

Strong negative covariation between toxoplasmosis and CoVID-19 at a global scale: a spurious indirect effect?

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Abstract

Coronaviruses may exert severely negative effects on the mortality and morbidity of birds and mammals including humans and domestic animals. Most recently CoVID-19 has infected over 2,360,000 humans and killed more than 165,000. Susceptibility to this disease appears to differ markedly across different societies but the factors underlying this variability are not known. Given that toxoplasmosis exerts both direct and immune-mediated antiviral effects, we hypothesize a negative covariation between toxoplasmosis and measures of the Covid-19 pandemic across countries. We obtained aged-adjusted toxoplasmosis prevalence of pregnant women from the literature. Since the differences in the CoVID-19 morbidity and mortality may depend on the different timing of the epidemics in each country, we applied a standard measure, i.e. the date of first documented CoVID-19 in each country as a proxy of susceptibility, with a statistical control for population size effects. Using these two indices, we show a highly significant negative co-variation between the two pandemics across 86 countries. Then, considering that the wealth of nations often co-varies with the prevalence of diseases, we introduced Gross Domestic Product per capita into our model. The prevalence of toxoplasmosis co-varies negatively, while the date of first CoVID-19 cases co-varies positively with GDP across countries. Further, to control for the strong spatial autocorrelation among countries, we carried out a Spatial Structure Analyses of the relationships between the date of first COVID-19, prevalence of toxoplasmosis, and GDP. Results of this analysis did not confirm a direct causal relationship between toxoplasmosis and susceptibility to the COVID-19 pandemics. As far as an analysis of observational data let us to suggest, it appears that the interaction between CoVID-19 and toxoplasmosis is mediated by GDP and spatial effects. This prompts the question whether the formerly known covariation between BCG vaccination and CoVID-19 might have also emerged as a spurious indirect effect.

Introduction

Coronaviruses are positive-stranded RNA viruses that may exert severely negative effects on the mortality and morbidity of a broad range of birds and mammals including humans and domestic animals. The strain called SARS-CoV-2 host-switched from bats to humans in Wuhan, China in November 2019 and subsequently gave rise to a devastating global pandemic called CoVID-19¹⁻³. Susceptibility of human societies appear to be markedly heterogeneous ranging from modest to very high morbidity. Contrary to general expectations, more developed, wealthier communities living under better hygienic conditions appear to be more threatened than others. Thus, Austria is more severely hit than Hungary, the Czech Republic than Slovakia, and Israel than Palestine or Jordan.

Sala and Miyakawa⁴ suggested that the different BCG vaccination policies across countries may partly explain such differences in susceptibility to CoVID-19. Indeed, higher morbidity and mortality is observed in societies with no obligatory BCG vaccination. However, vaccination schemes tend to be uniform within countries, thus this hypothesis cannot explain the huge within-country differences that are often observed, such as those between Northern vs. Southern Italy.

In accordance with the so-called ‘hygiene hypothesis’⁵, we hypothesize that certain common infections coming together with a less hygienic lifestyle facilitate some protection against CoVID–19.

Toxoplasmosis is a candidate infection for this purpose because it is one of the most widespread latent infections of humanity and its causative agent, the eukaryotic protozoan *Toxoplasma gondii*, is known to exhibit both direct and indirect antiviral effects^{6,7}.

T. gondii is an intracellular parasite that infects birds and mammals as intermediate hosts, while the sexual phase of its life cycle can only be completed in feline definitive hosts, most often in domestic cats. It is distributed in human societies mostly by semi-domestic, partly-feral cats that depredate on infected rodents and birds and then eat their prey.

Subsequently, the infective spores are released through their faeces and may get into direct contact with humans to cause infections. Alternatively, domestic animals may be infected by these spores and the consumption of their infected meat transmit *T. gondii* to humans. Thus, humans act like intermediate hosts, although they are not depredated by cats, and thus this is a dead-end for the parasites. ‘Luxury cats’ living on canned pet-food throughout their life may not transmit this infection. Asymptomatic infections are common in humans, especially among those living in the proximity of semi-feral domestic cats⁸.

