COS-D407. Scientific Modeling and Model Validation

Hands-on excercises

Week 6

University of Helsinki, Finland

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 $Source: \ https://github.com/christina-bohk-ewald/2021-COS-D407-scientific-modeling-and-deling-an$

model-validation

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1. Some preparations in R

- 1.1 Open a new script for week 6 in R (e.g., week-6.R) and save it to a folder of your choice (e.g., course-COS-D407).
- 1.2 Create a filepath to this folder from where you would like to load data and to where you would like to save your outcome. For example,

```
the_course_COS_D407_path <- c("C:/course-COS-D407")
```

1.3 You can then set the working directory to this path

```
setwd(the_course_COS_D407_path)
```

2. Find suitable model for predicting IFR by age based on all raw data

Levin and colleagues (2020) have introduced an exponential model for predicting IFR, in %, by age. As they provide the raw data, you can test how well this exponential model and other alternative models fit these raw data.

2.1 Load raw IFR_x data of Levin et al. (2020)

Comprehensive tracing:

Please go to the preprint of Levin et al. (2020) on medRxiv and download the Supplementary Data Spreadsheets. You can rename and save this file in your course folder and load the respective data in R:

```
require(openxlsx)
## Representative sample:
levin_ifr_rs <- read.xlsx("LevinEtAl2020-spreadsheet.xlsx", sheet = 2, startRow = 1)</pre>
levin_ifr_rs[1:2,]
##
       Study AgeGroup Median_Age Population InfectionRate infrate_ci95_low infrate_ci95_high Deaths
                                       402362
                                                                                                         1 0.0
## 1 Atlanta
                  0 - 17
                               8.5
                                                          0.0
                                                                            0.0
                                                                                                1.0
## 2 Atlanta
                 18-49
                              33.5
                                       867468
                                                          3.3
                                                                            1.6
                                                                                               6.4
                                                                                                        20 0.0
##
     ifr_ci95_high
## 1
## 2
         0.1440975
## Convenience sample:
levin_ifr_cs <- read.xlsx("LevinEtAl2020-spreadsheet.xlsx", sheet = 3, startRow = 1)</pre>
levin_ifr_cs[1:2,]
##
       Study AgeGroup Median_Age Population InfectionRate infrate_ci95_low infrate_ci95_high Deaths
## 1 Belgium
                  0 - 24
                                12
                                      3228894
                                                     6.686021
                                                                       4.666667
                                                                                          9.555556
                                                                                                         1 0.0
                                35
                                      2956684
                                                                       4.666667
                                                                                                        30 0.0
## 2 Belgium
                 25 - 44
                                                    6.562703
                                                                                          9.222222
     ifr_ci95_high
## 1
            0.0007
## 2
            0.0240
```

```
levin_ifr_ct <- read.xlsx("LevinEtAl2020-spreadsheet.xlsx", sheet = 4, startRow = 1)
levin_ifr_ct[1:2,]</pre>
```

```
Study AgeGroup Median_Age Population Infections Deaths InfectionRate infrate_ci95_low infrate_
##
## 1 Australia
                   0-39
                           20.55027
                                      13533252
                                                      2800
                                                                0
                                                                     0.06206934
                                                                                       0.03103467
                  40-59
                                                                3
                                                                     0.06486020
## 2 Australia
                           50.00000
                                       6413795
                                                      2080
                                                                                       0.03891612
##
     ifr_ci95_low ifr_ci95_high
## 1
       0.0000000
                              NA
## 2
       0.04807692
                       0.1201923
```

Brief data description. The data objects $levin_ifr_rs$, $levin_ifr_cs$, and $levin_ifr_ct$ contain IFR estimates (in %) in column 7 by median age in column 2.

