Evaluating alternative hypotheses to explain the downward trend in the indices of the COVID-19 pandemic death rate

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Background: In the ongoing Covid-19 pandemic, in the global data on the case fatality ratio (CFR) and other indices reflecting death rate, there is a consistent downward trend from mid-April to mid-November. The downward trend can be an illusion caused by biases and limitations of data or it could faithfully reflect a declining death rate. A variety of explanations for this trend are possible, but a systematic analysis of the testable predictions of the alternative hypotheses has not yet been attempted. Methodology: We state six testable alternative hypotheses, analyze their testable predictions using public domain data and evaluate their relative contributions to the downward trend. Results: We show that a decline in the death rate is real; changing age structure of the infected population and evolution of the virus towards reduced virulence are the most supported hypotheses and together contribute to major part of the trend. The testable predictions from other explanations including altered testing efficiency, time lag, improved treatment protocols and herd immunity are not consistently supported, or do not appear to make a major contribution to this trend although they may influence some other patterns of the epidemic. Conclusion: The fatality of the infection showed a robust declining time trend between mid April to mid November. Changing age class of the infected and decreasing virulence of the pathogen were found to be the strongest contributors to the trend.
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Abstract:

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Conclusion: The fatality of the infection showed a robust declining time trend between mid April to mid November. Changing age class of the infected and decreasing virulence of the pathogen were found to be the strongest contributors to the trend.

Keywords: Covid-19, case fatality rate, infection fatality rate, evolution of virulence, evolutionary medicine
Introduction

A consistent global trend in the current Covid-19 pandemic is that of decreasing case fatality rate (CFR). Whether this is an illusion created by some biases and limitations of data and if not, what are the possible alternative causes of the decline is the question we address here. In an on-going epidemic, an estimate of true death rate is rather difficult for several reasons including inappropriate diagnosis, time lag in diagnosis and mortality, and rapidly changing population dynamics. Nevertheless, a number of indices can reflect death rates with some limitations. The case fatality rate, a cumulative index of the number of deaths attributed to the virus divided by the confirmed positive cases so far, is most commonly used (Ghani et al., 2005; Reich et al., 2012). This index has shown a decline in global data [1,2]. (Note that we use square brackets for citing public domain data). However, being cumulative, this index is less sensitive to time trends and is dominated by the phase having more number of cases. We therefore use two other ratios here which are more faithful to the time trend, although somewhat more sensitive to stochasticity. The ratio of the number of new deaths in a day (ND) to the number of new cases registered (NC) on that day (ND/NC) is one index and the ratio of ND to the number declared recovered (NR) on that day (ND/NR) the other. The pros and cons of these indices are detailed in the supplementary material.

It can be seen in the global picture that from mid-April to mid-November, both the ratios show a consistent monotonic decline although not quite linearly [1-3] (fig1). The ND/NC ratio was close to 10 in mid-April, which came down to between 1 and 2 by mid-November. ND/NR also declined in similar proportion. So among the recorded global data there is a 4 to 5 fold difference in these indices between mid-April and mid-November.

It is well recognized from the early phase of the epidemic that CFR is a substantial underestimate of the infection fatality rate (IFR). IFR estimates the proportion of death from infected individuals [7] not all of which may be detected and diagnosed. There have been a number of attempts to estimate the IFR which are consistently smaller than CFR. However, the methods of estimating IFR vary, the IFR estimates are spatially and temporally fragmentary and therefore plotting time trends in IFR is not possible. If we assume that the bias in CFR is more or less constant over time, the time trend in CFR may reflect the time trend in IFR as well. However, it is quite possible that the bias itself has an increasing or
decreasing time trend. Therefore it is necessary to test for a temporal trend in the bias as well. The IFR estimates till June were between the ranges of 0.09 to 1.6 % with a mean of 0.68% (Meyerowitz-Katz and Merone 2020). However, in later serosurveys the estimate of the incidence increases substantially (Bobrovitz et al., 2020; Bhattacharyya et al., 2020). Therefore it is possible that even IFR is much smaller than the one estimated earlier or is declining considerably in time similar to CFR.

While underreporting of deaths has also been a serious problem, death underreporting has been disproportionately smaller than case underreporting (Bhattacharyya et al., 2020; Chakravarty 2020). Therefore any correction for the reporting bias will reveal a death rate much lower than the CFR. Using CFR along with the ND/NC and ND/NR ratios, wherever appropriate, we will first list the possible explanations, examine the differential testable predictions and evaluate their relative contributions in the declining trend. Further, since the different explanations are not mutually exclusive, we explore the possibility of their interactions.

