COS-D407. Scientific Modeling and Model Validation

Hands-on excercises

Week 4

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 4 in R (e.g., week-4.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. Download and load required input data

In week 4 we apply the demographic scaling model of Bohk-Ewald et al. (2020) for estimating total numbers of COVID-19 infections in Finland. We will start with downloading and loading required input data in R, continue with reading basic functions of the demographic scaling model in R, and finally applying it to estimate the number of people who probably are and have been infected with COVID-19 so far.

2.1 Download confirmed cases and reported deaths attributable to COVID-19 as of today

Please go to the website of the Johns Hopkins University CSSE. The files

- \bullet time_series_covid19_confirmed_global.csv
- \bullet time_series_covid19_deaths_global.csv

contain confirmed cases and reported deaths, respectively, for many countries on a daily basis since January 22, 2020. Please download these two files and save them in your project folder.

2.2 Load COVID-19 data

Please load the numbers of confirmed cases and reported deaths from COVID-19 in R using the function read.csv of the R-package openxlsx.

```
require(openxlsx)
confirmed <- read.csv("time_series_covid19_confirmed_global.csv",header=TRUE,</pre>
stringsAsFactors = FALSE)
confirmed[1:2,1:6]
##
     Province.State Country.Region
                                          Lat
                                                   Long X1.22.20 X1.23.20
## 1
                        Afghanistan 33.93911 67.70995
                                                                         0
                                                                         0
## 2
                            Albania 41.15330 20.16830
deaths <- read.csv("time_series_covid19_deaths_global.csv",header=TRUE,</pre>
stringsAsFactors = FALSE)
deaths[1:2,((ncol(deaths)-5):ncol(deaths))]
     X9.10.20 X9.11.20 X9.12.20 X9.13.20 X9.14.20 X9.15.20
## 1
         1420
                   1420
                            1420
                                      1420
                                               1425
                                                         1426
## 2
          324
                    327
                             330
                                       334
                                                 338
                                                          340
```

2.3 Download and load abridged life tables

Please go to the UNWPP2019 website, download abridged life tables for both sexes together, save them into your project folder, and then load them into R.

```
lt_1950_2020 <- read.xlsx("WPP2019_MORT_F17_1_ABRIDGED_LIFE_TABLE_BOTH_SEXES.xlsx",
sheet = 1,startRow = 17)</pre>
```

Brief data description. The data object lt_1950_2020 contains a bridged life tables for both sexes for all UN countries.

Explore this data object and find out how large Finnish remaining life expectancy at birth has been 1950-55 through 2015-19.

```
## lt_1950_2020[1:2,]
colnames(lt_1950_2020)
    [1] "Index"
##
##
       "Variant"
    [2]
##
    [3] "Region, .subregion, .country.or.area.*"
    [4] "Notes"
##
##
    [5]
        "Country.code"
##
    [6] "Type"
##
    [7] "Parent.code"
   [8] "Period"
##
##
    [9] "Age.(x)"
## [10] "Age.interval.(n)"
## [11] "Central.death.rate.m(x,n)"
## [12] "Probability.of.dying.q(x,n)"
## [13] "Probability.of.surviving.p(x,n)"
## [14] "Number.of.survivors.l(x)"
## [15] "Number.of.deaths.d(x,n)"
  [16] "Number.of.person-years.lived.L(x,n)"
  [17] "Survival.ratio.S(x,n)"
## [18] "Person-years.lived.T(x)"
## [19] "Expectation.of.life.e(x)"
## [20] "Average.number.of.years.lived.a(x,n)"
lt_1950_2020[which(lt_1950_2020[,"Region,.subregion,.country.or.area.*"] == "Finland" &
  lt_1950_2020["Age.(x)"]==0),c("Period","Expectation.of.life.e(x)")]
```

```
##
            Period Expectation.of.life.e(x)
## 4823 1950-1955
                         66.401073999999994
## 10349 1955-1960
                         68.189361000000005
## 15875 1960-1965
                         69.067266000000004
## 21401 1965-1970
                         69.71624799999993
## 26927 1970-1975
                         70.930030000000002
## 32453 1975-1980
                         72.714505000000003
## 37979 1980-1985
                         74.326365999999993
## 43505 1985-1990
                         74.788223000000002
## 49031 1990-1995
                         75.84142799999993
                         77.144569000000004
## 54557 1995-2000
## 60083 2000-2005
                         78.400789000000003
## 65609 2005-2010
                         79.548135000000002
## 71135 2010-2015
                         80.705350999999993
## 76661 2015-2020
                         81.646075999999994
```

2.4 Load global pattern over age of COVID-19 deaths

Dudel et al. (2020) provide in their supplementary material data for age-specific death counts attributable to COVID-19, which has been served as a basis for calculating a global average pattern over age for total death counts as input for the demographic scaling model. You can download this *global average pattern over age* from the GitHub repository for this course.

```
global_age_dist_deaths <- source("global_age_dist_deaths.R")
## global_age_dist_deaths</pre>
```

Brief data description. The data object *global_age_dist_deaths* contains the global pattern over 10-year age groups of COVID-19 deaths.

