Introduction to Handling Data

ECON20222 - Lecture 2

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Aim for today

- Become familiar with handling some Covid-19 data
- Produce some graphical representation of data
- Become familiar with merging datasets
- Review hypothesis testing
- Review simple regression analysis
- Understand the limitation of regression as a causal analysis tool
- Become more familiar with R

Preparing your workfile

We add the basic libraries needed for this week's work:

```
#library(sets)
                       # used for some set operations
library(forecast)
                      # used for some data smoothing
library(readxl)
                      # enable the read excel function
library(tidyverse)
                      # for almost all data handling tasks
library(ggplot2)
                      # plotting toolbox
library(utils)
                      # for reading data into R
library(httr)
                      # for downloading data from a URL
library(stargazer)
                      # enables well formatted regression out;
```

New Dataset - Covid

The CORE-ECON Covid-19 Collection contains a more detailed version of this example.

The dataset is published by the European Centre for Disease Control (ECDC)

- Weekly data for Covid cases and deaths
- more than 200 countries

```
#download the dataset from the ECDC website to a local temporary file ("tf")
GET("https://opendata.ecdc.europa.eu/covid19/nationalcasedeath/csv",
    authenticate(":", ":", type="ntlm"),
    write_disk(tf <- tempfile(fileext = ".csv")))
#read the Dataset sheet into "R". The dataset will be called "data".
data_cov <- read_csv(tf,col_types = "fffnfncnnc") # coltypes presets the types</pre>
```

- This will load data into your environment.
- We are tapping directly into the ECDC's datafile. Everytime you do this you will get the most recent data (on 3 Feb 2022 this delivered 44002 observations)

Covid Data - Explore

After some name changes and turning dates into date format:

```
str(data_cov) # prints some basic info on variables
## tibble[,16] [21,357 x 16] (S3: tbl_df/tbl/data.frame)
   $ country
                              : Factor w/ 224 levels "Afghanistan",..: 1 1 1 1 1 1
##
##
   $ country_code
                              : Factor w/ 218 levels "AFG", "ALB", "DZA", ...: 1 1 1 1
   $ continent
                              : Factor w/ 5 levels "Asia". "Africa"...: 1 1 1 1 1 1
##
   $ popData2019
                              : num [1:21357] 38928341 38928341 38928341 38928341 3
##
##
   $ year
                              : chr [1:21357] "2020" "2020" "2020" "2020" ...
   $ week
                              : chr [1:21357] "01" "02" "03" "04" ...
##
##
   $ cases weekly
                              : num [1:21357] 0 0 0 0 0 0 0 1 3 ...
                              : num [1:21357] 0 0 0 0 0 0 0 0 0 0 ...
##
   $ deaths weekly
   $ cases_weekly_cumulative : num [1:21357] 0 0 0 0 0 0 0 1 4 ...
##
   $ deaths_weekly_cumulative: num [1:21357] 0 0 0 0 0 0 0 0 0 0 ...
##
   $ cases_14_day
                              : num [1:21357] NA 0 0 0 0 ...
##
##
   $ deaths_14_day
                              : num [1:21357] NA 0 0 0 0 0 0 0 0 0 ...
##
   $ source cases
                              : chr [1:21357] "Epidemic intelligence national data"
##
   $ source_deaths
                              : chr [1:21357] "Epidemic intelligence national data"
##
   $ dates
                              : Date[1:21357], format: "2020-01-02" "2020-01-09" ...
##
    $ geoID
                              : chr [1:21357] "AF" "AF" "AF" "AF" ...
```

Covid Data - Explore

Let's find out what one observation represents.

```
data_cov[678,] # 678 is just an arbitrary row in the dataset
## # A tibble: 1 x 16
##
     country
                        country code continent popData2019 year week cases weekly
##
     <fct>
                        <fct>
                                     <fct>
                                                     <dbl> <chr> <chr>
                                                                              <dbl>
    Antigua And Barbu~ ATG
                                     America
                                                     97928 2021 18
     ... with 9 more variables: deaths weekly <dbl>,
## #
       cases_weekly_cumulative <dbl>, deaths_weekly_cumulative <dbl>,
       cases_14_day <dbl>, deaths_14_day <dbl>, source_cases <chr>,
## #
       source deaths <chr>, dates <date>, geoID <chr>
## #
```

Covid Data - Explore

Let's find out how many observations we have for a set of countries. Say for China, the UK ("United_Kingdom") and the Bahamas.