The alternative hypotheses:

A. We first consider the possibility that the downward trend is illusionary for one or more of the following reasons.

(i) **Time lag in diagnosis and death:** Because of the inevitable but unpredictable time lag, by the time deaths are recorded, the number of cases might have gone up and therefore the indices of death rates are an underrepresentation of true death rate.

(ii) **Increased testing detected more asymptomatic cases:** The assumption behind this explanation is that the death rate was always as low as it is apparent today. The IFR that can be calculated from recent serosurveys is of the order of 0.02 to 0.07 % (Bhattacharyya et al., 2020; Ioannidis 2020; Pune Municipal Corporation 2020). If this was the true death rate right from the beginning, then the social implication is serious. The perceived death rate was the basis on which a number of measures were imposed by different state administrations, which have seriously affected the livelihood of a large population. If people develop an impression that it was a false alarm, they may lose trust in international and national health authorities including WHO. This may have serious long term
consequences. Therefore it is extremely important to evaluate this possibility carefully.

(iii) The age class of patients changed: Covid 19 is known to cause disproportionately higher deaths in the elderly (Ioannidis 2020; Bonanad et al., 2020; Williamson et al., 2020). So if more individuals of the younger age class are infected in the later phases of the epidemic, there would be an apparent decline in the indices of death rates.

B. An alternative possibility is that the downward trend in death rate is real and because of one or more of the following reasons.

(i) Increased efficiency of treatment regime brings down the death rate.

(ii) Increased immunity in the population reduces the death rate.

(iii) The virus progressively lost its virulence: A number of evolutionary epidemiology models indicate the possibility that under certain set of conditions progressive evolution of a newly invading virus leads to reduced virulence (Ewald 1994; Anderson and May 1982; Read 1994, Levin and Bull 1994, Kerr et al., 2012). This might be applicable to Covid-19.

We would now comparatively evaluate the alternative hypotheses for the apparent decline in the indices of death rate using data from public domain.

Methodology

Sources of data:

We use data from sources available in the public domain, mainly from WHO, CDC [1, 7, 8] and other open sources giving raw data as well as patterns seen in it. A systematic review was unwarranted because data were available from very standard and trusted sources such as WHO, CDC and other official national and international sources. The other sources mainly include World meter [3] Our World in Data [2], and Covid19India [4].

The data sources are listed below and specifically cited in the text with the appropriate hyperlink. There are certain inevitable limitations in the data. Data collection from different countries has subtle differences in the method of collection and accordingly some inevitable biases. We use pooled global data whenever available but some numbers are not available from all countries. For example, data on the number of tests performed is not available
globally. Furthermore some countries report the number of tests performed and others report
the number of individuals tested. The two are not inter-convertible and their implications
can be different. Whenever, getting global figures is not possible, we take the four countries
with maximum number of cases reported so far, namely the United States, Brazil, India and
Russia and perform country specific analysis.

**Testable predictions and evaluation of the hypotheses:**

**Hypothesis A(i): The time-lag between diagnosis, death and recovery:** This leads to
under or over-estimates of death rates. Simultaneous use of two ratios, ND/NC and ND/NR
(fig 1) can resolve the issue (see supplementary material S1). Since in the global data we
see a consistent decline in both the ratios, it is unlikely to be an illusion created by the time
lag effect. Further along the course of the epidemic, Rt, which measures the virus
transmission rate has been declining globally, (Contreras et al., 2020). When Rt is declining,
ND/NC tends to be overestimated. A decline in ND/NC in spite of a declining Rt is a robust
indication of a true decline. Therefore the time lag bias alone is unable to explain the
consistent decline in ND/NC.

**Hypothesis A(ii): Increased testing detected more asymptomatic cases:** If we assume
that the death rate was always low but in the initial phases of the epidemic, it was
overestimated because of inadequate testing efficiency. If the testing effort effectively
increased, the proportion of positives among the tested could have decreased. In contact
tracing, the number tested is expected to increase with the number of positive cases found at
a given time. Therefore rather than the absolute number of tests or the number of
individuals tested, the proportion of positives is a better inverse indicator of the effective
testing effort.

