1. Let, 
$$X \sim N_3$$
 (  $M = \begin{pmatrix} -2 \\ 1 \\ 2 \end{pmatrix}$  ,  $S = \begin{pmatrix} 4 & 1 & 0 \\ 1 & 2 & 1 \\ 0 & 1 & 3 \end{pmatrix}$ )

(a) Let, 
$$y = \begin{pmatrix} x_1 \\ x_3 \end{pmatrix}$$

So, 
$$\frac{y}{x} \sim N_2 \left( \frac{y}{y} = \begin{pmatrix} -3 \\ 2 \end{pmatrix} \right) = \begin{pmatrix} 4 & 0 \\ 0 & 3 \end{pmatrix}$$

D

we know, 
$$\frac{1}{2}NN_2\left(lly=\left(-\frac{3}{2}\right), \frac{2}{2}y=\left(40\atop 03\right)\right)$$
 and  $\frac{1}{2}NN(1,2)$ 

So, 
$$\begin{pmatrix} \chi \\ \chi_2 \end{pmatrix} \sim N_3 \begin{pmatrix} \begin{bmatrix} uy \\ 1 \end{bmatrix} \end{pmatrix}$$
 Snew =  $\begin{bmatrix} 40 & 1 \\ 03 & 1 \end{bmatrix}$   $= \begin{bmatrix} 2 & 2 \\ 1 & 1 \end{bmatrix}$   $= \begin{bmatrix} 2 & 2 \\ 2 & 2 \end{bmatrix}$ 

Using the result of conditional multivariate normal distribution,

Y ( Xa=n2 ~ N2 ( My + E12 Sat ( 22-1), E11 - E12 Sat E21)

$$\equiv N_{\lambda} \left( \begin{pmatrix} -3 \\ 2 \end{pmatrix} + \begin{pmatrix} 1 \\ 1 \end{pmatrix} \cdot \begin{pmatrix} 24 - 1 \\ 2 \end{pmatrix} \right) \cdot \begin{pmatrix} 40 \\ 03 \end{pmatrix} - \begin{pmatrix} 5 \\ 5 \\ 5 \end{pmatrix} \right)$$

$$= N_{2} \left( \left( \frac{-3 + \frac{2y-1}{2}}{2} \right) \right) = \frac{2y_{2}}{2} \left( \frac{3\cdot 5}{-0\cdot 5} - 0\cdot 5 \right)$$

$$= N_2 \left( \frac{\alpha_1 - 7}{\alpha_2 + 3} \right) = \chi_1 \chi_2 = \left( \frac{3.5}{-0.5} \right),$$

(c) Similarly,
$$X_{3} | X = X \quad N \quad (M_{3} + \sum_{j=1}^{j-1} (X - M_{j})_{j} \leq_{23} - \sum_{j=1}^{j} N \leq_{17}^{j-1} \leq_{12})$$

$$= N \left( 1 + (1 + 1) \left( \frac{4}{0} + \frac{0}{0} \right)^{-1} \left[ \frac{n_{4}}{n_{3}} \right] - \frac{1}{2} \left( \frac{1}{0} + \frac{1}{0} \right) \right]$$

$$= N \left( 1 + (0 \cdot 25 + 0 \cdot 33) \left( \frac{n_{4} + 3}{n_{3} - \lambda} \right)_{j} + 1 \cdot 42 \right)$$

$$= N \left( 1 + (0 \cdot 25 + 0 \cdot 33 n_{3} - 0 \cdot 66)_{j} + 1 \cdot 42 \right)$$

$$= N \left( (0 \cdot 25 n_{4} + 0 \cdot 33 n_{3} + 1 \cdot 09)_{j} + 1 \cdot 42 \right)$$

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$$= N \left( (0 \cdot 25 n_{4} + 0 \cdot 33 n_{4} + 1 \cdot 09)_{j} + 1 \cdot 42 \right)$$

$$= N \left( (0 \cdot 25 n_$$

Let, 
$$Z = \frac{1}{2} \times X = (130) \begin{pmatrix} x_1 \\ x_2 \\ x_3 \end{pmatrix}$$
  
So,  $Z \sim N \begin{pmatrix} 2 & 1 \\ 2 & 1 \end{pmatrix}$ ,
  
where,  $2 = (130) \begin{pmatrix} -3 \\ 1 \\ 2 \end{pmatrix} = 0$ 
  
 $2 = 28$ 

Thus, ZNN(0,28)

2. Let I be the concentration of a certain substance in liver tissue after the administration of a dray at 30 days (cindicated by 1/2) and 60 days (cindicated by 1/3) and 1000 1/1 cindicating the first administration.