2.2 Plot IFR by median age

To get a feeling for these raw data, you can just plot them.

```
par(fig = c(0,1,0,1), las=1, mai=c(0.4,0.8,0.8,0.4))

plot(x=-100,y=-100,xlim=c(0,100),ylim=c(0,30),xlab="Medin age",ylab="",
    cex.main=0.9,main="Estimates of the infection fatality rate, in %,\nbased on samples
    of Levin et al. (2020)",axes=FALSE)

## names(levin_ifr_rs)

points(x=levin_ifr_rs[,"Median_Age"],y=levin_ifr_rs[,"IFR"],pch=19,col="black",lwd=2)

points(x=levin_ifr_cs[,"Median_Age"],y=levin_ifr_cs[,"IFR"],pch=19,col="blue",lwd=2)

points(x=levin_ifr_ct[,"Median_Age"],y=levin_ifr_ct[,"IFR"],pch=19,col="red",lwd=2)

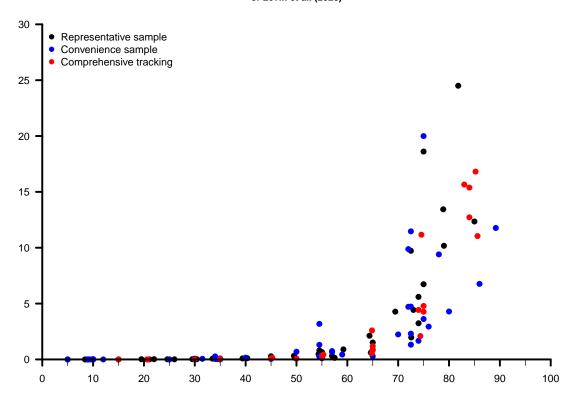
axis(side=1,at=seq(0,100,5),labels=FALSE,lwd=1,pos=0)

axis(side=1,at=seq(0,100,10),labels=TRUE,lwd=3,pos=0)

axis(side=2,at=seq(0,30,5),labels=TRUE,lwd=3,pos=0)

legend(0,30,c("Representative sample","Convenience sample","Comprehensive tracking"),
pch=rep(19,3),col=c("black","blue","red"),bty="n")
```

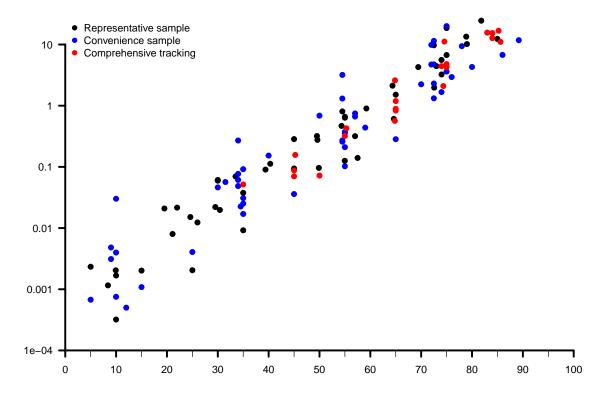
Estimates of the infection fatality rate, in %, based on samples of Levin et al. (2020)



... and on the logarithmic scale:

```
par(fig = c(0,1,0,1), las=1, mai=c(0.4,0.8,0.8,0.4))
plot(x=-100,y=-100,xlim=c(0,100),ylim=c(log(0.0001,base=exp(1)),log(30,base=exp(1))),
xlab="Medin age", ylab="", cex.main=0.9, main="Estimates of the infection fatality rate,
in %, on the log scale\nbased on samples of Levin et al. (2020)",axes=FALSE)
## names(levin_ifr_rs)
points(x=levin_ifr_rs[,"Median_Age"],y=log(levin_ifr_rs[,"IFR"],base=exp(1)),pch=19,
col="black",lwd=2)
points(x=levin_ifr_cs[,"Median_Age"],y=log(levin_ifr_cs[,"IFR"],base=exp(1)),pch=19,
col="blue",lwd=2)
points(x=levin ifr ct[,"Median Age"],y=log(levin ifr ct[,"IFR"],base=exp(1)),pch=19,
col="red", lwd=2)
axis(side=1,at=seq(0,100,5),labels=FALSE,lwd=1,pos=log(0.0001,base=exp(1)))
axis(side=1,at=seq(0,100,10),labels=TRUE,lwd=3,pos=log(0.0001,base=exp(1)))
axis(side=2,at=log(c(0.0001,0.001,0.01,0.1,1,10),base=exp(1)),
labels=c(0.0001,0.001,0.01,0.1,1,10),lwd=3,pos=0)
legend(0,log(30,base=exp(1)),c("Representative sample","Convenience sample",
"Comprehensive tracking"),pch=rep(19,3),col=c("black","blue","red"),bty="n")
```