T. gondii excretes Dense Granule Protein–7 (GRA–7) into the host cell that inhibits viral replication. Its effect has been proven both in vitro and in vivo against Indiana vesiculovirus, influenza A virus, Coxsackie virus, and herpes simplex virus. Overall, GRA–7 exhibits immune-stimulatory and a broad spectrum of antiviral activities via type I interferons signaling⁹. Moreover, in response to *T. gondii* infection, laboratory mice highly upregulate Immune Responsive Gene 1 in their lungs¹⁰. This is an interferon-stimulated gene that mediates antiviral effects against RNA viruses like the West Nile and Zika Viruses through its product named itaconate¹¹.

It is not known whether such direct and indirect antiviral mechanisms may have a significant negative effect on the coexistence of toxoplasmosis and viral epidemics on an epidemiological scale. Therefore, below we set out to test whether there is a negative co-variation between levels of toxoplasmosis and CoVID–19 pandemic at a global scale.

Results

The linear regression model without a spatial component indicates that toxoplasmosis ($N = 86$) is positively related to CoVID–19 Delay, while GDP per capita is negatively related to CoVID–19 Delay (Table 1, model 1, Figs. 1B, 1C). The total variation explained by these environmental variables is 36.8%.

Results of the linear model with spatial covariates are presented in Table 1, model 2. Akaike information criterion indicates a better fit of this model. Total variation of all environmental variables plus the spatial variable is 52.3%. Variance partitioning indicates that the total variation explained by GDP is 31.0%, by

toxoplasmosis prevalence is 23.6%, and by the spatial component it is 39.5%. However, variation explained only by the spatial competent was 18.9%, only by GDP was 10.8% (with the spatial component it was 13.2%) and by toxoplasmosis only it was 0.08% (with spatial component 5.9%).

Discussion

As we predicted, there was a negative correlation between measures of toxoplasmosis and CoVID–19 pandemics that would be considered ‘highly significant’ by conventional measures. Moreover, as it is the case with many other diseases¹², GDP per capita values co-vary negatively with toxoplasmosis prevalence. Opposite to this pattern, however, CoVID–19 emerged earlier in wealthier societies, most probably due to their more intensive participation of the global tourism and traffic industries.

Introducing a spatial component into the analysis, however, modified this result. It appears that CoVID–19 delay is greatly influenced by GDP per capita and spatial position of each country, while the effect of toxoplasmosis—contrary to our expectation—is negligible. This indicates that the strong negative covariation we have documented between prevalence of toxoplasmosis and the emergence of CoVID–19 on a global scale is likely to be a spurious side-effect mediated by GDP and spatial effects. However, this is not at all a proof for the lack of interaction between CoVID–19 and other pathogens. Perhaps the pathogen species, *Toxoplasma gondii*, we have chosen to investigate, or our methodologies, or the types of rough data we have analyzed were insufficient to verify such interactions.

Coronaviruses surround us, and we most probably get into contact with human coronaviruses (other than the SARS-CoV-2) like HCoV-OC43 and HCoV-229E, and animal coronaviruses like the porcine and feline coronaviruses (PEDV, FCoV) quite frequently, but these contacts are most often symptomless and unnoticed in humans. We do not exclude the scenario that the variability in human populations’ resistance vs. susceptibility to the current CoVID–19 pandemic is greatly influenced by interactions with such widespread but asymptomatic pathogens. Further, after the decline of the current CoVID–19 pandemic, more reliable indicators of population resistance vs. susceptibility will be available for our purposes, like the morbidity or mortality rates.

Our results may also serve as a cautionary factor regarding the apparent interaction between BCG vaccination and the CoVID–19 pandemics. Although statistically highly significant, that interaction was not controlled for GDP effects, neither for geographical context⁴. Thus we cannot exclude the possibility that the apparent influence of BCG vaccination on the current coronavirus pandemic is a spurious indirect effect, just as in our case with toxoplasmosis.