Use piping technique of the tidyverse

```
sel_countries <- c("China", "United Kingdom", "Bahamas")

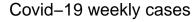
# select only countries in sel_countries
table1 <- data_cov %>% filter(country %in% sel_countries) %>%
    group_by(country) %>% # groups by Wave and Country
    summarise(n = n()) %>% # calculating number of obs
    print()
```

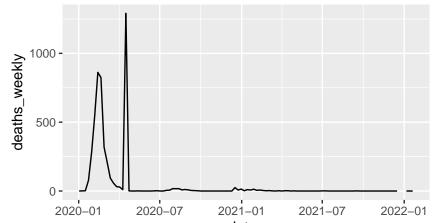
```
## # A tibble: 3 x 2
## country n
## <fct> <int>
## 1 Bahamas 97
## 2 China 107
## 3 United Kingdom 107
```

Smaller countries tend to start reporting later than bigger ones.

Data - Some graphical representation

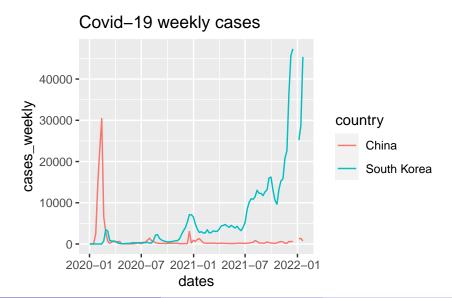
Essentially we have a dataset which combines cross-section (different countries) with time-series (consequitive weeks for each country). We call such a dataset a panel.





Data - Some graphical representation

Weekly cases for two countries



Data - Some summary stats

Summarise data by country.

- cases_weekly: average number of weekly cases across all weeks
- deaths_weekly: average number of weekly deaths across all weeks

```
## # A tibble: 6 x 3
##
    country Avg wc Avg wd
    <fct>
               <dbl> <dbl>
##
## 1 Afghanistan 1502. 69.5
## 2 Albania
            2505. 33.5
## 3 Algeria 2147. 60.2
## 4 American Samoa 1
                      0
               308.
                      1.47
## 5 Andorra
                       19.5
## 6 Angola
                830
```

Table 2 now contains an observation for each country with the average of weekly cases and deaths. Why is there such a stark difference between Algeria and Andorra?

Data - When can you compare data

```
## # A tibble: 6 x 3
##
    country
          Avg_wc Avg_wd
    <fct>
          <dbl> <dbl>
## 1 Afghanistan 1502. 69.5
## 2 Albania
             2505. 33.5
           2147. 60.2
  3 Algeria
## 4 American Samoa 1 0
## 5 Andorra
             308.
                     1.47
## 6 Angola
               830
                       19.5
```

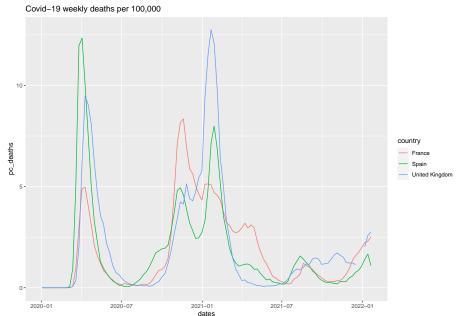
Comparing absolute numbers of any variable can be extremely misleading.

- Adjusting by population is often appropriate (other standardisation variables may be appropriate in other situations)
- Some variables may be the result of different testing strategies in countries (affects weekly_cases more than weekly_deaths)

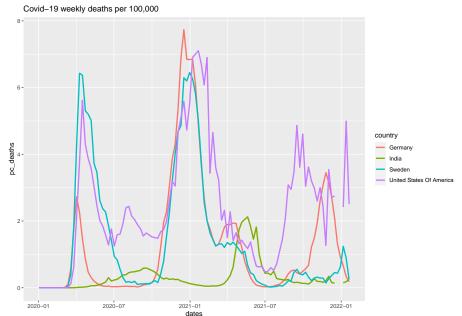
Data - Standardise the case data

Let's standardise by population (popData2019) which is included as a variable into the dataset. Note that this is constant through the weeks. These are often reported as cases per 100,000 (although for Deaths sometimes per 1,000,000 - see Our World in Data).

Data - Standardise the case data



Data - Standardise the case data



Data - Importing and Merging Data

Is it the case that countries with larger population density find it more difficult to control the spread of Covid?

