


BMJ Open Investigating the effect of macro-scale estimators on worldwide COVID-19 occurrence and mortality through regression analysis using online country-based data sources

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ABSTRACT

Objective To investigate macro-scale estimators of the variations in COVID-19 cases and deaths among countries.

Design Epidemiological study.

Setting Country-based data from publicly available online databases of international organisations.

Participants The study involved 170 countries/territories, each of which had complete COVID-19 and tuberculosis data, as well as specific health-related estimators (obesity, hypertension, diabetes and hypercholesterolaemia).

Primary and secondary outcome measures The worldwide heterogeneity of the total number of COVID-19 cases and deaths per million on 31 December 2020 was analysed by 17 macro-scale estimators around the health-related, socioeconomic, climatic and political factors. In 139 of 170 nations, the best subsets regression was used to investigate all potential models of COVID-19 variations among countries. A multiple linear regression analysis was conducted to explore the predictive capacity of these variables. The same analysis was applied to the number of deaths per hundred thousand due to tuberculosis, a quite different infectious disease, to validate and control the differences with the proposed models for COVID-19.

Results In the model for the COVID-19 cases ($R^2=0.45$), obesity ($\beta=0.460$), hypertension ($\beta=0.214$), sunshine ($\beta=-0.157$) and transparency ($\beta=0.147$); whereas in the model for COVID-19 deaths ($R^2=0.41$), obesity ($\beta=0.279$), hypertension ($\beta=0.285$), alcohol consumption ($\beta=0.173$) and urbanisation ($\beta=0.204$) were significant factors ($p<0.05$). Unlike COVID-19, the tuberculosis model contained significant indicators like obesity, undernourishment, air pollution, age, schooling, democracy and Gini Inequality Index.

Conclusions This study recommends the new predictors explaining the global variability of COVID-19. Thus, it might assist policymakers in developing health policies and social strategies to deal with COVID-19.

Trial registration number ClinicalTrials.gov Registry (NCT04486508).

INTRODUCTION

COVID-19 caused by SARS-CoV-2 has spread over countries worldwide. On 11 March

Strengths and limitations of this study

- This study might represent recent macro-scale data of 170 countries and COVID-19 cases and deaths for 1 year.
- It is free from the vaccine effect.
- It contains a reliable predictive model incorporating social, economic and political indicators as well as health-related ones.
- Some macro-scale estimators were not recently published but we assume that there are no remarkable changes annually.
- There might be some interruption or reliability issues in COVID-19 cases and deaths data flow from countries.

2020, the WHO declared the condition as a public health emergency. With new variants emerging, experts now assume that SARS-CoV-2 would not only remain an endemic virus but would also continue to circulate in communities and would make a massive burden of disease and death for years.¹

The cumulative incidence (total number of cases per million) of COVID-19 (Total-Case-COVID19) and mortality (total number of deaths per million) of COVID-19 (Total-Death-COVID19) vary significantly among countries. Besides, old age, diabetes, high blood pressure, obesity, pregnancy, immunosuppression, cancer, and cardiovascular, respiratory (asthma and chronic obstructive pulmonary disease, etc), and chronic kidney disease are all well-known health-related factors that contribute to COVID-19 morbidity and mortality.^{2,3} Aside from these clinical factors and comorbidities affecting infection progression, there might be additional demographic, social, economic and environmental factors at the macro-level

differentiating countries in terms of TotalCase-COVID19 and TotalDeath-COVID19.

The term 'macro-level data' (eg, Human Development Index (HDI) 2020 by United Nations (UN)) could be defined as periodical (eg, weekly, monthly, quarterly or annual) and long-term data that were gathered, summarised, and published by the government, private, public, national and international organisations about a specific subject (eg, health, energy, society, economy, politics) in the world, continents, regions, countries, cities or territories. This kind of data shows the current position and trend of any subject in a comparable manner and guides the policymakers in countries and organisations for setting or updating their strategies and helps them take an action around that subject. Although healthcare executives and regulatory authorities have implemented regulations and recommendations such as wearing masks, social distancing, staying at home, isolation and lockdowns to limit the spread of SARS-CoV-2, considering these macro indicators will help steer national and international health policymakers to combat the COVID-19 pandemic.