Different pathogen species utilizing the same host population as a shared resource tend to form an interactive pathogen community. Within this community, the ecological network of immune-mediated pathogen-pathogen interactions may define the emergence of disease¹³. How SARS-CoV–2 will be positioned in the global ecological network of human pathogens (probably the most species-rich pathogen community on Earth¹⁴) is yet to be seen.

Methods

Toxoplasmosis is routinely screened in pregnant women in several countries, thus nation-level prevalence values are available from the literature. We obtained data from Flegr and Dama¹⁵. Since the prevalence of this parasite is known to be age-dependent, these values were adjusted to a standard age following Lafferty¹⁶.

Currently, the CoVID–19 pandemic is still in its growing phase in most countries, but possibly saturated or already declining in others. Thus the actual measures of morbidity or mortality would be misleading for comparison among countries. Therefore, we compared the starting date, i.e. the date of the first documented case of the CoVID–19 disease in each country. For obvious reasons, countries with large population sizes are more likely to have earlier dates of the first case of the disease. To control for population size differences, we applied residuals taken from the (first date ~ log [population size]) regression (Fig. 1A; $\beta_0 = 178.241 \pm 18.258$ s.e., $\beta_{\log(\text{population})} = -10.673 \pm 2.527$ s.e., t value = -4.223 , $P < 0.001$). Positive residuals mean that countries had later start of the pandemic as expected (more resistant), while negative residuals indicate that countries had earlier starting dates (more susceptible). This variable is interpreted as a population-size-corrected time delay of first documented case of CoVID–19 in each country, and hereafter we called it 'CoVID–19 Delay'. Population sizes of each country were taken from UNData¹⁷, with values of countries that recently split to become independently treated accordingly. The first date of disease from CoVID–19 in each country originated from WHO¹⁸.

The prevalence of numerous human diseases depends on GDP per capita as an estimate of resource availability, and hence the ability to live a healthy life without exposure to zoonoses, and/or in the absence of untreated diseases¹⁹. To control for this, both in the case of toxoplasmosis and CoVID–19, we introduced Gross Domestic Product (GDP 2018) per capita and it was derived from World Bank data²⁰.

Further, statistical autocorrelation is a general feature of ecological variables measured across geographic space²¹. Since this effect, called spatial autocorrelation, violates the assumption of the independence of data required by most standard statistical procedures, we applied a Spatial Structure Analyses of the relationships between the date of first COVID–19 documented case (controlled for human population size), prevalence of toxoplasmosis (age adjusted), and GDP per capita.

Using a linear regression model, we analyzed the relationship between these three variables. We found no issue with multicollinearity between toxoplasmosis prevalence and GDP (VIF = 1.325). However, there were negative spatial correlation (Pearson's r = -0.495 , Corrected Pearson's r F = 14.598, Corrected D. F. = 44.921, $p < 0.001$; Fig. 2 and 3) between these two variables.

We tested all used variables for spatial autocorrelation with Moran's local indicator²¹. The index greater than 0 indicates how the pairs of locations are more similar, lower than 0 shows the pairs are less similar than expected by random pairs of observations. Moran's Is were computed for 11 distance classes and

were showed on spatial correlograms. Analyzes of these correlograms indicate (Supplementary materials) significant spatial similarity of neighbouring countries according to GDP, toxoplasmosis prevalence and CoVID-19 Delay (Figs. 2–4). In order to account for spatial autocorrelation, we used spatial eigenvector mapping (SEVM)²². This tool allows us to select an eigenvector (spatial filters) that minimizes Moran's I in model residuals. Then selected filters can be used as explanatory variables in the linear model. We used partial regression analyses to quantify how much of the variation of the response variable is explained by the spatial structure versus by the environmental variables.

All analyses were computed using SAM software²². Maps were created using QGIS software (version 3.8.3-Zanzibar).

Declarations

Acknowledgments

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Conflicts of interest

The authors declare no conflict of interest.