Need to import land area data and merge them into our dataset (CountryIndicators.csv). This also imports two further country indicators (Health Expenditure and GDP per capita).

```
## # A tibble: 2 x 4

## country Avg_cases Avg_deaths PopDen

## <fct> <dbl> <dbl> <dbl> <dbl> 59.6

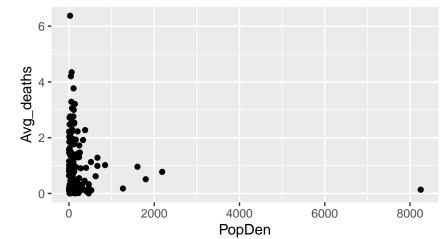
## 2 Albania 88.0 1.18 104.
```

table3 now includes a column for the population density along the average weekly case and deaths (per capita).

Data - Scatter Plot

```
ggplot(table3,aes(PopDen,Avg_deaths)) +
geom_point() +
ggtitle("Population Density v Per Capita Deaths")
```

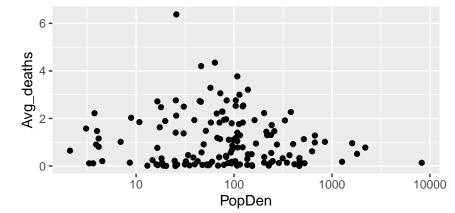
Population Density v Per Capita Deaths



Data - Scatter Plot

```
ggplot(table3,aes(PopDen,Avg_deaths)) +
  geom_point() +
  scale_x_log10() +
  ggtitle("Population Density v Per Capita Deaths")
```

Population Density v Per Capita Deaths



Data - Correlation

An important statistic which is used to measure the strength of a relationship is the correlation coefficient.

Is there a relationship between PopDen and Avg_deaths?

$$Corr_{PopDen,Avg_deaths} = \frac{Cov(PopDen,Avg_deaths)}{s_{PopDen} \ s_{Avg_deaths}}$$

Correlations are in the [-1,1] interval. They are standardised covariances. Ensure you revise how to calculate sample s.d. and covariances! R does it using the cor function.

[1] -0.1087693

So if at all, there is a negative relationship but close to 0.

Data on Maps

Geographical relationships are sometimes best illustrated with maps.

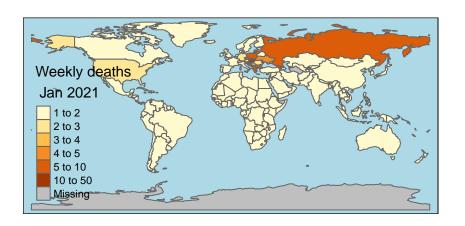
Sometimes these will reveal patterns which are not very obvious in other ways..

R can create great maps (but it requires a bit of setup - see the additional file on BB). You need the following

- A shape file for each country
- The statistics for each country, like Avg_deaths
- a procedure to merge these bits of information in one data-frame (merge)

Let's look at the distribution of weekly deaths across the globe as of Jan 2021.

Data on Maps



Hypothesis Testing - Introduction

Hypothesis testing is a core technique used in empirical analysis. Use sample data to infer something about the population mean (or correlation, or variance, etc). Hence inference.

It is crucial to understand that the particular sample we have is one of many different possible samples. Whatever conclusion we arrive at is not characterised by certainty.

Example

Is the average number of (per capita) weekly cases in Europe the same as that in America (as per 1 Feb 2021).

$$H_0$$
: $\mu_{c,EU,1Feb21} = \mu_{c,AM,1Feb21}$
 H_A : $\mu_{c,EU,1Feb21} \neq \mu_{c,AM,1Feb21}$

The truth is either represented by H_0 or H_A .

Here c represents the variable pc_cases,

When performing a test we need to calibrate some level of uncertainty. We typically fix the Probability with which we reject a correct null hypothesis (Type I error). This

Introduction to Handling Data

The data

Let's first look at the averages across the continents (continentExp)

```
## # A tibble: 5 x 4
##
   continent Avg_cases Avg_deaths
   <fct>
              <dbl> <dbl> <int>
##
             54.6 0.465
## 1 Asia
                              42
  2 Africa
             19.1 0.403
                              55
  3 Europe 164. 3.58
                              55
## 4 America
             83.5 1.53
                              49
## 5 Oceania
              7.64 0.0932 14
```

Hypothesis Testing - Introduction

Depending on the type of hypothesis there will be a test statistic which will be used to come to a decision.

Assuming that H_0 is true this test statistic has a random distribution (frequently t, N, χ^2 or F). We can then use this distribution to evaluate how likely it would have been to get the type of sample we have if the null hypothesis was true () or obtain

Decision Rule 1: If that probability is smaller than our pre-specified significance level, then we H_0 . If, however, that p-value is larger than our pre-specified significance level then we will H_0 .