Using this principle, a testable prediction of this hypothesis is that there should be a positive
correlation between the proportion of positives among the tested and ND/NC. If changing
testing effort is the main reason for apparent change in death rates, then the time trend in
the proportion of positives should match the time trend in the ratio at least qualitatively.
That is if the proportion of positives is declining, the ratio should also decline.
Unfortunately global data on the number of tests are not available. In country level analysis
of the focal countries reporting maximum incidence, namely India, US and Russia we find
different and mutually inconsistent patterns (Fig 2). Across 91 countries for which the testing
data at least as recent as August 2020 was available at the time of analysis [5]. We do not find a
positive correlation between the proportion of positives and CFR which was expected if
testing efficiency was the main factor deciding the apparent CFR (Fig 3).

Furthermore, there have been many attempts to estimate the true proportion of infected
individuals in a population vis-a-vis the registered cases. This is attempted using a variety
of methods, at different locations and different phases of the epidemic. So a rigorous
comparative analysis is not possible. Grossly, while the estimated seroprevalences during
the trials completed by April are all less than 10 % with only one exception from Iran
(Bobrovitz et al., 2020), the July and August trials have estimates in the range of 15 to 57 %
(Pune Municipal Corporation 2020; Chakravarty 2020; Majiya et al., 2020). In areas where
a comparison of the fold increase in cases to fold increase in seroprevalence during the same
period is possible, in some countries such as in US [8] (Havers et al., 2020; Sutton, Cieslak
and Linder, 2020) and Brazil (Silveira et al., 2020) the ratio of the rise in seroprevalence to
rise in cases declined with time as expected by the hypothesis. But in many other countries
it increased (Pune Municipal Corporation 2020; Chakravarty 2020; Murhekar et al., 2020;
Nisar, et al., 2020; Stringhini et al., 2020; BBC news 2020). Collectively there is no
consistent evidence that the overestimation bias in the CFR and ND/NC ratio reduced with
time. Thus there is no consistent evidence that the bias in CFR estimates has been
consistently reducing with time. Therefore the hypothesis that the downward trend is caused
by a greater bias in the earlier phases of the epidemic and gradual removal of the bias
subsequently is not supported by evidence.

**Hypothesis A (iii): The age class of patient changed:** The age class distribution among
the diagnosed cases has evidently changed with time in global data [6]. Also the differential
case fatality across age classes is well known (Ioannidis 2020; Bonanad et al., 2020;
Williamson et al., 2020). Therefore it is very likely that at least qualitatively the changed
age class distribution may explain the apparent decline in death rates. We need to estimate
to what extent the changed age distribution explains the decline.

The > 65 age group has declined from 28% to 10 % among the infected population between
mid-April to end-July during which time ND/NC in the registered cases declined by 75-80% to remain at 20-25%. If we take a limiting assumption that all deaths are only in the >65 group, the death rate would have declined by 64% to come down to 36%. By this assumption a changing age distribution explains a substantial part but not the entire reduction in death rate. This is a limiting estimate assuming all deaths are in the >65 age class. The age class distribution of deaths is different in different countries since the age class distribution of the population itself is widely different. In the US, about 20% deaths were in the class < 65 and in India, 47% deaths were among the < 60% age class [9]. Considering that the 64 % reduction was only in the > 65 or > 60 class, the expected decline is 51.2 % and 33.92 % in the two countries respectively. This indicates that changing age class distribution may explain a substantial part of the apparent decline in death rate but still leaves a considerable decline unexplained. Further among the elderly age class itself, up to 2/3rd decline in death rate is shown in some countries [10] which also means that the decline is not entirely contributed by changing age distribution.

**Hypothesis B(i): Increased efficiency of treatment regime brings down the death rate:**

Global trends show that there is substantial reduction in the proportion of patients under critical care but there is only a marginal or no reduction in the proportion of deaths among patients under critical care (Fig 4) [2].

This need not necessarily mean that there is no improvement in treatment success. It is likely that the age distribution of the hospitalized has itself changed. If the mean age in this class has gone up, this may mask the expected improvement in the treatment. Even in that case the success rate of treatment does not explain the downward trend in death rates. Moreover the documented improvements in the success rates are limited. Clinical trials of all mainstream drugs have failed to show any significant reduction in mortality (WHO Solidarity Trial Consortium 2020; Spinner et al., 2020; Beigel et al., 2020; Li et al., 2020; Burki 2020; Mitjà et al., 2020; Simonovich et al., 2020). More relevant for our question is the fact that there is a considerable decline in the proportion needing critical care. Rather than improvement in treatment results, the need for treatment itself appears to have gone down.
Hypothesis B(ii): Increased immunity in the population reduces the death rate: Going by the registered cases a very small fraction of the population is exposed to the infection to see any major population level immunity change. However, going by seroprevalence, a much larger fraction of the population appears to have been sub clinically infected. It is possible that a substantial fraction of the population has indeed been exposed and presumably became immune. This immunity may be partially responsible for the reduced mortality. However, a critical question here is what made the large proportion of asymptomatic cases possible? If the virus was as virulent as initially perceived, the epidemic wouldn’t have progressed to cause so many asymptomatic infections. The low mortality and mild clinical course therefore should be a cause, rather than a consequence of the increasing population immunity (Johansson et al., 2021).