Credit author statement

Łukasz Jankowiak: Methodology, Software, Visualization, Formal analysis; **Lajos Rozsa:** Conceptualization, Writing - Original Draft, **Piotr Tryjanowski:** Conceptualization; Writing- Reviewing and Editing; **Anders Pape Møller:** Conceptualization, Supervision.

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Table

Table 1. Linear regression models explaining CoVID-19 Delay in different countries and due to toxoplasmosis and Gross Domestic Product (GDP) per capita.

Model 1 - not spatial covariate, $AIC_c = 701.163$, $r^2 = 0.368$

	Estimate	Std. Error	t value	P
Intercept	-2.054	4.548	-0.452	0.653
Toxoplasmosis	0.259	0.093	2.775	0.007
GDP	-0.316	0.076	-4.166	<.001

Model 2 - with spatial covarites¹, $AIC_c = 682.523$, $r^2 = 0.523$

Intercept	2.785	4.101	0.679	0.499
Toxoplasmosis	0.104	0.089	1.162	0.249
GDP	-0.292	0.067	-4.354	<.001

¹ – for spatial covariates details, see supplementary materials

Figures

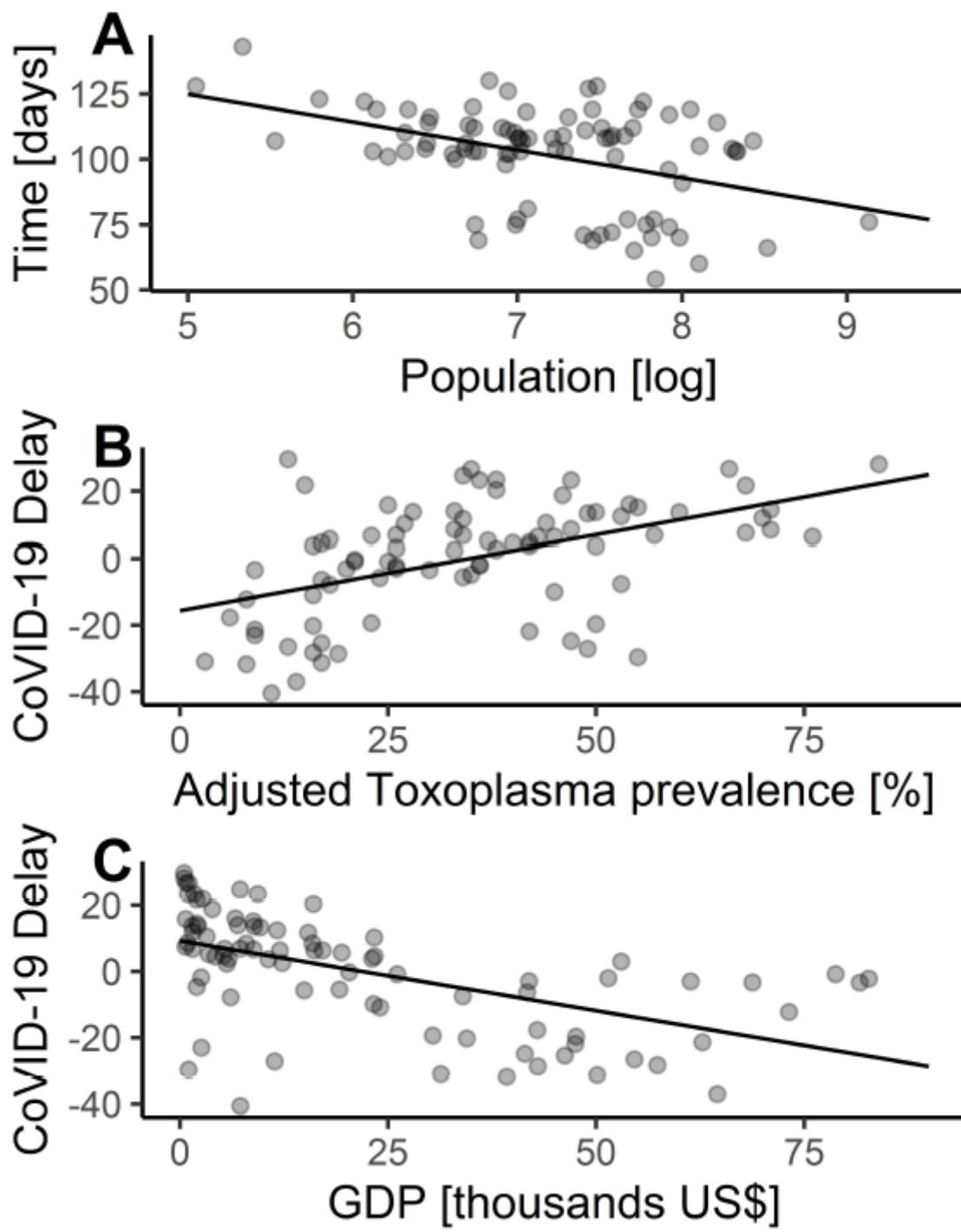


Figure 1

Linear regressions. A – starting date of epidemic counted since first case in China (China not included in analysis) with relationship to population size and residuals of this regression are used as dependent variable (CoVID-19 Delay). B - relationship of CoVID-19 Delay in days (population size corrected time delay of first case of CoVID-19 in a given country; negative values = CoVID-19 faster, positive values =

CoVID-19 later) and adjusted Toxoplasma prevalence. C – relationship of CoVID-19 Delay and Gross Domestic Product per capita (GDP).