Now 
$$y = \begin{pmatrix} 14 \\ 24 \\ 24 \end{pmatrix}$$
 and  $S = \begin{pmatrix} 30 & 10 & 14 \\ 10 & 15 & 4 \\ 14 & 4 & 37 \end{pmatrix}$ 

(a) Here, we want to test:

Ho: llg = llg vs Hj: not Ho.

So, we use the test statistic:

for we use the test statistic:

$$f = \frac{u_{A} - u_{A}}{(u_{A} - 1)u_{A}} T^{2}, \text{ where, } T^{2} = u(e_{X})'(e_{S}e')^{-1}(e_{X})$$

$$(u_{A} - 1)u_{A}$$

$$u_{A} = 25, (=(1 - 1 0), n = nauk(e)$$

$$= 2.$$

$$\left[\begin{pmatrix} 1 & -1 & 0 \\ 0 & 1 & -1 \end{pmatrix} \begin{pmatrix} 14 \\ 24 \\ 22 \end{pmatrix}\right]$$

= 111.43

Thus, 
$$f = \frac{(25-2)}{(25-1)} 11.43 = 53.39$$

The critical region is F<0.025 and F>4.349. So, as Fetat> 4349 we reject the well hypathesis at 5% level of significance and conclude that the mean consentrations of the substance et different times after the administration of the drag are not same.

(b) Let 
$$\frac{2}{3}$$
 denote the results for drug  $\frac{1}{3}$ .

So,  $\frac{2}{3} = \begin{pmatrix} 15 \\ 20 \\ 24 \end{pmatrix}$  and  $W = \begin{pmatrix} 26 & 12 & 10 \\ 12 & 17 & 9 \\ 10 & 8 & 43 \end{pmatrix}$ 

(i) Let the population covariance variance matrices for drugs A and B se the same.

b, an estimate of the proled sample variance is:

$$S = \frac{1}{proled} \left( \frac{(u_A - 1)}{s} \right) + \left( \frac{(u_B - 1)}{s} \right) W$$

$$\left( \frac{(u_A + u_B - 2)}{s} \right)$$

$$\left( \frac{(u_A + u_B - 2)}{s} \right)$$

$$= \frac{24}{40} \begin{pmatrix} 30 & 10 & 14 \\ 10 & 15 & 4 \\ 14 & 4 & 37 \end{pmatrix} + \frac{16}{40} \begin{pmatrix} 26 & 12 & 10 \\ 12 & 17 & 8 \\ 10 & 8 & 43 \end{pmatrix}$$

$$= \begin{pmatrix} 28.4 & 10.8 & 12.4 \\ 10.8 & 15.8 & 5.6 \\ 12.4 & 5.6 & 39.4 \end{pmatrix}$$

(ii) To perform bartletts test, we test 
$$H_0$$
:  $\Xi_A = \Xi_B \times S + H_1$ : not to. First, we find  $M = (N-K) \log |S| - \frac{S}{i-1} (n_i - 1) \log |S_i|$ , where,  $N = S_{ii} = 42$ ,  $K = 2B$ 

Next, 
$$e^{-1} = 1 - \frac{2p^2 + 3p - 1}{6(p+1)(k-1)} \left[ \sum_{i=1}^{k} \left( \frac{1}{w_i - 1} \right) - \frac{1}{N-k} \right]$$
 where,  $p = 3$ 

The test statistic is:  $Me^{-1} = -109.28$ Now, the critical pregion is greater than  $\chi^2_{1-d}$ ; (K-1) P(PH)  $= \chi^2_{0.95;6} = 12.59$ 

As, Me-1 \$ 12:59, we fail to reject Ho at 5% level of significance and thus conclude that the population variance-covariances metrices are not same.

(ill) colo Mahalanolis distance is estimated as:

$$= \begin{pmatrix} -1 \\ 4 \\ -2 \end{pmatrix} \begin{pmatrix} 28.4 & 10.8 & 12.4 \\ 10.8 & 15.8 & 5.6 \\ 12.4 & 5.6 & 39.4 \end{pmatrix} \begin{pmatrix} -1 \\ 4 \\ -2 \end{pmatrix}$$

= 1.78

D

(iv) To test Ho: W= lb vs H,: not Ho.