Note that we follow here the original methodology of the demographic scaling model that has been introduced in the paper of Bohk-Ewald et al. (2020). Another way is to use COVID-19-related death counts by age that have been reported to the COVerAGE database (as presented on Monday).

2.5 Load infection fatality rates from Verity et al. (2020)

Verity and colleagues (2020, page 5) report infection fatality rates by 10-year age groups for Hubei, China, on page 5. Please create a data object *ifr_by_age_china_verity* that contains these data or download it from the GitHub repository for this course.

```
ifr_by_age_china_verity <- read.table("infection-fatality-rates-by-age-china-Verity.txt",
header=FALSE, stringsAsFactors = FALSE)
ifr_by_age_china_verity</pre>
```

```
## V1 V2 V3 V4
## 1 0 1.6e-05 1.85e-06 0.000249
## 2 10 7.0e-05 1.50e-05 0.000500
## 3 20 3.1e-04 1.40e-04 0.000920
## 4 30 8.4e-04 4.10e-04 0.001850
## 5 40 1.6e-03 7.60e-04 0.003200
## 6 50 6.0e-03 3.40e-03 0.013000
## 7 60 1.9e-02 1.10e-02 0.039000
## 8 70 4.3e-02 2.50e-02 0.084000
## 9 80 7.8e-02 3.80e-02 0.133000
```

Brief data description. The data object *ifr_by_age_china_verity* contains the modal estimate as well as the lower and upper bound of the 95 percent credible interval for the infection fatality rates of Hubei, China, by 10-year age groups.

3. Estimate COVID-19 infections in Finland

3.1 Source basic functions of the demographic scaling model

You can find the basic functions of the demographic scaling model in the file basic-functions-week-4.R in the GitHub repository for this course. They contain the functions:

- to ungroup to interpolate IFR estimates of Verity et al. (2020) into single years of age
- get_ungrouped_ex_2015_2020 to ungroup remaining life expectancy
- map_ifr_betw_ref_and_one_coi_thanatAge to scale IFRs from a reference country (here: China; Verity et al. (2020)) onto a country of interest based on remaining life expectancy
- $aggregate_mapped_ifr_10y$ to aggregate scaled IFRs into 10-year age groups
- disaggregate_deaths_one_coi_10y to disaggregate total deaths into 10-year age groups

You may have a look at these basic functions if you wish. But it is also fine to just source (or load) them via the file basic-functions-week-4.R:

3.2 Apply the demographic scaling model with data for Finland

```
#
## 1. Ungroup China's IFR:
ungrouped_mode_ifr_by_single_age_china_sp <- to_ungroup(to_ungroup=
                ifr_by_age_china_verity[,2],nr_grouped_years=10)
ungrouped_low95_ifr_by_single_age_china_sp <- to_ungroup(to_ungroup=
                ifr_by_age_china_verity[,3],nr_grouped_years=10)
ungrouped_up95_ifr_by_single_age_china_sp <- to_ungroup(to_ungroup=
                ifr_by_age_china_verity[,4],nr_grouped_years=10)
## 2. Scale IFRs from a RC onto a COI via remaining life expectancy:
mapped_mode_ifr_thanatAge <- map_ifr_betw_ref_and_one_coi_thanatAge(coi="Finland",</pre>
    lt_1950_2020=lt_1950_2020,
    ungrouped_ifr_by_single_age_china_sp=ungrouped_mode_ifr_by_single_age_china_sp)
## and fill in the few NA values:
pos_na <- which(is.na(mapped_mode_ifr_thanatAge[1,]))</pre>
    if(length(pos_na)>0){
        for(pos in 1:length(pos_na)){
            if(pos_na[pos] < 6){</pre>
                mapped_mode_ifr_thanatAge[1,pos_na[pos]] <-</pre>
                    min(mapped mode ifr thanatAge[1,],na.rm=TRUE)
            if(pos_na[pos] >= 6){
                mapped_mode_ifr_thanatAge[1,pos_na[pos]] <-</pre>
                    mapped_mode_ifr_thanatAge[1,pos_na[pos]-1]
        } ## for pos
    } ## if
#
## 3. Put scaled IFRs into 10-year age groups:
mapped_mode_ifr_thanatAge_10y <- aggregate_mapped_ifr_10y(disaggregated_mapped_ifr=</pre>
                         mapped_mode_ifr_thanatAge)
## 4. Disaggregate total COVID-19-related deaths into 10-year age groups:
deaths_by_age <- disaggregate_deaths_one_coi_10y(coi="Finland")</pre>
```

```
#
## 5. Estimate COVID-19 infections over time:
#
inf_mode <- colSums( deaths_by_age / mapped_mode_ifr_thanatAge_10y )</pre>
```

