatment == "RHC", "death30"], w = rhc[rhc$treatment ==  
  "RHC", "ipw2"])  
EY0 = weighted.mean(x = rhc[rhc$treatment == "no RHC", "death30"], w = rhc[rhc$treatment ==  
  "no RHC", "ipw2"])  
ATE = EY1 - EY0  
print(c(EY1 = EY1, EY0 = EY0, ATE = ATE))
```

```
##           EY1           EY0           ATE  
## 0.35799263 0.32073365 0.03725898
```

h

```
d.w = svydesign(~1, weights = rhc$ipw2, data = rhc)  
w_reg = svyglm(death30 ~ treatment, design = d.w)  
print(summary(w_reg))
```

```
##  
## Call:  
## svyglm(formula = death30 ~ treatment, design = d.w)  
##  
## Survey design:  
## svydesign(~1, weights = rhc$ipw2, data = rhc)  
##  
## Coefficients:  
##              Estimate Std. Error t value Pr(>|t|)  
## (Intercept)   0.32073    0.00851  37.689 < 2e-16 ***  
## treatmentRHC  0.03726    0.01432   2.602  0.00929 **  
## ---  
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1  
##  
## (Dispersion parameter for gaussian family taken to be 0.2238068)  
##  
## Number of Fisher Scoring iterations: 2
```

```
print(confint(w_reg))
```

```
##              2.5 %    97.5 %  
## (Intercept) 0.304050792 0.3374165  
## treatmentRHC 0.009187664 0.0653303
```

The 95% confidence interval of ATE is (0.009, 0.065)

i



```
summary(rhc$ipw2)
```

```
##      Min. 1st Qu.  Median    Mean 3rd Qu.    Max.
##    1.035   1.340   1.643   1.985   2.240  15.452
```

The individual with very large weights can have big effect on the ATE. If they are outliers, the final results will be biased. If weights are very large, they are usually truncated.

j

```
ipw2_99th = quantile(rhc$ipw2, probs = 0.99)
rhc$ipw2_trunc = ifelse(rhc$ipw2 > ipw2_99th, ipw2_99th, rhc$ipw2)
EY1 = weighted.mean(x = rhc[rhc$treatment == "RHC", "death30"], w = rhc[rhc$treatment ==
  "RHC", "ipw2_trunc"])
EY0 = weighted.mean(x = rhc[rhc$treatment == "no RHC", "death30"], w = rhc[rhc$treatment ==
  "no RHC", "ipw2_trunc"])
ATE = EY1 - EY0
print(c(EY1 = EY1, EY0 = EY0, ATE = ATE))
```

```
##           EY1           EY0           ATE
## 0.35962584 0.32077150 0.03885434
```

ATE becomes a little bit larger than weighting by ipw2 but still very similar.

## Exercise 3

```
load("rhc_exercise.RData")
```

```
library(MatchIt)
```

```
##
## Attaching package: 'MatchIt'

## The following object is masked from 'package:cobalt':
##
##      lalonde
```

a

```
match_out1 = matchit(treatment ~ transhx + age + scoma1 + hrt1 + bili1 + wtkilo1 +
  cat1 + surv2md1 + aps1, data = rhc, method = "nearest", distance = "glm")
print(summary(match_out1))
```

```
##
## Call:
## matchit(formula = treatment ~ transhx + age + scoma1 + hrt1 +
##      bili1 + wtkilo1 + cat1 + surv2md1 + aps1, data = rhc, method = "nearest",
##      distance = "glm")
##
## Summary of Balance for All Data:
```

	Means Treated	Means Control	Std. Mean Diff.	Var. Ratio
distance	0.4658	0.3286	0.8097	1.0208
transhx	0.1497	0.0943	0.1552	.
age	60.7498	61.7609	-0.0647	0.8175
scoma1	18.9734	22.2532	-0.1160	0.8116
hrt1	118.9281	112.8730	0.1460	1.0260
bili1	2.7057	1.9973	0.1329	1.4504
wtkilo1	78.2336	73.2753	0.2298	0.9297
cat1ARF	0.4162	0.4452	-0.0589	.
cat1CHF	0.0957	0.0696	0.0889	.
cat1COPD	0.0266	0.1124	-0.5337	.
cat1Cirrhosis	0.0224	0.0493	-0.1813	.
cat1Colon Cancer	0.0005	0.0017	-0.0576	.
cat1Coma	0.0435	0.0960	-0.2575	.
cat1Lung Cancer	0.0023	0.0096	-0.1524	.
cat1MOSF w/Malignancy	0.0723	0.0679	0.0173	.
cat1MOSF w/Sepsis	0.3205	0.1484	0.3688	.
surv2md1	0.5685	0.6072	-0.1954	1.0663
aps1	60.7390	50.9335	0.4837	1.1609