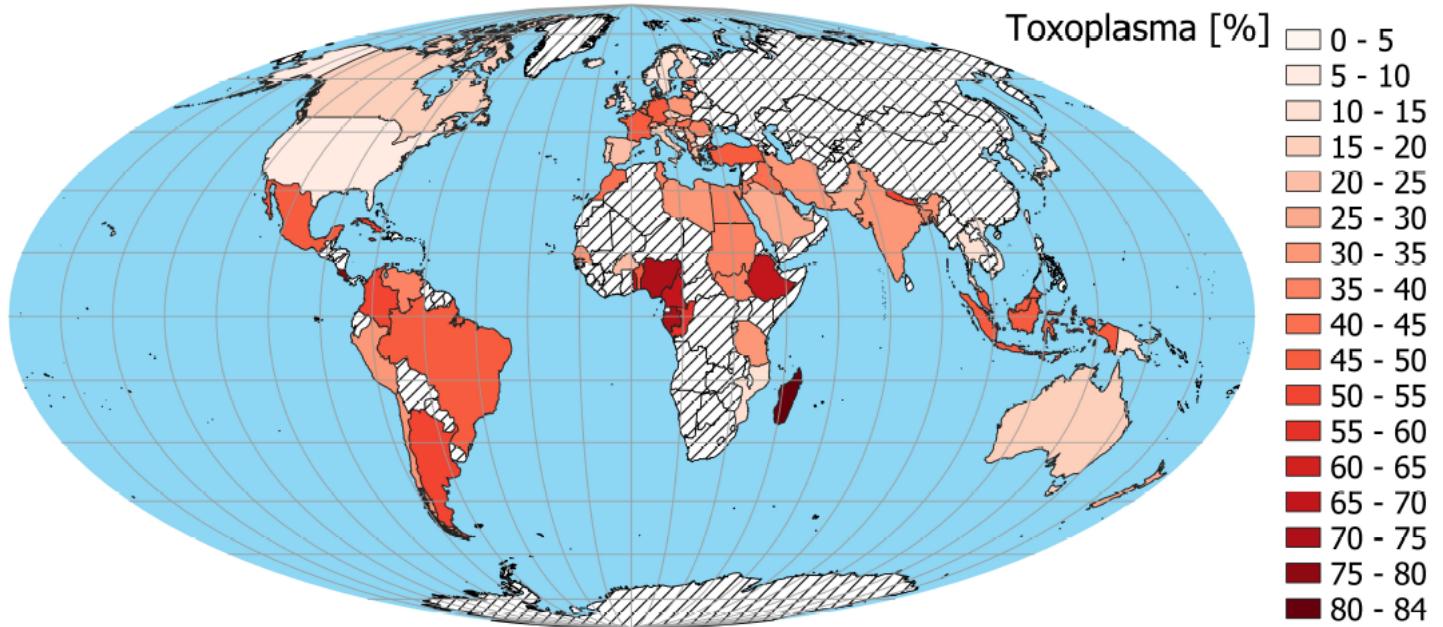


Figure 2

Linear regressions. A – starting date of epidemic counted since first case in China (China not included in analysis) with relationship to population size and residuals of this regression are used as dependent variable (CoVID-19 Delay). B - relationship of CoVID-19 Delay in days (population size corrected time delay of first case of CoVID-19 in a given country; negative values = CoVID-19 faster, positive values = CoVID-19 later) and adjusted Toxoplasma prevalence. C – relationship of CoVID-19 Delay and Gross Domestic Product per capita (GDP). The designations employed and the presentation of the material on this map do not imply the expression of any opinion whatsoever on the part of Research Square concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. This map has been provided by the authors.