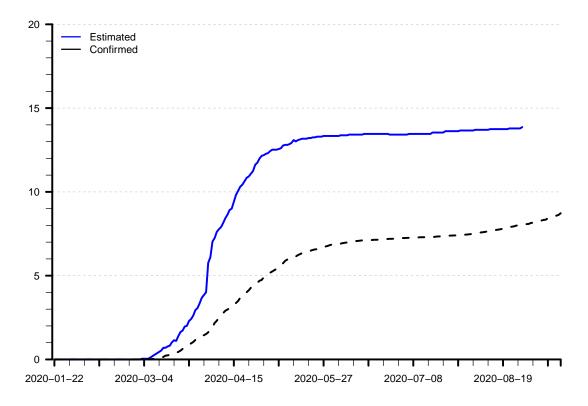
Think about how you could extend this R-code in order to estimate the total numbers of COVID-19 infections for other countries and how to provide uncertainty estimates for them.

3.3 Visualize COVID-19 infecton estimates for Finland

Visualize the numbers of COVID-19 infections in Finland over time, accounting for an average time to death of 18 days.

```
dates <- seq(as.Date("22/01/2020", format = "%d/%m/%Y"),
by = "days", length = (ncol(deaths)-4) )
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,length(5:ncol(deaths))),ylim=c(0,20),xlab="Date",ylab="",cex.main=0.9,
main="Total numbers of COVID-19 infections, in thousand, in Finland", axes=FALSE)
segments(x0=rep(0,4),x1=rep(length(5:ncol(deaths)),4),y0=seq(5,20,5),y1=seq(5,20,5),
lty=2,col=grey(0.8))
lines(x=1:length(5:ncol(deaths)), y=c(inf_mode[-c(1:18)], rep(NA, 18))/1000, col="blue", lwd=3)
lines(x=1:length(5:ncol(deaths)),y=confirmed[which(confirmed[,"Country.Region"]=="Finland"),
5:ncol(confirmed)]/1000,col="black",lty=2,lwd=3)
legend(0,20,c("Estimated","Confirmed"),col=c("blue","black"),bty="n",lwd=2,lty=1)
axis(side=1,at=seq(1,length(5:ncol(deaths)),7),labels=FALSE,lwd=1,pos=0)
axis(side=1,at=c(seq(1,length(5:ncol(deaths)),21),length(5:ncol(deaths))),
labels=dates[c(seq(1,length(5:ncol(deaths)),21),
length(5:ncol(deaths)))],lwd=3,pos=0)
axis(side=2,at=seq(0,20,1),labels=FALSE,lwd=1,pos=0)
axis(side=2,at=seq(0,20,5),labels=TRUE,lwd=3,pos=0)
```

Total numbers of COVID-19 infections, in thousand, in Finland



Please describe the level and the temporal development of the estimated numbers of COVID-19 infections in Finland, also compared to (1) the numbers of confirmed cases in Finland, and to (2) the corresponding figures in other European countries. As a source of inspiration, you may want to have a look at the press release *How many Finns have really been infected with COVID-19?*.

Again, please think about how you could extend this R-code in order to estimate the total numbers of COVID-19 infections for other countries and how to provide uncertainty estimates for them.

4. Time for you to think both creatively and critically about these COVID-19 infection estimates for Finland.

How would you evaluate the process of the demographic scaling model?

How plausible are the infection estimates, also considering, e.g., the quality of the input data and the key assumptions of the demographic scaling model? As a source of inspiration, you may want to have a look at the paper of Bohk-Ewald et al. (2020).