Estimates of the infection fatality rate, in %, on the log scale based on samples of Levin et al. (2020)



2.3 Fit linear and nonlinear models to all raw data in order to predict IFR by age

In this section you can fit various models to all raw IFR_x data provided by Levin et al. (2020).

2.3.1 Build data object that contains all raw data In a first step, you can build the data objects levin_ifr and levin_age that contain all raw data (i.e., 134 data points).

2.3.2 Fit models 1 through 6 to *all* raw data and analyze their summary statistics In a second step you can fit six (or even more?) models to these raw data that differ in their complexity.

```
lm_fit <- lm(levin_ifr~levin_age)
## lm_fit
summary(lm_fit)

##
## Call:
## lm(formula = levin_ifr ~ levin_age)
##
## Residuals:
## Min  1Q Median  3Q Max
## -4.429 -2.706 -0.883  2.114 17.701</pre>
```

```
##
## Coefficients:
##
              Estimate Std. Error t value Pr(>|t|)
## (Intercept) -3.35483
                         0.69036 -4.860 3.28e-06 ***
## levin_age
               0.12412
                         0.01295
                                   9.585 < 2e-16 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## Residual standard error: 3.679 on 132 degrees of freedom
## Multiple R-squared: 0.4104, Adjusted R-squared: 0.4059
## F-statistic: 91.87 on 1 and 132 DF, p-value: < 2.2e-16
lm_fit_2 <- lm(levin_ifr~I(levin_age^2))</pre>
## lm_fit_2
summary(lm_fit_2)
##
## Call:
## lm(formula = levin_ifr ~ I(levin_age^2))
##
## Residuals:
##
      Min
               1Q Median
                              3Q
                                     Max
## -4.9030 -2.3955 -0.0029 1.6351 16.0712
##
## Coefficients:
                   Estimate Std. Error t value Pr(>|t|)
##
                 ## (Intercept)
## I(levin age^2) 0.0015348 0.0001235 12.424 < 2e-16 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 3.253 on 132 degrees of freedom
## Multiple R-squared: 0.539, Adjusted R-squared: 0.5356
## F-statistic: 154.4 on 1 and 132 DF, p-value: < 2.2e-16
lm_fit_3 <- lm(levin_ifr~I(levin_age^3))</pre>
## lm_fit_3
summary(lm_fit_3)
##
## Call:
## lm(formula = levin ifr ~ I(levin age^3))
##
## Residuals:
##
      Min
               1Q Median
                              3Q
## -5.1039 -1.7190 0.3719 1.1707 14.9290
##
## Coefficients:
                   Estimate Std. Error t value Pr(>|t|)
##
## (Intercept)
                 -1.194e+00 3.609e-01 -3.308 0.00121 **
## I(levin_age^3) 1.966e-05 1.347e-06 14.598 < 2e-16 ***
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## Residual standard error: 2.963 on 132 degrees of freedom
```

```
## Multiple R-squared: 0.6175, Adjusted R-squared: 0.6146
## F-statistic: 213.1 on 1 and 132 DF, p-value: < 2.2e-16
lm_fit_4 <- lm(levin_ifr~I(levin_age^4))</pre>
## lm_fit_4
summary(lm_fit_4)
##
## Call:
## lm(formula = levin_ifr ~ I(levin_age^4))
## Residuals:
                1Q Median
                                3Q