The test statistic is 72 mm d2(x,2)

The critical value is greater than  $(n_A + n_B - a) P F_{1-x}; P, n_A + n_B - P^{-1} = 0.1a$ 

As T2 > 0'12, we reject 16 at 5% level of significance and conclude that the means for the two doings are considered unequal. (V) Here, we test: c(ely-ela) = 0 vs Hz: not Ho, where,  $C = \begin{pmatrix} 1 & -1 & 0 \\ 0 & 1 & -1 \end{pmatrix}$ ,  $\alpha = \operatorname{reauk}(e) = \lambda$ . The test statistic is (14+16-2) 4 T2 where  $T^2 = \frac{\mu_A \mu_B}{\mu_A + \mu_B} \left( \frac{1}{\lambda} - \frac{1}{\lambda} \right) e'$ (ege!)  $T_C(\frac{1}{\lambda} - \frac{1}{\lambda})$ = 25+17-2-1 X14-27 (25+17-2) X2 = 14.27 The critical value is fix, y + 4 - 1 = 0.05 Now, as 6.96 > 0.05, we reject to at 5% level of significance and woulde that there is presence of doing by time interaction. 3. n=16, n=11,p=4. (a) ll, and lla supresents the mean vectors of the groups of Loys and giods respectively. itere, we tex Ho: 14=lla vs Hi: not Ho = 25x4 fors; 4,22 = 0.77 As The profiles of the loops and girls are not same. I

(6) To test: Ho: e(M1-Ma)=0 ve H1: wot Ho

where, 
$$C = \begin{cases} 1 & -1 & 0 & 0 \\ 0 & 1 & -1 & 0 \\ 0 & 0 & 1 & -1 \end{cases}$$
, scank $(C_1) = 3 = x(C_2)$ 

The test statistic is 4+42-9-1 T2

(4442-2)2

$$= \frac{16+11-3-1}{(16+11-2)3}$$
 8:3

= 2.70

The critical value is greater than  $f_{1-\alpha}$ ; x,  $y_1+y_2-y_{-1}=0$  0.12 As,  $\alpha.70>0.12$ , we reject the at 5% level of significance and thus combude that there is presence of group by time interaction. (c) It we reject the null hypothesis in part (b), we might further do post-we tasts such as pairwise comparison or even perform profile analysis to understand the nature of the interaction.

## PROBLEM 4.

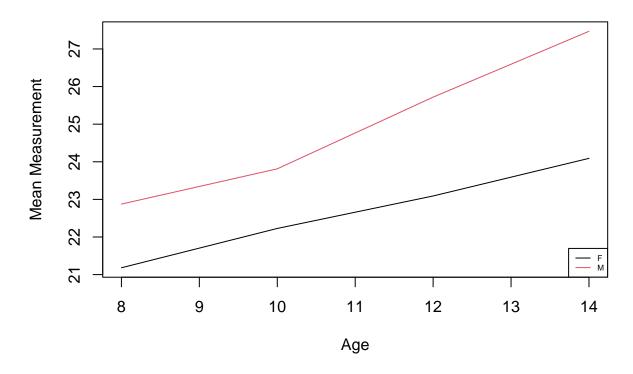
Part (a)

```
library(ggplot2)
library(nlme)
library(dplyr)
library(reshape2)
library(profileR)
data_4 = read.table("C:/Users/Jayaditya Nath/Documents/PothoffRoy1964.dat")

mean_profiles_4 = aggregate(. ~ data_4$V2, data = data_4[, -c(1, 2)], FUN = mean)

# Plot mean profiles
plot(seq(8,14,2), mean_profiles_4[1, -1], type = 'l', col = 1, ylim = c(min(mean_profiles_4[-1]), max(m lines(seq(8,14,2), mean_profiles_4[2, -1], col = 2)
legend("bottomright", legend = c("F", "M"), col = 1:2, lty = 1,cex = 0.5)
```

## Mean Profiles for different ages



From the profile plots, it is clearly visible that there is a difference between the dental measurements of males and females across different ages. The profiles are not at all similar to each other and males tend to have greater dental measurements compared to females at different ages.

Part (b)

```
# Performing MANOVA
anova(lm(cbind(V3,V4,V5,V6)~V2,data = data_4))
```

```
## Analysis of Variance Table
##
##
                 Df Pillai approx F num Df den Df Pr(>F)
## (Intercept) 1 0.9943
                              960.17
                                             4
                                                    22 < 2e-16 ***
                  1 0.3977
                                 3.63
                                                    22 0.02034 *
## Residuals
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
Here, we are interested in testing:
H_0: C(\mu_{males} - \mu_{females}) = \tilde{\mathbf{0}} \text{ vs } H_a: \text{not } H_0
```

The Pillai's test statistic for performing MANOVA is :

$$V = \sum_{i=1}^{p} \frac{\lambda_i}{1 + \lambda_i}$$

, where  $\lambda_i$  is the number of non-zero eigen values of the product of the inverse of the within-group variance-covariance matrix with the between-group variance-covariance matrix.