To the best of our knowledge, there has been limited information on how the nature and dynamics of SARS-CoV-2 vary across the world.⁴⁻⁸ These studies have revealed that demographic, climatic, environmental, socioeconomic, and political indicators could help predict or clarify variations in TotalCase-COVID19 and TotalDeath-COVID19 among countries. In particular, the elderly ratio, the prevalence of comorbidities, population size, and health expenditure were shown to have been associated with the incidence and mortality of COVID-19 at various levels according to the aforementioned studies. However, these findings have been based on the early phase of the outbreak. The current study has been based on data (the cumulative number of cases and deaths) just before starting vaccinations around the world. Additionally, the findings might provide more consistent details about the progression of the disease because of covering data for a longer period. Therefore, the current study aims to identify health-related and socioeconomic macro-scale indicators that could explain some of the substantial variations in TotalCase-COVID19 and TotalDeath-COVID19 worldwide.

METHODS

Publicly accessible data about TotalCase-COVID19 and TotalDeath-COVID19 were obtained from the John Hopkins University–Center for Science and Engineering.⁹ While the process of the pandemic was still ongoing, mass immunisation has started in several countries as of December 2019. To eliminate the effect of immunisation, in this study, TotalCase-COVID19 and TotalDeath-COVID19 data were censored by 31 December 2020, when vaccination has not yet become widespread. Our dataset is available in Open Science Framework. Health-related indicators from the WHO and socioeconomic, climatic, and political

indicators from UN World Inequality Database and World Bank explaining variability in TotalCase-COVID19 and TotalDeath-COVID19 among countries were summarised in [table 1](#). Our dataset involves 17 macro-scale variables of 170 countries.

Statistical analyses

In this study, we conducted multiple linear regression analyses to explain variations in our target variables (ie, TotalCase-COVID19 and TotalDeath-COVID19). Before the analysis, we tested all the assumptions of multiple linear regression analysis by:

- ▶ Drawing 'actual target variable versus predicted target value' scatter plot diagram for testing linearity.
- ▶ Calculating standardised residual values for each case for detecting outlier as outside +3.
- ▶ Drawing a partial regression plot diagram for each predictor versus target variable for detecting the variance of predictors.
- ▶ Calculating variance inflation factor (VIF) for each predictor variable in the model for testing multicollinearity.
- ▶ Calculating Durbin-Watson test statistics for detecting autocorrelation of consequent errors.
- ▶ Drawing 'regression predicted value versus regression standardised residual' plot diagram for detecting heteroscedasticity.
- ▶ Drawing 'observed cumulative probability versus expected cumulative probability (P–P)' plot diagram of standardised regression residual for testing normality of residuals.

For eliminating the variance inflation and suppressing potential interactions among predictors, we have transformed all variables into the standardised form $[(x - \text{mean}(x))/\text{SD}(x)]$, so that we could get a more acceptable VIF ($\text{VIF} = 1/(1 - R^2) < 2.0$) values where R^2 is obtained by regression analysis in that the inspecting variable is regarded as a dependent variable and others are regarded as its predictor. We have applied Durbin-Watson statistics to test the autocorrelation in consequent error terms in the model and we have found all error terms are independent in all models since test statistics were between lower critical value and 4-lower critical value.¹⁰ After having inspected the plot diagrams and histograms, it could be said that all assumptions of multiple linear regression were almost met with the help of using standardised values of all predictors and target variables in all models. The exception for slight violation was observed as variances in error terms in P–P plots and predicted values against observed values scatter plot for homoscedasticity that could be questionable if other test results were unsatisfactory.