## -5.9851 -1.1885 0.3942 0.7453 14.1965
##
## Coefficients:
##
                    Estimate Std. Error t value Pr(>|t|)
                 -7.514e-01 3.164e-01 -2.375
## (Intercept)
                                                  0.019 *
## I(levin_age^4) 2.468e-07 1.543e-08 15.998
                                                 <2e-16 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 2.795 on 132 degrees of freedom
## Multiple R-squared: 0.6597, Adjusted R-squared: 0.6572
## F-statistic: 255.9 on 1 and 132 DF, p-value: < 2.2e-16
lm_fit_exp <- lm(log(levin_ifr,base=exp(1))~levin_age, subset=levin_ifr>0)
## lm_fit_exp
summary(lm_fit_exp)
##
## Call:
## lm(formula = log(levin_ifr, base = exp(1)) ~ levin_age, subset = levin_ifr >
##
       0)
##
## Residuals:
        Min
                  1Q
                     Median
                                    3Q
                                            Max
## -1.88568 -0.53207 0.03272 0.52574 2.65947
##
## Coefficients:
##
                Estimate Std. Error t value Pr(>|t|)
## (Intercept) -7.345377
                           0.182781 -40.19
## levin_age
               0.118387
                           0.003271
                                     36.20
                                             <2e-16 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## Residual standard error: 0.8258 on 119 degrees of freedom
## Multiple R-squared: 0.9167, Adjusted R-squared: 0.916
## F-statistic: 1310 on 1 and 119 DF, p-value: < 2.2e-16
lm_fit_exp_2 <- lm(log(levin_ifr,base=exp(1))~I(levin_age^2), subset=levin_ifr>0)
## lm_fit_exp_2
summary(lm_fit_exp_2)
##
## Call:
```

```
## lm(formula = log(levin_ifr, base = exp(1)) ~ I(levin_age^2),
##
       subset = levin ifr > 0)
##
## Residuals:
##
                 1Q
                      Median
                                   3Q
  -3.12701 -0.63996 0.09893 0.78511
                                       2.65380
##
##
## Coefficients:
##
                   Estimate Std. Error t value Pr(>|t|)
## (Intercept)
                 -5.039e+00 1.632e-01
                                        -30.87
                                                 <2e-16 ***
## I(levin_age^2)
                  1.193e-03 4.263e-05
                                         27.99
                                                  <2e-16 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 1.039 on 119 degrees of freedom
## Multiple R-squared: 0.8681, Adjusted R-squared: 0.867
## F-statistic: 783.5 on 1 and 119 DF, p-value: < 2.2e-16
```

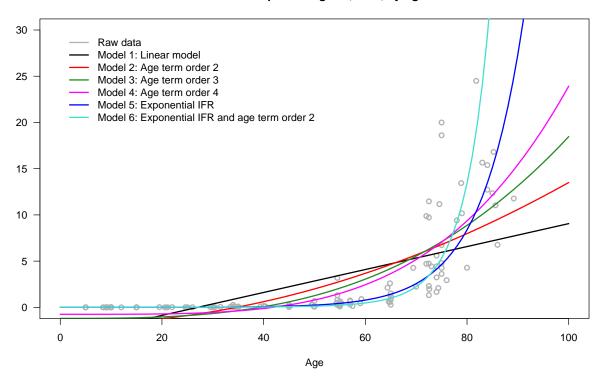
You can compare how well these models fit *all* raw data using standard diagnostic tools that R provides, e.g., via the function *summary()*. What do you think: what is the most suitable model for predicting IFR by age based on these summary statistics?

Something to think about. You could also extend this analysis and add a model of your choice here.