The p-value for the associated test is 0.02.

Thus, we are able to reject the null hypothesis at 5% level of significance and conclude that the dental measurements of femlaes and males vary across time.

Part (c)

```
## Linear mixed-effects model fit by REML
##
     Data: data_4_long
##
          AIC
                  BIC
                         logLik
##
     447.7572 466.268 -216.8786
##
## Random effects:
   Formula: ~1 | ID
##
##
           (Intercept) Residual
              1.816214 1.386382
## StdDev:
##
## Correlation Structure: Compound symmetry
  Formula: ~1 | ID
## Parameter estimate(s):
## Rho
## Fixed effects: Measurement ~ Gender * Age
##
                   Value Std.Error DF t-value p-value
```

```
## (Intercept) 17.372727 1.1835071 79 14.679023
               -1.032102 1.5374208 25 -0.671321
## GenderM
                                                  0.5082
                0.479545 0.0934698 79
                                       5.130483
                                                  0.0000
## GenderM:Age 0.304830 0.1214209 79 2.510520
                                                  0.0141
##
   Correlation:
##
               (Intr) GendrM Age
## GenderM
               -0.770
               -0.869 0.669
## Age
## GenderM: Age 0.669 -0.869 -0.770
##
## Standardized Within-Group Residuals:
##
           Min
                        Q1
                                   Med
                                                            Max
## -3.59804400 -0.45461690
                            0.01578365
                                        0.50244658
                                                     3.68620792
##
## Number of Observations: 108
## Number of Groups: 27
```

From the linear mixed model with compound symmetry variance-covariance structure, we can see that the main effect **Gender** is not significant, though **Age** seems to be quite significant. The interaction term is also significant in the model. I feel that somehow the MANOVA and the LMM model results are at agreement with each other in terms of the fact that the interaction between age and gender is significant, however, they disagree on the fact that **Gender** itself is significant or not.

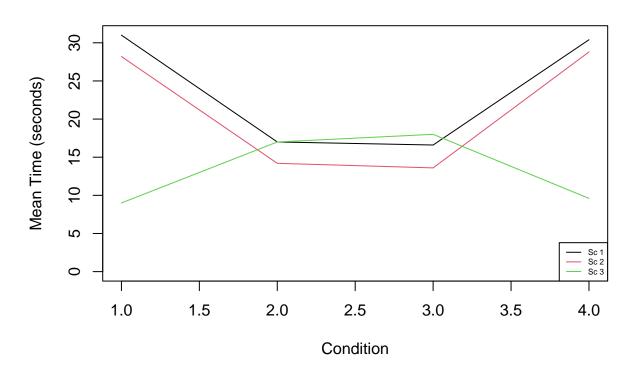
## PROBLEM 5.

```
# Define the data
data_5 = data.frame(
    Reinforcement_Schedule = rep(1:3, each = 5),
    Rat = rep(1:5, times = 3), Cond_1 = c(29, 24, 31, 41, 30, 25, 20, 35, 35, 26, 10, 9, 7, 8, 11),
    Cond_2 = c(20, 15, 19, 11, 20, 17, 12, 16, 8, 18, 18, 10, 18, 19, 20),
    Cond_3 = c(21, 10, 10, 15, 27, 19, 8, 9, 14, 18, 16, 18, 19, 20, 17),
    Cond_4 = c(18, 8, 31, 42, 53, 17, 8, 28, 40, 51, 14, 11, 12, 5, 6))

# Calculate mean time for each condition within each reinforcement schedule
mean_profiles = aggregate(. ~ data_5$Reinforcement_Schedule, data = data_5[, -c(1, 2)], FUN = mean)

# Plot mean profiles
plot(1:4, mean_profiles[1, -1], type = '1', col = 1, ylim = c(0, max(mean_profiles[-1])), xlab = "Condilines(1:4, mean_profiles[2, -1], col = 2)
lines(1:4, mean_profiles[3, -1], col = 3)
legend("bottomright", legend = c("Sc 1", "Sc 2", "Sc 3"), col = 1:3, lty = 1,cex = 0.5)
```

## **Mean Profiles for Reinforcement Schedules**



From the profiles plot, it seems that the reinforcement schedules 1,2 and 3 vary vastly among themselves across different conditions although there might be some visible similarity among the schedules 1 and 2 over different conditions.

