# Reanalysis of 03-Zecca

Simon Schwab

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#### Reference

Zecca, E., Brunelli, C., Centurioni, F., Manzoni, A., Pigni, A., & Caraceni, A. (2017). Fentanyl Sublingual Tablets Versus Subcutaneous Morphine for the Management of Severe Cancer Pain Episodes in Patients Receiving Opioid Treatment: A Double-Blind, Randomized, Noninferiority Trial. *Journal of Clinical Oncology*, 35(7), 759–765. https://doi.org/10.1200/JCO.2016.69.9504

#### Notes from reading methods section

- Dependent variable: Pain intensity (PI) (average across follow-ups, AVP 30)
- Independant variable: treatment group
  - Fentanyl sublingual tablets FST (n=57, 1 from 58 was excluded)
  - Subcutaneous morphine SCM (n=56)
- Covariate: basline PI
- Design: 1-way ANCOVA with group as IV and baselie PI as covariate

# Reading data

Data is loaded, reshaped if necessary, and factors are specified.

```
PATH = file.path(path.expand("~"), "Data", "ancova") # ancova project folder

data = read_excel(file.path(PATH, "dataPrimaryStudies", "03-Zecca", "03-Zecca.xls"))

data = as.data.frame(data)

data$tratt = as.factor(data$tratt)
```

# Descriptives

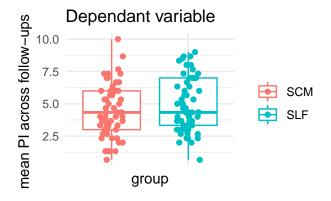
#### Dependant variable

Number of samples and mean (SD) in levels of the independent variables.

```
colnames(tab.dv) = c("group", "n", "mean (SD)")
print(tab.dv)

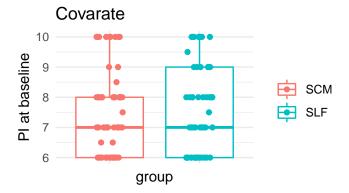
## group n mean (SD)
## [1,] "SCM" "56" "4.48 (2.03)"
## [2,] "SLF" "57" "4.97 (2.12)"

ggplot(data, aes(y=pain_mea_30, x=tratt, color=tratt)) +
    geom_boxplot() +
    geom_point(position = position_jitter(width = 0.15, height = 0)) +
    theme_minimal() +
    theme(axis.text.x = element_blank(), legend.title = element_blank()) +
    xlab("group") + ylab("mean PI across follow-ups ") + ggtitle("Dependant variable")
```



#### Covariate(s)

```
tab.cv = array(NA, dim=c(2,3))
tab.cv[,1] = levels(data$tratt)
tab.cv[,2] = summary(data$tratt)
tab.cv[,3] = tapply(data$pain_mea_bas, data$tratt,
                  function (x) sprintf("%0.2f (%0.2f)", mean(x), sd(x)))
colnames(tab.cv) = c("group", "n", "mean (SD)")
print(tab.cv)
##
                   mean (SD)
        group n
## [1,] "SCM" "56" "7.54 (1.39)"
## [2,] "SLF" "57" "7.53 (1.45)"
ggplot(data, aes(y=pain_mea_bas, x=tratt, color=tratt)) +
  geom_boxplot() +
  geom_point(position = position_jitter(width = 0.15, height = 0)) +
  theme_minimal() +
  theme(axis.text.x = element_blank(), legend.title = element_blank()) +
  xlab("group") + ylab("PI at baseline") + ggtitle("Covarate")
```



### Main analysis ANCOVA

```
# Orthogonal contrasts
contrasts(data$tratt) = contr.helmert(2)
fit.ancova = aov(pain_mea_30 ~ pain_mea_bas + tratt, data = data)
result = Anova(fit.ancova, type=3) # Type III
print(result)
## Anova Table (Type III tests)
## Response: pain_mea_30
##
                 Sum Sq Df F value
                                       Pr(>F)
                 24.426
                          1 9.9081 0.002118 **
## (Intercept)
## pain_mea_bas 207.658
                          1 84.2329 2.989e-15 ***
                             2.8047 0.096830 .
## tratt
                  6.914
                          1
## Residuals
                271.182 110
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
The paper used ANCOVA to to a non-inferiority analysis. We calculate ANCOVA adjusted means and
95%-CL
means.adj = effect("tratt", fit.ancova, se=TRUE)
MD = means.adj$fit[1] - means.adj$fit[2]
# SE is smaller as covarate reduced noise
SE = sqrt(means.adj$se[1]^2 + means.adj$se[2]^2)
ci95 = c(MD - 1.96*SE, MD + 1.96*SE)
sprintf("%.2f (%.2f, %.2f)", MD, ci95[1], ci95[2])
## [1] "-0.49 (-1.07, 0.08)"
```