Hypothesis B(iii): The virus progressively lost its virulence: The concept that in the process of host-parasite co-evolution, a pathogen often evolves towards reduced virulence is quite old, but evolution towards loss of virulence is conditional. Not all pathogens appear to have reduced virulence when they coexist with a host population for a long time (Kerr et al., 2012; Ewald 1994; Anderson and May 1982; Bull 1994). Many evolutionary epidemiology models for optimum virulence were built and the continued theoretical development was backed up by epidemiological (Cressler et al., 2016; Best and Kerr 2000; Kale, Chanda and Wate 2002) as well as experimental studies (Berngruber et al., 2013, Tardy et al., 2019). With respect to the Covid-19 pandemic there are a multitude of reasons why evolution towards reduced virulence can be expected.

A fundamental assumption behind the evolution of virulence models is that the severity of symptoms and fatality is at least partly decided by the virulence of the virus. On the other hand, a virus that kills its host rapidly, gets less time to spread from the infected individual. Secondly the quarantine measures applied all over the world are likely to have created a selective force upon the virus. Since a symptomatic case is more likely to undergo testing and subsequently quarantined, a virulent variant causing more serious symptoms is more likely to be quarantined. A milder variant, has a greater chance of escaping detection and subsequent quarantine. Thirdly, if virulence is tightly correlated to viral loads and thereby transmission success, the virulent variant can have a greater selective advantage (Ewald 1994; Read 1994; Anderson and May 1982). If virulence does not have a direct correlation
with infection intensity and pathogen transmission, it is likely to be selected against (Levin
and Bull 1994). If the viral loads are not consistently high in serious cases, this advantage can be
assumed to be marginal and not sufficient to compensate the quarantine disadvantage. In
SARS Cov-2 infections there is a large overlap in the viral loads of symptomatic or fatal
versus asymptomatic cases and even in studies where there is a statistically significant
difference, the effect size or the magnitude of difference is not very large (Lennon et al.,
2020; Pujadas et al., 2020). Some mutants such as Gly614 showed higher viral loads but are
not more virulent (Long et al., 2020). The absence of viral load and virulence correlation
makes this virus an ideal case for evolution towards reduced virulence. Further at least
some components of the immune response are expressed in proportion to the extent of
invasion by the pathogen (Read 1994; Spooner and Yilmaz 2011). If the host response is
proportional to the extent of invasion, a milder virus may survive better in a more resistant
host, while a virulent one may do better in a susceptible host. If this is true, host immunity
and viral virulence are expected to interact in a positive feedback loop. As the population
acquires greater immunity, a milder virus can experience a selective advantage. Thus there
are multiple reasons why SARS-Cov-2 may have experienced a selective pressure for
reduced virulence.

Rejection or quantitatively inadequate explanation by other hypotheses is an indirect support
to the evolution hypothesis. But a true test of the hypothesis is to show evolutionary changes
in the genome indicating reduced virulence. Over 50,000 genomes have been sequenced in
various parts of the world during the pandemic. There are on average over 7 mutations per genome (Van et al., 2020; Mercatelli and Giorgi 2020). So the mutation rate can be
assumed to be sufficient to generate the required variation for natural selection. Among the
single nucleotide mutations there is a high proportion of recurrent non-synonymous
mutations (Spooner and Yilmaz 2011). The distribution of mutations along the genome is
highly non-random and the genes responsible for the pathogenicity have accumulated
significantly higher frequency of mutations (Choi et al., 2020). Both these patterns indicate
strong selective pressure for infectivity and/or virulence related genes. Genomic signatures
of strong selection coupled with the declining death rates not explained completely by other
hypotheses makes the evolution towards reduced virulence hypothesis more promising. One
of the mutations, D614G is suspected to increase the cell adhesion but whether it affects the
infectivity or virulence or both is not clearly known (Mercatelli and Giorgi 2020; Brufsky
2020). On the contrary, there are many other mutations in the region of the spike protein S1,
S2 and docking studies show that they reduce the stability of the host cell binding complex.
Furthermore in a comparative study of four regions of India, the ones with lower average
stability of mutants in the spike protein correlated negatively with local CFR (Banerjee et
al., 2020). These mutations are likely candidates responsible for the loss of virulence. There
are many other mutations in structural and non-structural proteins (Paul et al., 2020) which
are also likely to play a role in determining virulence. Virulence is a complex phenomenon
and from previous studies it is apparent that a large number of genes may contribute to viral
virulence (Kerr et al., 2012). Unfortunately as yet we do not have sufficient knowledge
linking specific mutations to their phenotypic effects. There is no standardized empirical test
of virulence to examine the effects of specific mutations on virulence. Of more direct
relevance is the observation that the mutational set observed among samples coming from
symptomatic versus asymptomatic cases is significantly different (Paul et al., 2020). This is
the most direct indicator that the asymptomatic clinical course is likely to be at least partly
driven by changes in the viral genome.