We conducted multiple linear regression analyses for different situations/scenarios and created predictive models to determine indicators aligned with these scenarios. After having tested the dataset against the assumptions of multiple linear regression, we investigate the dataset using some considerations for each scenario/

Table 1 The list and the definitions of indicators and the outcome variable used in the analysis

Factors	Variables	Description	Year	N	Mean±SD	Min	Max
Health related	Obesity ⁴¹	Obesity (defined as body mass index ≥ 30 , crude estimate %) prevalence in ≥ 18 -year-old people	2016	170	10.59±5.84	2.10	37.30
	Hypertension ⁴²	High blood pressure (systolic blood pressure ≥ 140) or (diastolic blood pressure ≥ 90) prevalence in ≥ 18 -year-old people	2015	170	23.45±5.31	12.60	41.00
	Alcohol ⁴³	Annual alcohol consumption per capita (litre) in ≥ 15 -year-old people	2018	170	4.96±3.87	0.00	20.05
	Diabetes ⁴⁴	Diabetes, raised fasting blood glucose (≥ 7.0 mmol/L) prevalence in ≥ 18 -year-old people	2014	170	7.70±3.99	1.00	22.10
	Cholesterol ⁴⁵	Hypercholesterolaemia, raised total cholesterol (≥ 6.2 mmol/L) prevalence in ≥ 25 -year-old people	2008	170	10.59±5.84	2.50	29.10
	Air pollution ⁴⁶	Mortality attributable to joint effects of household and ambient air pollution in adults such as acute respiratory infections, cerebrovascular diseases, ischaemic heart diseases, chronic obstructive pulmonary disease and lung cancer in adults (ambient and household air pollution attributable death rate (per 100 000 population)	2016	169	10.59±5.84	9.00	184.00
	Climatic	Sunshine ⁴⁷	Total sunshine hours (duration of sunlight in a year for countries)	2019	170	2519.18±576.3	1177.00

Continued

Table 2 Parameter testing results for linear regression for predicting ‘total cases per million people’

	Standardised coefficients	95% CI for β		Collinearity statistics	
	Beta	Lower bound	Upper bound	VIF	P value
Transparency	0.147	0.012	0.264	1.363	0.032
Obesity	0.460	0.311	0.547	1.213	0.001
Sunshine	-0.157	-0.266	-0.026	1.266	0.018
Hypertension	0.214	0.085	0.312	1.134	0.001

TotalCase-COV19=0.214×hypertension+0.460×obesity−0.157×sunshine+0.147×transparency.
 $R^2=0.449$, SE=0.70, Durbin-Watson=2.01.
 VIF, variance inflation factor.

model through a best subset approach that analyses data in terms of a predictor’s subset that could create applicable models that illuminate the next step for finding the best one. We could decide the better one by inspecting each model in terms of cost–benefit ratios such as R^2 , SE or Mallows’ Cp (min is preferred), Akaike Information Criterion (min is preferred) and Bayesian Information Criterion (min is preferred).¹¹

After determining the incorporated variables in a model, we applied a hierarchical regression model with forward selection and stepwise methods. There is no constant in our models because they were all insignificant statistically. Therefore, they were discarded. Using the information from best subset, literature and best practices, we tried a variety of models by evaluating the usage of intercept, handling missing values, handling outliers, applying the best technique for regression analysis (eg, enter, stepwise, forward/backward selection), determining thresholds for alpha to enter (0.05 through 0.15) and remove (0.05 through 0.15), and for accepting or rejecting the next variable to the incorporated model. We used the SPSS V.24 and Minitab V.19 trial version for analysing the data.

RESULTS

The descriptive statistics of all variables were provided in table 1. We have considered possible different models, explaining the TotalCase-COV19 and TotalDeath-COV19, including both common and different estimators that could explain the variations in countries. Therefore, we

have listed all possible significant models via the best subset method widely used in regression analyses for TotalCase-COV19 and TotalDeath-COV19 as demonstrated in online supplemental tables A and B.