```
##
##      eCDF Mean eCDF Max
## distance      0.2181  0.3183
## transhx      0.0554  0.0554
## age          0.0285  0.0703
## scoma1       0.0346  0.0683
## hrt1         0.0344  0.0782
## bili1       0.0244  0.1119
## wtkilo1     0.0535  0.1252
## cat1ARF     0.0290  0.0290
## cat1CHF     0.0261  0.0261
## cat1COPD    0.0858  0.0858
## cat1Cirrhosis 0.0268  0.0268
## cat1Colon Cancer 0.0012  0.0012
## cat1Coma    0.0525  0.0525
## cat1Lung Cancer 0.0073  0.0073
## cat1MOSF w/Malignancy 0.0045  0.0045
## cat1MOSF w/Sepsis 0.1721  0.1721
## surv2md1   0.0475  0.0957
## aps1       0.0797  0.2127
##
## Summary of Balance for Matched Data:
```

	Means Treated	Means Control	Std. Mean Diff.	Var. Ratio
distance	0.4658	0.4209	0.2650	1.4091
transhx	0.1497	0.1241	0.0719	.
age	60.7498	60.8612	-0.0071	0.7863
scoma1	18.9734	18.7161	0.0091	0.9883
hrt1	118.9281	117.8668	0.0256	0.9871
bili1	2.7057	2.3190	0.0726	1.1183

```

## wtkilo1          78.2336      75.8427      0.1108      0.8740
## cat1ARF          0.4162      0.4858      -0.1412      .
## cat1CHF          0.0957      0.1067      -0.0374      .
## cat1COPD         0.0266      0.0201      0.0399      .
## cat1Cirrhosis    0.0224      0.0270      -0.0309      .
## cat1Colon Cancer 0.0005      0.0000      0.0214      .
## cat1Coma         0.0435      0.0467      -0.0157      .
## cat1Lung Cancer  0.0023      0.0014      0.0192      .
## cat1MOSF w/Malignancy 0.0723      0.0783      -0.0230      .
## cat1MOSF w/Sepsis 0.3205      0.2340      0.1854      .
## surv2md1         0.5685      0.5867      -0.0921      0.9837
## aps1             60.7390      56.9679      0.1860      1.1814
##
## eCDF Mean eCDF Max Std. Pair Dist.
## distance         0.0593      0.1461      0.2651
## transhx          0.0256      0.0256      0.6442
## age              0.0285      0.0641      1.1914
## scoma1           0.0074      0.0247      0.9372
## hrt1             0.0089      0.0266      1.0963
## bili1            0.0144      0.0943      0.5661
## wtkilo1          0.0297      0.0723      1.0410
## cat1ARF          0.0696      0.0696      0.8880
## cat1CHF          0.0110      0.0110      0.6257
## cat1COPD         0.0064      0.0064      0.2050
## cat1Cirrhosis    0.0046      0.0046      0.3154
## cat1Colon Cancer 0.0005      0.0005      0.0214
## cat1Coma         0.0032      0.0032      0.4108
## cat1Lung Cancer  0.0009      0.0009      0.0766
## cat1MOSF w/Malignancy 0.0060      0.0060      0.5355
## cat1MOSF w/Sepsis 0.0865      0.0865      0.6387
## surv2md1         0.0228      0.0582      1.1004
## aps1             0.0320      0.0966      0.9154
##
## Sample Sizes:
##           Control Treated
## All           3551      2184
## Matched       2184      2184
## Unmatched     1367         0
## Discarded         0         0

```

ATT is being targeted. 2184 treated individuals and 2184 control individuals are matched. After matching, there is still imbalance in some covariates. For example, the standard mean difference of wtkilo1 on matched data is  $0.1108 > 0.1$ .

**b**

```

match_out2 = matchit(treatment ~ transhx + age + scoma1 + hrt1 + bili1 + wtkilo1 +
  cat1 + surv2md1 + aps1, data = rhc, method = "nearest", distance = "glm", caliper = 0.1)
print(summary(match_out2))

```