Conclusion

In summary, based on the available evidence, the hypotheses that there is a time lag in
diagnosis and death [A(i)]and increased testing detected more symptomatic cases [A(ii)]
fail to get any supportive evidence, increased efficiency of treatment regime brings down
the death rate [B(i)] does not appear to have made a strong contribution to the trend. The
change in age class of patients [A(iii)], death rate reduction due to increased immunity in
the population [B(ii)] and loss of virulence [B(iii)] are likely to be important causes of the
decline in death rates, out of which [B(ii)] increased immunity in the population needs prior
initiation by either or both the others. Therefore a combination of changed age class [A(iii)]
and loss of virulence [B(iii)] are most likely the primary causes of the declining trend and a
combination of both appears to be necessary to explain the trend quantitatively. The three
are not mutually exclusive and in fact may interact with each other. The cause of changed
age class in the infected population could be that the older age classes are being effectively
protected by the prevalent preventive measures, but it is also likely that the virus has evolved to infect younger age classes. It is of particular relevance here that some of the mutants are non-randomly represented in different age classes (Paul et al., 2020). This is possible if certain mutants are more likely to invade younger age classes.

An infectious epidemic is a complex process and multiple factors decide the outcomes. Our efforts to differentially support alternative hypothesis is limited by some constraints on data. We did not find a single data set giving all necessary parameters. Data on some variables is not available from all countries. Nevertheless, despite these limitations, the emerging patterns are strong enough to support some of the hypotheses. Previous studies on the changing parameters of an epidemic have mostly been published post facto (e.g. Best and Kerr 2000; Kale, Chanda and Watve 2002; Kerr et al., 2012). Analyzing patterns during an ongoing epidemic has a different set of challenges. Therefore we used a set of indices and methods different from earlier studies. But the trend in apparent reduction in virulence is similar to the earlier findings (Best and Kerr 2000; Kale, Chanda and Watve 2002; Kerr et al., 2012).

The specific contribution of our analysis is to suggest the evolutionary changes in a virus can be detected during an ongoing epidemic and policies to minimize population loss should take into account the ongoing evolution. Understanding the nature of selection on the virus can lead to virulence management strategies. Viruses have short generation times and high mutation rates and therefore can evolve very fast. Evolution on the background of host physiology, immunity, behavior, public health policies and available treatments should be an intrinsic part of epidemiological theories and models, which is likely to deepen our understanding of the disease process at different levels.

Data sources

1] WHO Coronavirus disease (COVID-19) Weekly Epidemiological Update and Weekly Operational Update https://www.who.int/emergencies/diseases/novel-coronavirus-2019/situation-reports

2] Our world in data: Statistics and research. Mortality rate in Covid-19. https://ourworldindata.org/mortality-risk-covid
3] Worldometer: Coronavirus worldwide graphs https://www.worldometers.info/coronavirus/

4] WHO Corona Disease (COVID 19). Covid19India.org

5] Wikipedia: Covid-19 testing https://en.wikipedia.org/wiki/COVID-19_testing

6] WHO coronavirus situation report, 5th August 2020 https://www.who.int/docs/default-source/coronaviruse/situation-reports/20200805-covid-19-sitrep-198.pdf?sfvrsn=f99d1754_2

7] World Health Organization. Estimating mortality from COVID-19. Available from https://www.who.int/news-room/commentaries/detail/estimating-mortality-from-covid-19

8] Interactive Serology Dashboard for Commercial Laboratory Surveys https://www.cdc.gov/coronavirus/2019-ncov/cases-updates/commercial-labs-interactive-serology-dashboard.html