Part (b)

```
# Reshape data into long format
# Perform one-way ANOVA
anova_res_5 = aov(formula = Time ~ Reinforcement_Schedule + Condition, data = data_5_long)
summary(anova res 5)
##
                   Df Sum Sq Mean Sq F value Pr(>F)
## Reinforcement Schedule
                    1
                       1071
                            1071.2
                                 10.382 0.00211 **
## Condition
                    1
                         0
                              0.3
                                  0.003 0.95938
```

By performing an Analysis of Variance, we can infer that the reinforcement schedule has a significant overall effect. The associated p-value is approximately 0.0021.

103.2

0 '\*\*\* 0.001 '\*\* 0.01 '\* 0.05 '.' 0.1 ' 1

57

5881

Part (c)

## ---

## Residuals

## Signif. codes:

```
# Perform profile analysis comparing schedules 1 and 2 across conditions
mod_5_12 = pbg(data = as.matrix(subset(data_5[,3:6],data_5$Reinforcement_Schedule%in%c(1,2))),group = a
summary(mod 5 12)
## Call:
## pbg(data = as.matrix(subset(data_5[, 3:6], data_5$Reinforcement_Schedule %in%
       c(1, 2))), group = as.matrix(subset(data_5[, "Reinforcement_Schedule"],
##
       data_5$Reinforcement_Schedule %in% c(1, 2))), original.names = T)
##
## Hypothesis Tests:
## $'Ho: Profiles are parallel'
    Multivariate.Test
                         Statistic
                                      Approx.F num.df den.df
                                                               p.value
## 1
                Wilks 0.995637701 0.008762825
                                                    3
                                                           6 0.9987461
## 2
               Pillai 0.004362299 0.008762825
                                                    3
                                                           6 0.9987461
## 3 Hotelling-Lawley 0.004381412 0.008762825
                                                    3
                                                           6 0.9987461
                   Roy 0.004381412 0.008762825
## 4
                                                    3
                                                           6 0.9987461
##
## $'Ho: Profiles have equal levels'
              Df Sum Sq Mean Sq F value Pr(>F)
                    16.3
                           16.26
                                 0.394 0.548
## group
                1
                8 330.3
## Residuals
##
## $'Ho: Profiles are flat'
           F df1 df2
                         p-value
## 1 7.314892
              3
                   6 0.01982651
```

On the basis of profile analysis, we can conclude at 5% level of significance on the basis of p-value that the schedules 1 and 2 behave similarly across different conditions and evidently they are parallel to each other.

```
# Perform profile analysis comparing schedules 1,2 with 3 across conditions
data_5_grouped = data_5 %>% mutate(Reinforcement_Schedule= case_when(Reinforcement_Schedule==1~1.2,
                                                                      Reinforcement_Schedule==2~1.2,
                                                                      Reinforcement_Schedule==3~3))
mod_512_3 = pbg(data = as.matrix(data_5_grouped[,3:6]),group = as.matrix(data_5_grouped$Reinforcement_S
summary(mod_512_3)
## pbg(data = as.matrix(data_5_grouped[, 3:6]), group = as.matrix(data_5_grouped$Reinforcement_Schedule
##
       original.names = T)
##
## Hypothesis Tests:
## $'Ho: Profiles are parallel'
    Multivariate. Test Statistic Approx. F num. df den. df
                                                             p.value
## 1
                 Wilks 0.2519382 10.88717
                                               3
                                                     11 0.001275056
## 2
               Pillai 0.7480618 10.88717
                                               3
                                                     11 0.001275056
## 3 Hotelling-Lawley 2.9692270 10.88717
                                               3
                                                     11 0.001275056
                   Roy 2.9692270 10.88717
                                               3
                                                     11 0.001275056
##
## $'Ho: Profiles have equal levels'
               Df Sum Sq Mean Sq F value Pr(>F)
##
                1 274.5 274.52 10.19 0.00708 **
## group
```

```
## Residuals 13 350.3 26.94
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## $'Ho: Profiles are flat'
## F df1 df2 p-value
## 1 4.154268 3 11 0.03395195
```

After grouping the schedules 1 and 2 and comparing with the schedule 3, it is evident at 5% level of significance that the profiles for the combined groups behave significantly different from the profile of the third schedule and thus are not parallel to each other.