Based on best subset analysis of TotalCase-COV19 and TotalDeath-COV19 modelling, the indicators ‘sunshine’, ‘obesity’, ‘hypertension’, ‘urbanisation’, ‘schooling’, ‘alcohol’, ‘democracy’, ‘transparency’ and ‘HDI’ are commonly found significant for the most of models. Additionally, for modelling the ‘TotalDeath-COV19’, ‘age’, ‘GINI’ (Gini Inequality Index) and ‘undernourishment’ were found significant; and for modelling ‘TotalCase-COV19’, ‘cholesterol’ was found significant. However, it was remarkable that the best subset model shows the results for only 139 countries with complete data for all variables at issue. That is why the recommended model could not be significant as in best subset because of missing values and their handling strategies. In all these regression analyses, we found that regression model fit and all parameter fits are significant ($p<0.05$).

Urbanisation, schooling, HDI, transparency, democracy and GINI are all inter-related factors, despite the lack of their multicollinearity. Thus, countries with desirable socioeconomic conditions are likely to have high levels of urbanisation, education, human development, transparency and democracy, as well as low-income inequality.

Table 2 shows the results of the regression model indicating the determinants of the TotalCase-COV19 worldwide. Here is the regression model indicating that transparency, sunshine, obesity and hypertension in

Table 3 Parameter testing results for linear regression for predicting ‘total deaths per million people’

	Standardised coefficients	95% CI for β		Collinearity statistics	
	Beta	Lower bound	Upper bound	VIF	P value
Alcohol	0.173	0.030	0.315	1.461	0.018
Urban	0.204	0.040	0.368	1.920	0.015
Obesity	0.279	0.116	0.442	1.915	0.001
Hypertension	0.285	0.146	0.423	1.380	0.000

TotalDeath-COV19=0.204×urban+0.285×hypertension+0.173×alcohol+0.279×obesity.
 $R^2=0.409$, SE=0.78, Durbin-Watson=2.29.
 VIF, variance inflation factor.

**Table 4** Parameter testing results for linear regression for predicting 'deaths due to tuberculosis'

	Standardised coefficients	95% CI for β		Collinearity statistics	
	Beta	Lower bound	Upper bound	VIF	P value
Obesity	-0.246	-0.427	-0.085	2.129	0.004
GINI	0.23	0.096	0.389	1.528	0.001
Air pollution	0.284	0.148	0.44	1.566	0.001
Undernourishment	0.245	0.084	0.402	2.014	0.003
Age	-0.453	-0.690	-0.224	4.193	0.001
Democracy	0.280	0.110	0.477	2.399	0.002
Schooling	0.253	0.048	0.459	3.303	0.016

TBDeath=-0.246×obesity+0.230×GINI+0.284×air pollution+0.245×undernourishment-0.453×age+0.280×democracy+0.253×schooling.

R²=0.535, SE=0.72, Durbin-Watson=1.79.

GINI, Gini Inequality Index; VIF, variance inflation factor.

countries could help to explain the variability of the Total-Case-COVID-19 across countries. In this regression analysis, R² value shows that the model with these four indicators explains 44.9% of all variability. As a result, the generated model is acceptable and comes out with expected roles of variables as aligned with the literature.

Table 3 shows the results of the regression model indicating the determinants of the TotalDeath-COVID-19 worldwide. The regression model revealed that urbanisation and alcohol in addition to obesity and hypertension might help to explain the variation in the TotalDeath-COVID-19 across countries. In this regression analysis, the R² value shows that the model with these four predictors explains 40.9% of all variability. Durbin-Watson value shows that there is no evidence for autocorrelation at 0.05 significance level.

Finally, we analysed the TBDeath by using the same 17 macro-scale indicators for validating/controlling purposes of our model proposals for the TotalCase-COVID-19 and TotalDeath-COVID-19. As an infectious disease quite dissimilar from COVID-19, the TBDeath model was expected to be different from our proposed model. As parallel in our expectations, it was quite different in terms of their involved variables except for obesity where 'air pollution', 'undernourishment' and 'GINI' are other significant factors rather than our proposed models (table 4). Exceptionally, VIF values of age and schooling variables are large, relatively and still tolerable since their VIF values are not exceeding a threshold of 5.¹²

DISCUSSION

Principal findings

The present study, which aims to explain the heterogeneity in the TotalCase-COVID-19 and TotalDeath-COVID-19 worldwide, by using an analysis of the health-related, socioeconomic, climatic and political macro indicators of 170 countries, showed two regression models. Well-established comorbidities, obesity and hypertension, were significant in both these models. Additionally, indicators sunshine and transparency for the TotalCase-COVID-19

model, as well as indicators alcohol and urbanisation for the TotalDeath-COVID-19, were important.