# Comparing ANCOVA in original study with reanalysis

Independent variable

```
tab.IV = rbind(stats.orig.IV, stats.rep.IV)
rownames(tab.IV) = c("original Study", "reanalysis")
print(t(tab.IV))
```

```
##
           original Study reanalysis
                           "2.80"
## Fvalue
           NA
## df1
           NA
                           " 1"
## df2
           NA
                           "110"
## pvalue NA
                           "0.097"
           "-0.49"
## MD
                           "-0.49"
## lowerCI "-1.1"
                           "-1.07"
## upperCI "0.09"
                           "0.08"
```

#### Covariate

```
tab.CV = rbind(stats.orig.CV, stats.rep.CV)
rownames(tab.CV) = c("original Study", "reanalysis")
print(t(tab.CV))
## original Study reanalysis
```

```
original Study reanalysis
## Fvalue
                           "84.23"
                           " 1"
## df1
           NA
                           "110"
## df2
           NA
                           "< 0.0001"
## pvalue NA
## MD
           NA
                           NA
## lowerCI NA
                           NA
## upperCI NA
                           NA
```

# Assumptions

#### 1. Homogeneity of variance

- ANOVA/ANCOVA is fairly robust in terms of the error rate when sample sizes are equal.
- When groups with larger sample sizes have larger variances than the groups with smaller sample sizes, the resulting F-ratio tends to be conservative. That is, it's more likely to produce a non-significant result when a genuine difference does exist in the population.
- Conversely, when the groups with larger sample sizes have smaller variances than the groups with smaller samples sizes, the resulting F-ratio tends to be liberal and can inflate the false positive rate.

```
tapply(data$pain_mea_30, data$tratt, sd)

## SCM SLF

## 2.030992 2.121189

leveneTest(pain_mea_30 ~ tratt, data = data)

## Levene's Test for Homogeneity of Variance (center = median)

## Df F value Pr(>F)

## group 1 0.3764 0.5408

## 111
```

#### 2. Independence between covariate and IV

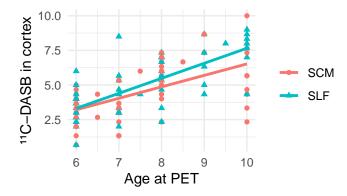
When the covariate and the experimental effect (independent variable) are not independent the treatment effect is obscured, spurious treatment effects can arise and the interpretation of the ANCOVA is seriously compromised.

We test whether our groups differ on the CV. If the groups do not significantly differ then is appropriate to use the covariate.

```
fit.cv = aov(pain_mea_bas ~ tratt, data = data)
Anova(fit.cv, type=3)
## Anova Table (Type III tests)
##
## Response: pain_mea_bas
                           F value Pr(>F)
##
              Sum Sq Df
## (Intercept) 6408.4
                      1 3180.7293 <2e-16 ***
                            0.0012 0.972
## tratt
                 0.0
                       1
## Residuals
               223.6 111
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
```

#### 3. Homogeneity of regression slopes

```
fit.hrs = aov(pain_mea_30 ~ pain_mea_bas*tratt, data = data)
Anova(fit.hrs, type=3)
## Anova Table (Type III tests)
##
## Response: pain_mea_30
##
                       Sum Sq Df F value
                                             Pr(>F)
## (Intercept)
                       23.374
                                1 9.5251
                                            0.00257 **
## pain_mea_bas
                      204.174
                                1 83.2032 4.304e-15 ***
## tratt
                        1.988
                                1
                                  0.8100
                                            0.37011
                        3.704
                                   1.5096
                                            0.22185
## pain_mea_bas:tratt
                                1
## Residuals
                      267.477 109
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
ggplot(data, aes(y=pain_mea_30, x=pain_mea_bas, color=tratt, shape=tratt)) +
  geom_point() +
  geom_smooth(formula = y ~ x,method=lm, se=FALSE, fullrange=TRUE) +
  theme_minimal() +
  theme(legend.title = element_blank()) +
  xlab("Age at PET") + ylab("11C-DASB in cortex")
```



Data was analyzed according to recommendations by Field, Miles, & Field (2012).