Decision Rule 2: If the absolute value of the test statistic is larger than the critical value (obtain from the Null distribution - see next slide), then we H_0 . If, however, the absolute value of the test statistic is smaller than the critical value, then we will H_0 .

Hypothesis Testing - Introduction

Example The test statistic

$$t = \frac{\bar{c}_{EU,1Feb21} - \bar{c}_{AM,1Feb21}}{\sqrt{\frac{s_{c,EU,1Feb21}^2}{n_{EU,1Feb21}} + \frac{s_{c,AM,1Feb21}^2}{n_{AM,1Feb21}}}}$$

How is this test statistic, t, distributed (assuming H_0 is true)? **If**

- 1 The two samples are independent
- ② The random variables $c_{EU,1Feb21}$ and $c_{AM,1Feb21}$ are either normally distributed or we have sufficiently large samples
- **3** The variances in the two samples are identical

then $t \sim$

The above assumptions are crucial (and they differ from test to test). If they are not met then the resulting p-value (or critical values) are not correct.

Hypothesis Testing - Example 1

Let's create a sample statistic:

```
test data EU <- data cov %>%
  filter(continent == "Europe") %>%  # pick European data
  filter(dates == "2021-02-04")  # pick the date
mean EU <- mean(test data EU$pc cases.rm.na = TRUE)
test data AM <- data cov %>%
  filter(continent == "America") %>% # pick European data
  filter(dates == "2021-02-04")  # pick the date
mean_AM <- mean(test_data_AM$pc_cases,rm.na = TRUE)</pre>
sample_diff <- mean_EU - mean_AM</pre>
paste("mean_EU =", round(mean_EU,1),", mean_A =", round(mean_AM,1))
## [1] "mean_EU = 163.6 , mean A = 83.5"
paste("sample diff =", round(sample diff,1))
## [1] "sample_diff = 80.1"
```

Is this difference

significant?

Hypothesis Testing - Example 1

Formulate a null hypothesis. Here that the difference in population means (mu) is equal to 0 using the t.test function. We deliver the pc_cases series for both countries to the t.test function.

```
t.test(test_data_EU$pc_cases,test_data_AM$pc_cases, mu=0) # testing that mu = 0

##

## Welch Two Sample t-test

##

## data: test_data_EU$pc_cases and test_data_AM$pc_cases

## t = 3.2867, df = 95.202, p-value = 0.00142

## alternative hypothesis: true difference in means is not equal to 0

## 95 percent confidence interval:

## 31.73067 128.52786

## sample estimates:

## mean of x mean of y

## 163.5844 83.4551
```

The p-value is very small and hence

Hypothesis Testing - Example 2

What about the difference between Asia and Africa though?

```
t.test(test_data_AF$pc_cases,test_data_AS$pc_cases, mu=0)  # testing that mu = 0

##

## Welch Two Sample t-test

##

## data: test_data_AF$pc_cases and test_data_AS$pc_cases

## t = -1.9057, df = 50.669, p-value = 0.06237

## alternative hypothesis: true difference in means is not equal to 0

## 95 percent confidence interval:

## -72.919547  1.903617

## sample estimates:

## mean of x mean of y
```

The p-value is 0.06237 and hence

19.07221 54.58018

##

Hypothesis Testing - To reject or to not reject

When comparing between Europe and America the p-value was smaller than 0.01:

When comparing between Africa and Asia the p-value was 0.062:

- Conventional significance levels are 10%, 5%, 1% or 0.1%
- But what do they mean?

Regression Analysis - Introduction

Tool on which most of the work in this unit is based

- Allows to quantify relationships between 2 or more variables
- It can be used to implement hypothesis tests
- However it does not necessarily deliver causal relationships!

It is very easy to compute for everyone! Results will often have to be interpretated very carefully.

Your skill will be to interpret correctly!!!!