Interpretation and comparison with previous studies

Previous studies proposed that the comorbidities (obesity, hypertension, pulmonary disease, etc) contributed to the risk of COVID-19 and progression to severe disease by increasing the expression of ACE2 and/or transmembrane protease serine 2 on host lung cells and heightening the permissiveness of viral infection.¹³⁻¹⁵ In this study, the TotalCase-COVID-19 and TotalDeath-COVID-19 were greater in countries with a higher prevalence of obesity and hypertension. Global analyses of the variability of COVID-19 mortality among countries, as in our study, revealed that TotalDeath-COVID-19 was positively correlated with deaths due to comorbidities, including cardiovascular, chronic respiratory and kidney diseases, obesity and cancer.^{5 7 16}

The association between sunshine and COVID-19 is arguable in the literature. Studies revealing sunshine hours positively linked to COVID-19 growth claimed that on sunny days, outdoor activities increased along with a decline in adherence to lockdown rules, resulting in increased virus exposure and transmission.¹⁷⁻¹⁹ Other studies, on the other hand, suggested that sunny weather could help prevent the spread of COVID-19 by minimising air pollution and increasing vitamin D production.²⁰⁻²² Those studies revealed that ultraviolet light from the sun can inactivate SARS-CoV-2, reduce outdoor transmission or increase immune resistance through vitamin D production. These results are supported by our research, which found a high TotalCase-COVID-19 in countries with low sunshine.

Public corruption and GINI reflecting corruption in the flow of products, resources and services inside a nation were found to be associated with the TotalCase-COVID-19 and TotalDeath-COVID-19.^{23 24} The present study further supports these studies by indicating that there are more TotalCase-COVID-19 in countries with higher transparency (ie, lower corruption) levels. Thus, not only socioeconomic indicators but also political indicators could play a critical role in the COVID-19 outbreak. Lack of access

to reliable sources of information, as well as misinformation and inadequate communication, might lead to people disregarding government health alerts. Policies that promote the effective distribution of government budgets in public goods and services like healthcare and education, as well as policies that encourage transparency and information flow, could serve information accuracy about COVID-19's spread.^{23 25}

There is no country-level analysis demonstrating an association between alcohol consumption and the TotalCase-COVID19 and TotalDeath-COVID19 in the literature. Recent studies have emphasised that increasing alcohol consumption is linked to excessive production of proinflammatory cytokines by hepatic cells, resulting in higher levels of inflammatory markers (interleukin-8 and tumour necrosis factor- α), which has also been observed in patients with COVID-19.^{26–28} As a result, it was hypothesised that acute and chronic alcohol consumption might suppress the immune system, leading to reduced resistance to acute respiratory disease and SARS-CoV-2 infection, as well as facilitating the progression of COVID-19.^{29 30} The present study supports these findings, demonstrating that TotalDeath-COVID19 is higher in countries with high alcohol consumption.

Urbanisation has the potential to exacerbate disease and mortality by creating a slew of challenges such as public transportation, health and economic disparities, substandard living facilities, insufficient freshwater supplies, and ineffective sanitation and ventilation systems.^{4 6} Existing literature has shown that dense urban populations could cause less social distance, more trade and more human mobility, leading to multiple infection routes and more rapid spread of COVID-19.^{31 32} However, no significant relation was observed between urbanisation and TotalCase-COVID19, whereas some of the best subset models involve it. One explanation for this may be the difficulty of identifying infected people in countries with low socioeconomic status due to inadequate test conditions for COVID-19. Therefore, the effect of urbanisation on the TotalCase-COVID19 can be hard to observe. Besides that, developed countries with a larger urban population and advanced healthcare systems have longer life expectancies and a higher percentage of the population over 65 years, which could lead to higher mortality rates in high-income countries.⁸ Moreover, as tougher restriction policies for elderly people have been implemented, urbanisation might be a significant factor in TotalDeath-COVID19 rather than TotalCase-COVID19. Our study thus showed that countries with higher urban populations have higher TotalDeath-COVID19 in line with other global COVID-19 studies.^{6 33}