2.3.3 Plot how well linear and nonlinear models fit raw data of IFR by age In a third step you can visualize how well the six (or more?) models fit the raw data:

```
par(fig = c(0,1,0,1), las=1, mai=c(1.4,0.8,0.8,0.4))
plot(levin_age,levin_ifr,col=gray(0.7),xlim=c(0,100),ylim=c(0,30),lwd=2, xlab="Age",
vlab="", main="Models for predicting IFR, in %, by age")
lines(x=seq(0,100,1),y=predict(lm_fit,data.frame(levin_age=(seq(0,100,1)))),lwd=2)
lines(x=seq(0,100,1),y=predict(lm_fit_2,data.frame(levin_age=(seq(0,100,1)))),
col="red",lwd=2)
lines(x=seq(0,100,1),y=predict(lm_fit_3,data.frame(levin_age=(seq(0,100,1)))),
col="forestgreen",lwd=2)
lines(x=seq(0,100,1),y=predict(lm_fit_4,data.frame(levin_age=(seq(0,100,1)))),
col="magenta",lwd=2)
lines(x=seq(0,100,1),y=exp(predict(lm_fit_exp,data.frame(levin_age=(seq(0,100,1))))),
col="blue",lwd=2)
lines(x=seq(0,100,1),y=exp(predict(lm_fit_exp_2,data.frame(levin_age=(seq(0,100,1))))),
col="turquoise",lwd=2)
legend(0,30, c("Raw data", "Model 1: Linear model", "Model 2: Age term order 2",
"Model 3: Age term order 3", "Model 4: Age term order 4", "Model 5: Exponential IFR",
"Model 6: Exponential IFR and age term order 2"),
col=c(gray(0.7), "black", "red", "forestgreen", "magenta", "blue", "turquoise"), lwd=2, bty="n")
```

Models for predicting IFR, in %, by age



Again, based on these new information, what do you think: what is the most suitable model for predicting IFR by age?

2.3.4 Alternative metric to assess model performance: the mean squared error As an alternative to the statistics provided in the R function summary(), you can also compute the mean squared error (MSE) between the fitted and raw data:

And then you can, of course, plot the mean MSE in order to compare how well the six (or even more?) models fit *all* raw data and predict IFR by age:

```
par(fig = c(0,1,0,1), las=1, mai=c(1.4,0.8,0.8,0.4))

plot(1:6,mse_all_data,col=c("black","red","forestgreen","magenta","blue","turquoise"),
pch=19,lwd=5,xlab="Model type",ylab="",
main="Mean squared error for the six models fitted to all raw data")
lines(1:6,mse_all_data,col=gray(0.7))
legend(1,70, c("Model 1: Linear model","Model 2: Age term order 2",
"Model 3: Age term order 3","Model 4: Age term order 4",
"Model 5: Exponential IFR","Model 6: Exponential IFR and age term order 2"),
col=c("black","red","forestgreen","magenta","blue","turquoise"),pch=rep(19,6),
```

70 Model 1: Linear model Model 2: Age term order 2 Model 3: Age term order 3 60 Model 4: Age term order 4 Model 5: Exponential IFR Model 6: Exponential IFR and age term order 2 50 40 30 20 10 2 3 5 6 Model type

Mean squared error for the six models fitted to all raw data

Which of these six (or more?) models has the smallest average MSE? How does this compare to the summary statistics of the six models in section 2.3.2? Based on these new information, what do you think: what is the most suitable method to predict IFR by age?

Something more to think about. What could possibly be wrong with fitting and evaluating the six models for predicting IFR by age using *all* raw data?

3. Find suitable model for predicting IFR by age based on training data and testing data

Thinking critically about the procedure adopted in section 2, it might not be so much about finding the model that fits best to all raw data, but rather about finding the model that predicts best values of the IFR by age the model has not seen yet (machine learning; generalization of underlying pattern). That is why we redo (or refine) the analysis above and split all raw data into training data and testing data in order to find the model that predicts best IFR by age. Following this approach, we use the training data to fit the models and we use the testing data to test how well the models predict IFR by age in a new environment.