Regression Analysis - Data Preparation

Create new dataset which contains for every country:

- the average per capita deaths throughout the sample, Avg_deaths,
- the continent (continentExp),
- the population density data (PopDen).
- the GDP per capita (GDPpc,2018, in US\$1,000), from the World Health Organisation, Global Health Expenditure Database
- Current Health Expenditure (HealthExp) as % GDP, 2018, from the World Health Organisation, Global Health Expenditure Database

table3 already contains pc_deaths and PopDen. We need to merge in the other info from data.

```
mergecont <- data_cov %>% dplyr::select(country,continent, GDPpc, HealthExp) %>% unique() %>% # this reduces each country to one line drop_na # this drops all countries which have incomplete inform table3 <- merge(table3,mergecont) # merges in continent information table3 <- table3 %>% mutate(GDPpc = GDPpc/1000) # convert pc GDP into units of $1,000
```

Now we run a regresison of the average pc_deaths (Avg_deaths in table3) against a constant only. Recall, one observation here is one country.

$$Avg_deaths_i = \alpha + u_i$$

We use the stargazer function to display regression results

```
##
##
                            Dependent variable:
##
                                Avg_deaths
##
                                 1.078***
  Constant
                                  (0.093)
##
##
  Observations
                                    143
                                   0.000
## Adjusted R2
                                   0.000
## Residual Std. Error 1.110 (df = 142)
## Note:
                        *p<0.1; **p<0.05; ***p<0.01
```

The estimate for the constant, $\hat{\alpha}$, is the sample mean. So on average, countries had an average rate of deaths of just under 1 per 100,000 per week due to Covid-19. (Note that in this average all countries have the same weight).

Testing $H_0: \mu_{Avg_deaths} = 0$ can be achieved by

```
##
## One Sample t-test
##
## data: table3$Avg_deaths
## t = 11.609, df = 142, p-value < 2.2e-16
## alternative hypothesis: true mean is not equal to 0
## 95 percent confidence interval:
## 0.8944023 1.2615262
## sample estimates:
## mean of x
## 1.077964</pre>
```

We can use the above regression to achieve the same:

$$t - test = \widehat{\alpha}/se_{\widehat{\alpha}}$$

We now estimate a regression model which also includes the GDP per capita (GDPpc) as an explanatory variable.

$$Avg_deaths_i = \alpha + \beta \ GDPpc_i + u_i$$

mod2 <- lm(Avg_deaths~GDPpc,data=table3)</pre>

How do we interprete the estimate of $\widehat{\beta}$?

What sign do you expect it to have?

```
stargazer(mod2, type="text")
```

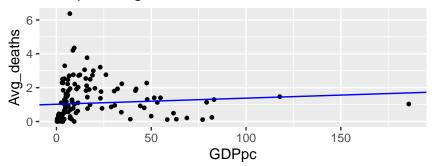
```
Dependent variable:
                             Avg_deaths
## GDPpc
                               0.004
                               (0.004)
                              1.019***
## Constant
                               (0.111)
## Observations
                                143
## R2
                                0.007
## Adjusted R2 -0.0002
## Residual Std. Error 1.111 (df = 141)
## F Statistic
                         0.975 (df = 1; 141)
                     *p<0.1: **p<0.05: ***p<0.01
## Note:
```

As the income increases by one unit (e.g. from \$1,000 to \$2,000 per capita) we should expect that the average number of deaths (per 100,000) increases by 0.004.

The effect is statistically significant (***) next to the estimated coefficient indicates a p-value <0.01.

Let's present a graphical representation.

GDPpc v Avg_deaths from Covid-19



We now estimate a regression model which includes the GDP per capita (GDPpc) and the measure of Health expenditure as a percentage of GDP (HealthExp) as an explanatory variable.

$$Avg_deaths_i = \alpha + \beta \ GDPpc_i + \gamma \ HealthExp_i + u_i$$

mod3 <- lm(Avg deaths~GDPpc+HealthExp,data=table3)</pre>

```
stargazer(mod3, type="text")
##
                           Dependent variable:
##
                               Avg_deaths
## GDPpc
                                 0.0002
                                 (0.004)
## HealthExp
                                0.151***
                                 (0.038)
## Constant
                                  0.117
                                  (0.250)
## Observations
                                   143
## R2
                                  0.108
## Adjusted R2
                                  0.095
## Residual Std. Error 1.056 (df = 140)
                         8.437**** (df = 2: 140)
## F Statistic
```

*p<0.1; **p<0.05; ***p<0.01

Note:

Regression Analysis - What does it actually do?

Two interpretations

- 1) Finds the regression line (via $\widehat{\alpha}$ and $\widehat{\beta}$) that minimises the residual sum of squares $\Sigma(Avg_deaths_i \widehat{\alpha} \widehat{\beta} \ GDPpc_i)^2$. \rightarrow Ordinary Least Squares (OLS)
- 2) Finds the regression line (via $\widehat{\alpha}$ and $\widehat{\beta}$) that ensures that the residuals ($\widehat{u}_i = Avg_deaths_i \widehat{\alpha} \widehat{\beta} \ GDPpc_i$) are uncorrelated with the explanatory variable(s) (here Inc_i).