Best subset regression analyses show that heterogeneity in the TotalCase-COVID19 and the TotalDeath-COVID19 across countries may be attributed to variations in macro indicators including obesity and hypertension, urbanisation, schooling, transparency, democracy and HDI, as well as alcohol consumption and sunshine. Additionally, while the elderly population, GINI and undernourishment

were related to the TotalDeath-COVID19, hypercholesterolaemia was related to the TotalCase-COVID19. While the elderly population was a major parameter in nearly all TotalDeath-COVID19 models in our analysis, it was left out of the proposed models because it suppresses other estimators.

Other worldwide and national studies suggested that socioeconomic, political, climatic, environmental and ecological factors are correlated with the TotalCase-COVID19 and the TotalDeath-COVID19 at different contexts and levels.^{4–8 34–37} These studies revealed that countries with a high elderly population and level of democracy, as well as low levels of schooling and human development, had the highest mortality rate. Socioeconomic inequalities like low socioeconomic status, inadequate schooling, limited access to healthcare, and income inequality induce difficulty in accessing and affording healthy food, use of tobacco and alcohol, low physical activity, and lower use of preventive medicine. This situation might increase the prevalence of comorbidities by altering the gut microbiome, increasing local inflammation and compromising immunity. Hence, the high-risk population becomes more susceptible to COVID-19.²

To control the COVID-19 models, we perform a regression analysis to estimate the TBDeath consisting of the same macro-scale estimators. In this regression model, tuberculosis is associated with GINI, air pollution, undernourishment, age, democracy, schooling and obesity following the literature.^{38–40} There were remarkable differences between our COVID-19 models and tuberculosis as we expected since they are different infections and they should have their own (disease-specific) characteristics.

Limitations

There are certain limitations of the present study. First, not all health-related, socioeconomic, climatic and political indicators were up to date like COVID-19 deaths and cases in 2020. At that point, it is reasonable to expect that macro-scale indicators would not change significantly over the few years. Second, just 139 countries out of 170 were involved in the best subset regression analyses since only these countries had complete data. Third, definitions of COVID-19 cases and illness of people based on symptoms and diagnosis varied by country over a certain time that led some country data on the number of cases to be biased, and so it increases the residual in the model. Additionally, some interruptions on data flow about COVID-19 deaths and cases from countries could lead the model to be biased. To some extent, these error impacts (ie, bias) were minimised by standardising the data's relative position to other countries' data by considering overall mean and SD as well. In deciding the cause of death, a similar condition occurred in that some deaths could be recorded such as due to respiratory problems, other infections or multiple organ failure. Finally, the reliability of the PCR test that is used for diagnosing COVID-19 cases as the most preferred procedure across the world could also be regarded as an important limitation of the

study since false positive and false negative results of PCR test might cause deviations both in the COVID-19 cases and models.

CONCLUSION AND IMPLICATIONS OF FINDINGS

In conclusion, this original study reveals that health-related, as well as social, economic, climatic and political macro indicators, have been noteworthy in the COVID-19 outbreak. The findings of this study do not claim that these macro indicators cause disease or death directly because it does not conduct any clinical or laboratory research. Essentially, the present study, demonstrating that these macro indicators explain the variation in the number of COVID-19 cases and deaths across 170 countries, may serve as a basis for future clinical trials. Additionally, if it is assumed that COVID-19 would have a lasting effect in the world for an extended period, this study might assist policymakers in developing short-term and long-term health and social strategies to enhance these factors associated with COVID-19. Following the complete vaccination in nations, further research can be done to show how the vaccine affects these models.