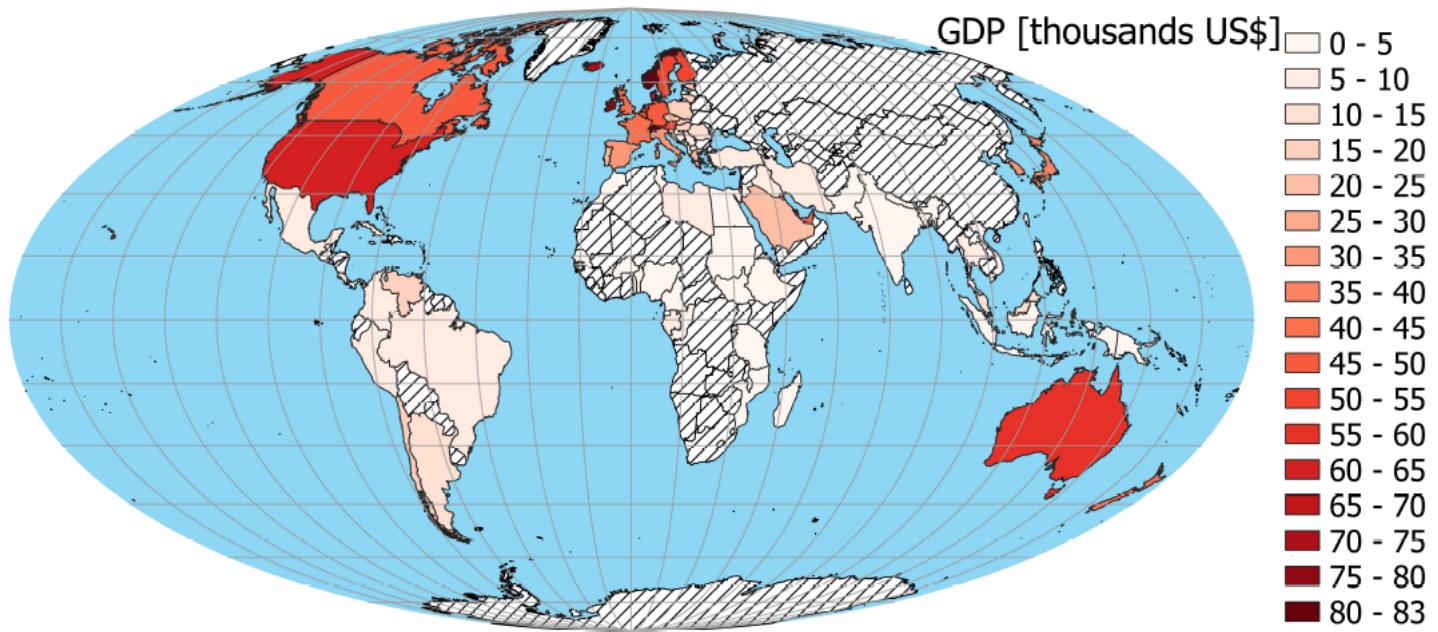


Figure 3

Spatial distribution of CoVID-19 Delay (population size corrected time delay of first case of CoVID-19 in given country; negative values = CoVID-19 faster, positive values = CoVID-19 later). China was not included in the analysis because it was treated as 1st day case. Selected were only countries with data on Toxoplasma prevalence. The designations employed and the presentation of the material on this map do not imply the expression of any opinion whatsoever on the part of Research Square concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. This map has been provided by the authors.