In many ways 2) is the more insightful one.

Regression Analysis - What does it actually do?

$$Avg_deaths = \alpha + \beta \ GDPpc + u$$

Assumptions

One of the regression assumptions is that the (unobserved) error terms u are uncorrelated with the explanatory variable(s), here GDPpc. Then we call GDPpc exogenous.

This implies that Cov(GDPpc, u) = Corr(GDPpc, u) = 0

In sample

$$Avg_deaths_i = \widehat{\alpha} + \widehat{\beta} \ GDPpc_i + \widehat{u}$$

Where $\widehat{\alpha} + \widehat{\beta} \ GDPpc_i$ is the regression-line.

In sample $Corr(GDPpc_i, \hat{u}_i) = 0$ (is ALWAYS TRUE BY CONSTRUCTION).

Regression Analysis - Underneath the hood?

$$Avg_deaths = \alpha + \beta \ GDPpc + u$$

What happens if you call

mod2 <- lm(Avg_deaths~GDPpc,data=table3)?</pre>

You will recall the following from Year 1 stats:

$$\begin{array}{lcl} \hat{\beta} & = & \dfrac{\widehat{Cov}(Avg_deaths,GDPpc)}{\widehat{Var}(GDPpc)} \\ \hat{\alpha} & = & \overline{Avg_deaths} - \hat{\beta} \; \overline{GDPpc} \end{array}$$

The software will then replace $\widehat{Cov}(Avg_deaths, GDPpc)$ and $\widehat{Var}(GDPpc)$ with their sample estimates to obtain $\hat{\beta}$ and then use that and the two sample means to get $\hat{\alpha}$.

Regression Analysis - Underneath the hood?

Need to recognise that in a sample $\hat{\beta}$ and $\hat{\alpha}$ are really

$$\begin{split} \hat{\beta} &= \frac{\widehat{Cov}(Avg_deaths,GDPpc)}{\widehat{Var}(GDPpc)} \\ &= \frac{\widehat{Cov}(\alpha + \beta \ GDPpc + u,GDPpc)}{\widehat{Var}(GDPpc)} \\ &= \frac{\widehat{Cov}(\alpha,GDPpc) + \beta \widehat{Cov}(GDPpc,GDPpc) + \widehat{Cov}(u,GDPpc)}{\widehat{Var}(GDPpc)} \\ &= \beta \frac{\widehat{Var}(GDPpc)}{\widehat{Var}(GDPpc)} + \frac{\widehat{Cov}(u,GDPpc)}{\widehat{Var}(GDPpc)} = \beta + \frac{\widehat{Cov}(u,GDPpc)}{\widehat{Var}(GDPpc)} \end{split}$$

So $\hat{\beta}$ is a function of the random term u and hence is itself a random variable. Once $\widehat{Cov}(Avg_deaths, GDPpc)$ and $\widehat{Var}(GDPpc)$ are replaced by sample estimates we get \underbrace{ONE} value which is draw from a random distribution.

Regression Analysis - The Exogeneity Assumption

Why is **assuming** Cov(GDPpc, u) = 0 important when, in sample, we are guaranteed $Cov(GDPpc_i, \hat{u}_i) = 0$?

If $Cov(GDPpc_i, u_i) = 0$ is **not true**, then

- 1) Estimating the model by OLS imposes an incorrect relationship
- 2) The estimated coefficients $\hat{\alpha}$ and $\hat{\beta}$ are biased (on average incorrect if we had many samples)
- 3) The regression model has no causal interpretation

As we cannot observe u_i , the assumption of exogeneity cannot be tested and we need to make an argument using economic understanding.

Regression Analysis - Outlook

$$y = \alpha + \beta \ x + u$$

Much of empirical econometric analysis is about making the exogeneity assumption (Corr(x, u) = 0) more plausible/as plausible as possible. But this begins with thinking why an explanatory variable x is endogenous.

- 1) Most models have more than one explanatory variable.
- 2) Including more relevant explanatory variables can make the exogeneity assumption more plausible.(*)
- 3) But fundamentally, if Cov(u, x) = 0 is implausible we need to find another variable z for which Cov(u, z) = 0 is plausible.
- (*) Including variables which are not explanatory variables can be very harmful. In particular variables which are determined by our explained and the explanatory variable (e.g. Health Expenditure in 2020!) can mask any relationship between the variables we are interested in.