Contributors SE is responsible for the overall content as guarantor. SE and SA had full access to all of the data in the study and take responsibility for the integrity and the accuracy of the data analysis. Concept and design—SE, SA and FI. Acquisition, analysis or interpretation of data—SE and FI. Drafting of the manuscript—SE and FI. Critical revision of the manuscript for important intellectual content—SE and FI. Statistical analysis—SE. Administrative, technical or material support—FI, AB, VG, EE, SM and AA. Supervision—SA.

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Supplementary Table A. Best Subset Models for Total Cases Per Million

Number of Variables	R ²	R ² (adjusted)	R ² (pred)	Mallows Cp	S	Age	Happiness	Gini	Transparency	Democracy	Urbanization	Schooling	AirPollution	HDI	Vegetable	SunShine	Undernourishment	Obesity	Cholesterol	Hypertension	Alcohol	Diabet		
1	38.5	38.1	36.1	16.0	0.78	X																		
1	38.5	38	36.7	16.1	0.78									X										
2	46.5	45.6	43.5	-0.4	0.73	X												X						
2	44.5	43.6	41.6	4.1	0.74									X					X					
3	48.1	46.8	44.3	-2.1	0.72	X									X			X						
3	47.9	46.7	44.4	-1.7	0.72									X				X	X					
4	49.5	47.9	45.2	-3.6	0.71									X	X			X	X					
4	49.2	47.6	44.2	-2.7	0.71	X									X			X	X					
5	50.0	47.9	45.1	-2.5	0.71					X				X	X			X	X					
5	49.9	47.9	45.0	-2.4	0.71									X	X			X	X			X		
6	50.4	48.0	44.5	-1.6	0.71					X				X	X			X	X	X				
6	50.4	48.0	44.5	-1.5	0.71									X	X			X	X	X	X			
7	50.7	47.9	44.3	-0.4	0.71					X				X	X			X	X	X	X			
7	50.6	47.8	44.0	-0.1	0.71					X	X			X	X			X	X	X				
8	50.9	47.7	43.8	1.2	0.71					X	X			X	X			X	X	X	X			
8	50.9	47.6	43.8	1.4	0.71					X			X	X	X			X	X	X	X			
9	51.1	47.4	43.3	2.8	0.72					X	X		X	X	X			X	X	X	X			
9	51.0	47.3	42.6	3.0	0.72					X	X	X		X	X			X	X	X	X			
10	51.2	47.1	42.2	4.6	0.72					X	X	X		X	X	X		X	X	X	X			
10	51.1	47.1	42.7	4.7	0.72			X		X	X		X	X	X			X	X	X	X			
11	51.2	46.7	41.5	6.5	0.72		X			X	X	X		X	X	X		X	X	X	X			
11	51.2	46.7	41.3	6.5	0.72					X	X	X		X	X	X		X	X	X	X	X		
12	51.3	46.3	40.7	8.4	0.72		X			X	X	X		X	X	X		X	X	X	X	X		
12	51.3	46.3	40.2	8.4	0.72	X	X			X	X	X		X	X	X		X	X	X	X	X		
13	51.3	45.9	40.0	10.3	0.73		X	X		X	X	X		X	X	X		X	X	X	X	X	X	
13	51.3	45.9	39.6	10.3	0.73	X	X	X		X	X	X		X	X	X		X	X	X	X	X		
14	51.4	45.5	38.6	12.2	0.73	X	X	X		X	X	X		X	X	X		X	X	X	X	X	X	
14	51.4	45.5	38.6	12.2	0.73		X	X		X	X	X		X	X	X	X	X	X	X	X	X	X	
15	51.4	45.1	37.7	14.1	0.73	X		X		X	X	X	X	X	X	X		X	X	X	X	X	X	
15	51.4	45.1	37.4	14.1	0.73	X	X	X		X	X	X	X	X	X	X		X	X	X	X	X	X	
16	51.4	44.6	37.0	16.0	0.73	X	X	X		X	X	X	X	X	X	X		X	X	X	X	X	X	
16	51.4	44.6	35.7	16.0	0.73	X	X	X		X	X	X	X	X	X	X	X	X	X	X	X	X	X	
17	51.4	44.1	35.3	18.0	0.74	X	X	X		X	X	X	X	X	X	X	X	X	X	X	X	X	X	

139 cases used; 31 cases contain missing values.