3.1 Validation set approach

According to the validation set approach, you randomly split *all* raw data into two parts of (roughly) equal size that are the training data and the testing data. You then fit the six models to the training data before you compare their IFR predictions with the actual IFR values of the testing data using the MSE.

```
train_MSE <- matrix(NA,nr=6,nc=10)</pre>
test_MSE <- matrix(NA,nr=6,nc=10)</pre>
for(run in 1:10){
    ## 3.1.1 Randomly split all data into training and testing data:
    set.seed(run)
    train <- sample(length(levin_ifr),length(levin_ifr)/2)</pre>
    test <- c(1:length(levin ifr))[-train]</pre>
    ## sort(train)
    ## sort(test)
    train_levin_ifr <- levin_ifr[train]</pre>
    train_levin_age <- levin_age[train]</pre>
    test_levin_ifr <- levin_ifr[test]</pre>
    test_levin_age <- levin_age[test]</pre>
    ## 3.1.2 Fit models to training data:
    model_1 <- lm(train_levin_ifr~train_levin_age)</pre>
    model_2 <- lm(train_levin_ifr~I(train_levin_age^2))</pre>
    model_3 <- lm(train_levin_ifr~I(train_levin_age^3))</pre>
    model_4 <- lm(train_levin_ifr~I(train_levin_age^4))</pre>
    model 5 <- lm(log(train levin ifr,base=exp(1))~train levin age,
subset=train_levin_ifr>0)
    model_6 <- lm(log(train_levin_ifr,base=exp(1))~I(train_levin_age^2),</pre>
subset=train_levin_ifr>0)
    ## 3.1.3 Calculate and save train MSE:
    train_MSE[1,run] <- mean((train_levin_ifr -</pre>
predict(model_1,data.frame(train_levin_age=train_levin_age)))^2)
    train_MSE[2,run] <- mean((train_levin_ifr -</pre>
predict(model_2,data.frame(train_levin_age=train_levin_age)))^2)
    train_MSE[3,run] <- mean((train_levin_ifr -</pre>
predict(model_3,data.frame(train_levin_age=train_levin_age)))^2)
    train_MSE[4,run] <- mean((train_levin_ifr -</pre>
predict(model_4,data.frame(train_levin_age=train_levin_age)))^2)
    train_MSE[5,run] <- mean((train_levin_ifr -</pre>
exp(predict(model_5,data.frame(train_levin_age=train_levin_age))))^2)
    train_MSE[6,run] <- mean((train_levin_ifr -</pre>
exp(predict(model_6,data.frame(train_levin_age=train_levin_age))))^2)
    ## 3.1.4 Apply fitted models to testing data and calculate and save test MSE:
    test_MSE[1,run] <- mean((test_levin_ifr -</pre>
```

```
predict(model_1,data.frame(train_levin_age=test_levin_age)))^2)

   test_MSE[2,run] <- mean((test_levin_ifr -
predict(model_2,data.frame(train_levin_age=test_levin_age)))^2)

   test_MSE[3,run] <- mean((test_levin_ifr -
predict(model_3,data.frame(train_levin_age=test_levin_age)))^2)

   test_MSE[4,run] <- mean((test_levin_ifr -
predict(model_4,data.frame(train_levin_age=test_levin_age)))^2)

   test_MSE[5,run] <- mean((test_levin_ifr -
exp(predict(model_5,data.frame(train_levin_age=test_levin_age))))^2)

   test_MSE[6,run] <- mean((test_levin_ifr -
exp(predict(model_6,data.frame(train_levin_age=test_levin_age))))^2)

} ## for run</pre>
```

The R-code above applies the validation set approach ten times (run 1 through ten in the for loop). The data objects *train_MSE* and *test_MSE* contain the MSE for each of the six models (rows) in each of the 10 runs (columns) when applied to the training data and the testing data, respectively.