```

##
## Call:

```

```
## matchit(formula = treatment ~ transhx + age + scoma1 + hrt1 +
##      bili1 + wtkilo1 + cat1 + surv2md1 + aps1, data = rhc, method = "nearest",
##      distance = "glm", caliper = 0.1)
##
## Summary of Balance for All Data:
##
```

	Means Treated	Means Control	Std. Mean Diff.	Var. Ratio
distance	0.4658	0.3286	0.8097	1.0208
transhx	0.1497	0.0943	0.1552	.
age	60.7498	61.7609	-0.0647	0.8175
scoma1	18.9734	22.2532	-0.1160	0.8116
hrt1	118.9281	112.8730	0.1460	1.0260
bili1	2.7057	1.9973	0.1329	1.4504
wtkilo1	78.2336	73.2753	0.2298	0.9297
cat1ARF	0.4162	0.4452	-0.0589	.
cat1CHF	0.0957	0.0696	0.0889	.
cat1COPD	0.0266	0.1124	-0.5337	.
cat1Cirrhosis	0.0224	0.0493	-0.1813	.
cat1Colon Cancer	0.0005	0.0017	-0.0576	.
cat1Coma	0.0435	0.0960	-0.2575	.
cat1Lung Cancer	0.0023	0.0096	-0.1524	.
cat1MOSF w/Malignancy	0.0723	0.0679	0.0173	.
cat1MOSF w/Sepsis	0.3205	0.1484	0.3688	.
surv2md1	0.5685	0.6072	-0.1954	1.0663
aps1	60.7390	50.9335	0.4837	1.1609

```
##
##      eCDF Mean eCDF Max
## distance      0.2181  0.3183
## transhx       0.0554  0.0554
## age           0.0285  0.0703
## scoma1        0.0346  0.0683
## hrt1          0.0344  0.0782
## bili1         0.0244  0.1119
## wtkilo1       0.0535  0.1252
## cat1ARF       0.0290  0.0290
## cat1CHF       0.0261  0.0261
## cat1COPD      0.0858  0.0858
## cat1Cirrhosis 0.0268  0.0268
## cat1Colon Cancer 0.0012  0.0012
## cat1Coma      0.0525  0.0525
## cat1Lung Cancer 0.0073  0.0073
## cat1MOSF w/Malignancy 0.0045  0.0045
## cat1MOSF w/Sepsis 0.1721  0.1721
## surv2md1     0.0475  0.0957
## aps1          0.0797  0.2127
##
## Summary of Balance for Matched Data:
##
```

	Means Treated	Means Control	Std. Mean Diff.	Var. Ratio
distance	0.4379	0.4296	0.0488	1.0901
transhx	0.1331	0.1259	0.0202	.
age	60.9422	61.0129	-0.0045	0.7935
scoma1	18.9162	18.8710	0.0016	0.9685
hrt1	118.0288	118.3258	-0.0072	0.9402
bili1	2.4512	2.3965	0.0103	0.8783
wtkilo1	77.0525	76.2229	0.0384	0.7715
cat1ARF	0.4419	0.4687	-0.0542	.

```

## cat1CHF                0.0961        0.1074        -0.0384        .
## cat1COPD               0.0298        0.0211         0.0543        .
## cat1Cirrhosis          0.0252        0.0272        -0.0139        .
## cat1Colon Cancer       0.0005        0.0000         0.0240        .
## cat1Coma               0.0483        0.0493        -0.0050        .
## cat1Lung Cancer        0.0026        0.0015         0.0215        .
## cat1MOSF w/Malignancy  0.0755        0.0776        -0.0079        .
## cat1MOSF w/Sepsis      0.2801        0.2472         0.0705        .
## surv2md1               0.5796        0.5815        -0.0093        0.9341
## aps1                   58.5149       57.7081         0.0398        1.0368
##
## eCDF Mean eCDF Max Std. Pair Dist.
## distance              0.0105     0.0421         0.0489
## transhx               0.0072     0.0072         0.6049
## age                   0.0269     0.0601         1.1907
## scoma1                0.0087     0.0252         0.9512
## hrt1                  0.0088     0.0334         1.0680
## bili1                 0.0053     0.0807         0.5481
## wtkilo1               0.0187     0.0550         1.0392
## cat1ARF               0.0267     0.0267         0.8924
## cat1CHF               0.0113     0.0113         0.6044
## cat1COPD              0.0087     0.0087         0.2205
## cat1Cirrhosis         0.0021     0.0021         0.3331
## cat1Colon Cancer      0.0005     0.0005         0.0240
## cat1Coma              0.0010     0.0010         0.4434
## cat1Lung Cancer       0.0010     0.0010         0.0860
## cat1MOSF w/Malignancy 0.0021     0.0021         0.5554
## cat1MOSF w/Sepsis     0.0329     0.0329         0.6166
## surv2md1              0.0128     0.0391         1.0847
## aps1                  0.0094     0.0344         0.8948
##
## Sample Sizes:
##           Control Treated
## All           3551    2184
## Matched       1946    1946
## Unmatched     1605    238
## Discarded           0      0

```

Matched individuals become less. 1946 treated individuals and 1946 control individuals are matched. After matching, there is no imbalance in covariates.

**c**