Supplementary Table B. Best Subset Models for Total Deaths Per Million

Number of Variables	R ²	R ² (adj)	R ² (pred)	Mallows Cp	S	Age	Happiness	Gini	Transparency	Democracy	Urbanization	Schooling	AirPollution	HDI	Vegetable	SunShine	Undernourishment	Obesity	Cholesterol	Hypertension	Alcohol	Diabet
1	35.6	35.1	33.2	28.1	0.86	X																
1	30.5	30.0	28.6	40.3	0.89									X								
2	42.3	41.4	39.3	13.8	0.81	X											X					
2	41.4	40.5	38.9	16.0	0.82										X		X					
3	45.6	44.3	41.9	7.9	0.79										X		X		X			
3	45.4	44.1	41.8	8.4	0.79	X									X		X					
4	48.4	46.7	43.8	3.3	0.78	X		X							X		X					
4	46.9	45.2	42.6	6.8	0.79		X								X		X		X			
5	49.7	47.6	44.5	2.2	0.77	X	X	X							X		X					
5	49.1	47.1	43.9	3.4	0.77	X		X	X						X		X					
6	50.5	48.1	45.0	2.1	0.77	X	X	X							X	X	X					
6	50.3	47.9	44.1	2.6	0.77	X	X	X							X		X		X			
7	51.2	48.4	44.7	2.5	0.76	X	X	X							X	X	X		X			
7	51.1	48.3	44.8	2.7	0.76	X	X	X		X					X	X	X					
8	51.6	48.4	45.2	3.5	0.76	X	X	X			X				X	X	X				X	
8	51.5	48.3	44.4	3.7	0.76	X	X	X	X						X	X	X		X			
9	52.0	48.4	44.8	4.6	0.76	X	X	X		X	X				X	X	X				X	
9	51.9	48.3	44.5	4.8	0.76	X	X	X			X				X	X	X		X	X		
10	52.3	48.3	44.4	5.9	0.76	X	X	X	X	X	X				X	X	X				X	
10	52.2	48.2	44.1	6.1	0.76	X	X	X	X		X				X	X	X		X	X		
11	52.5	48.1	44.1	7.4	0.77	X	X	X	X	X	X		X		X	X	X				X	
11	52.4	48.0	43.8	7.5	0.77	X	X	X	X		X		X		X	X	X		X	X		
12	52.7	47.9	43.4	8.8	0.77	X	X	X	X	X	X		X		X	X	X		X	X		
12	52.6	47.8	42.8	9.0	0.77	X	X	X	X		X	X	X	X	X	X	X		X	X		
13	52.9	47.6	42.4	10.5	0.77	X	X	X	X	X	X		X	X	X	X	X		X	X		
13	52.8	47.6	42.5	10.5	0.77	X	X	X	X	X	X		X		X	X	X		X	X	X	
14	53.0	47.3	41.4	12.2	0.77	X	X	X	X	X	X		X	X	X	X	X		X	X	X	
14	52.9	47.2	41.1	12.4	0.77	X	X	X	X	X	X		X	X	X	X	X	X	X	X	X	
15	53.0	46.9	40.1	14.1	0.77	X	X	X	X	X	X		X	X	X	X	X	X	X	X	X	X
15	53.0	46.8	40.8	14.2	0.77	X	X	X	X	X	X		X	X	X	X	X		X	X	X	X
16	53.0	46.5	39.5	16.0	0.78	X	X	X	X	X	X		X	X	X	X	X	X	X	X	X	X
16	53.0	46.4	39.4	16.1	0.78	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X
17	53.1	46.0	38.8	18.0	0.78	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X	X

139 cases used; 31 cases contain missing values.