These data allow you to eventually compute the average train MSE and the average test MSE across all ten runs for each of the six models:

```
rowMeans(train_MSE)
```

```
## [1] 12.955992 9.899420 8.036573 7.022064 8.565164 55.786678
rowMeans(test_MSE)
```

```
## [1] 14.003603 11.279971 9.657318 8.882602 11.759862 149.008199
```

Which model has the smallest / largest testing MSE? Which model has the smallest / largest training MSE? You can also plot your findings:

```
par(fig = c(0,1,0,1), las=1, mai=c(1.4,0.8,0.8,0.4))

plot(1:6,rowMeans(train_MSE),main="MSE based on validation set approach",
    col=c("black","red","forestgreen","magenta","blue","turquoise"),
    pch=19,cex=2,xlab="Model type",ylab="",ylim=c(0,150))

points(1:6,rowMeans(test_MSE),
    col=c("black","red","forestgreen","magenta","blue","turquoise"),pch=15,cex=2)

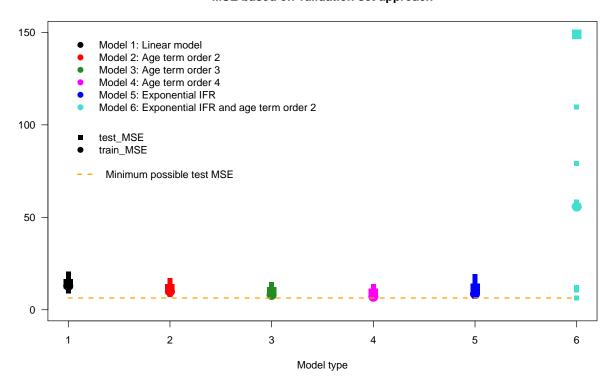
for(model in 1:6){
        points(x=rep(model,10),y=test_MSE[model,],
        col=c("black","red","forestgreen","magenta","blue","turquoise")[model],pch=15,cex=1)
}

lines(1:6,rep(min(test_MSE),6),col="orange",lty=2,lwd=2)

legend(1,150, c("Model 1: Linear model","Model 2: Age term order 2",
        "Model 3: Age term order 3","Model 4: Age term order 4",
        "Model 5: Exponential IFR","Model 6: Exponential IFR and age term order 2"),
        col=c("black","red","forestgreen","magenta","blue","turquoise"),
```

```
pch=rep(19,6),lty=NA,lwd=2,bty="n")
legend(1,100,col=c("black","black"),c("test_MSE","train_MSE"),
lty=NA,pch=c(15,19),lwd=2,bty="n")
legend(1,80,col=c("orange"),c("Minimum possible test MSE"),lty=c(2),lwd=2,bty="n")
```

MSE based on validation set approach



Something to think about. What do these information tell you about the suitability of each of these models for predicting the IFR by age? To answer this question it might be convenient to zoom into the plot above. What model(s) have comparably low bias and low variance?

Something more to think about. Why can low bias and low variance be regarded as indicators for a suitable model?

4. Time for you to think both creatively and critically about selecting the most suitable method for predicting IFR by age based on raw data provided by Levin and colleagues (2020).

You can consider here the pros and cons of the different procedures adopted to fit and evaluate the models with respect to their ability to accurately predict IFR by age. For example, what is the actual purpose of a model that has been designed to predict IFR by age? What does it mean when a model is fitted to training data instead of to *all* raw data? And what does it mean when a model's predictions are evaluated using testing data instead of *all* raw data?

When selecting the most suitable model (among those six models) for predicting the IFR by age, you may

| also want to consider t | the theoretical meaning | g of the models; e.g. | ., to distinguish betw | een M4 and M5? |
|-------------------------|-------------------------|-----------------------|------------------------|----------------|
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