```
library(dplyr)
```

```

##
## Attaching package: 'dplyr'

## The following object is masked from 'package:gridExtra':
##
## combine

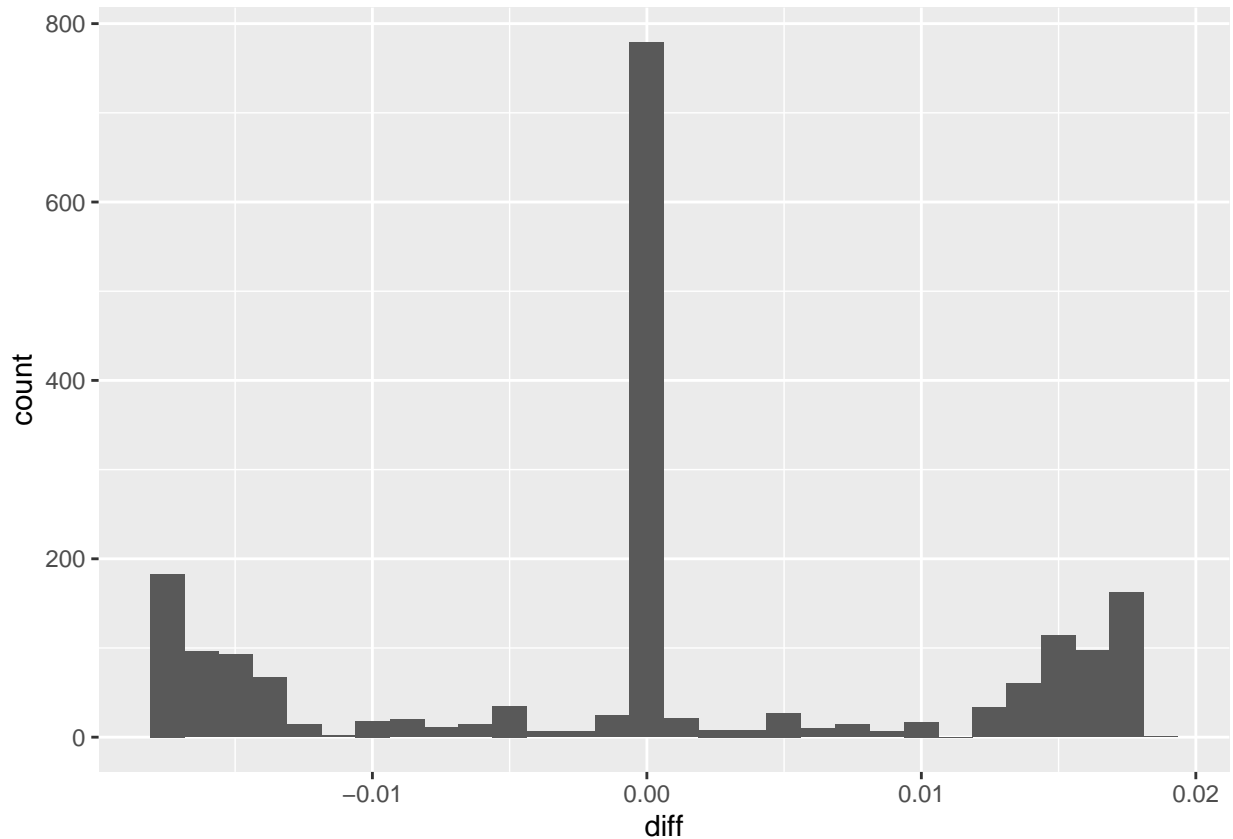
```

```
## The following objects are masked from 'package:stats':
##
##   filter, lag
```

```
## The following objects are masked from 'package:base':
##
##   intersect, setdiff, setequal, union
```

```
gen_diff = function(data) {
  return(data[1] - data[2])
}
match_rhc = match.data(match_out2)
match_rhc = match_rhc[order(match_rhc$subclass, decreasing = FALSE), ]
ps_diff = summarise(group_by(match_rhc, subclass), diff = gen_diff(distance))
ggplot(data = ps_diff, mapping = aes(x = diff)) + geom_histogram()
```

```
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
```



As shown on the histogram of propensity score difference within matched pairs, the propensity scores are very similar within matched pairs.

**d**

```

match_rhc$death30 = as.numeric(match_rhc$death30) - 1
EY1 = mean(match_rhc[match_rhc$treatment == "RHC", "death30"])
EY0 = mean(match_rhc[match_rhc$treatment == "no RHC", "death30"])
ATE = EY1 - EY0
print(c(EY1 = EY1, EY0 = EY0, ATE = ATE))

```

```

##           EY1           EY0           ATE
## 0.36742035 0.32785200 0.03956835

```

ATE is 0.040 which is larger than the result 0.039 of 2.j.