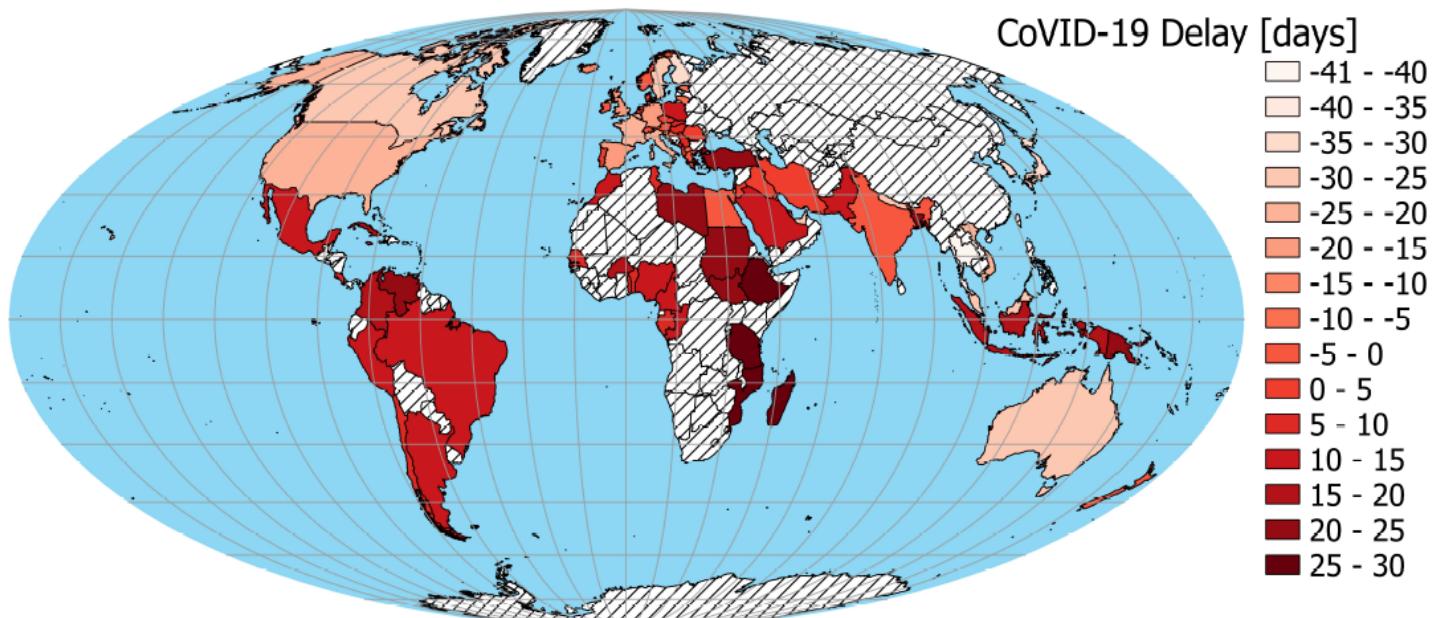


Figure 4

Spatial distribution of CoVID-19 Delay (population size corrected time delay of first case of CoVID-19 in given country; negative values = CoVID-19 faster, positive values = CoVID-19 later). China was not included in the analysis because it was treated as 1st day case. Selected were only countries with data on Toxoplasma prevalence. The designations employed and the presentation of the material on this map do not imply the expression of any opinion whatsoever on the part of Research Square concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. This map has been provided by the authors.

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