9] Age class distribution link https://www.hindustantimes.com/india-news/85-deaths-in-45-plus-age-bracket-govt-data/story-II4EFnB7APRnLuU3MPmyZK.html

10] Declining COVID-19 Case Fatality Rates: https://www.cebm.net/covid-19/declining-covid-19-case-fatality-rates-across-all-ages-analysis-of-german-data

References

1. Anderson RM, May RM. 1982. Coevolution of Hosts and Parasites. *Parasitology*. DOI: 10.1017/S0031182000055360.

2. Banerjee R, Basak K, Ghosh A, Rajachandran V. 2020. Spike protein mutational
landscape in India: Could Muller’s ratchet be a future game-changer for COVID-19?

bioRxiv preprint, 2020.08.18.255570 (Available at https://www.biorxiv.org/content/10.1101/2020.08.18.255570v1) (Assessed August 18, 2020).

3. BBC News: India coronavirus: 'More than half of Mumbai slum-dwellers had Covid-19 (https://www.bbc.com/news/world-asia-india-53576653)

4. Beigel JH, Tomashek KM, Dodd LE, Mehta AK, Zingman BS, Kalil AC, Hohmann E, Chu HY, Luetkemeyer A, Kline S, Lopez de Castilla D, Finberg RW, Dierberg K, Tapson V, Hsieh L, Patterson TF, Paredes R, Sweeney DA, Short WR, Touloumi G, Lye DC, Ohmagari N, Oh M, Ruiz-Palacios GM, Benfield T, Fätkenheuer G, Kortepeter MG, Atmar RL, Creech CB, Lundgren J, Babiker AG, Pett S, Neaton JD, Burgess TH, Bonnett T, Green M, Makowski M, Osinusi A, Nayak S, Lane HC. 2020. Remdesivir for the Treatment of Covid-19 — Final Report. New England Journal of Medicine 383:1813–1826. DOI: 10.1056/nejmoa2007764.

5. Berngruber TW, Froissart R, Choisy M, Gandon S. 2013. Evolution of Virulence in Emerging Epidemics. PLoSPathog9(3): e1003209.

6. Best SM, Kerr PJ. 2000. Coevolution of host and virus: The pathogenesis of virulent and attenuated strains of myxoma virus in resistant and susceptible european rabbits. Virology. DOI: 10.1006/viro.1999.0104.

7. Bhattacharyya R, Bhaduri R, Ritoban Kundu ;, Salvatore ; Maxwell, Mukherjee B. 2020. Reconciling epidemiological models with misclassified case-counts for SARS-CoV-2 with seroprevalence surveys: A case study in Delhi, India. medRxiv:2020.07.31.20166249. DOI: 10.1101/2020.07.31.20166249.

8. Bobrovitz N, Arora RK, Yan T, Rahim H, Duarte N, Boucher E, Wyk J Van, Evans TG. 2020. Lessons from a rapid systematic review of early SARS-CoV-2 serosurveys. medRxiv:2020.05.10.20097451. DOI: 10.1101/2020.05.10.20097451.

9. Bonanad C, García-Blas S, Tarazona-Santabalbina F, Sanchis J, Bertomeu-González V, Fácila L, Ariza A, Núñez J, Cordero A. 2020. The Effect of Age on Mortality in Patients With COVID-19: A Meta-Analysis With 611,583 Subjects. Journal of the American Medical Directors Association 21:915–918. DOI: 10.1016/j.jamda.2020.05.045.
10. Brufsky A. 2020. Distinct Viral Clades of SARS-CoV-2: Implications for Modeling of Viral Spread. *Journal of Medical Virology*. DOI: 10.1002/jmv.25902.

11. Bull JJ. 1994. Virulence. *Evolution*. DOI: 10.1111/j.1558-5646.1994.tb02185.x.

12. Burki TK. 2020. Completion of clinical trials in light of COVID-19. *The Lancet Respiratory Medicine* 8:1178–1180. DOI: 10.1016/S2213-2600(20)30460-4.

13. Chakravarty, S., Estimating missing deaths in Delhi’s COVID-19 data. *medRxiv Preprint*, 2020, 2020.07.29.20164392. (Available at https://www.medrxiv.org/content/10.1101/2020.07.29.20164392v2) (Assessed July 30, 2020)

14. Choi B, Choudhary MC, Regan J, Sparks JA, Padera RF, Qiu X, Solomon IH, Kuo HH, Boucau J, Bowman K, Adhikari UD, Winkler ML, Mueller AA, Hsu TY, Desjardins M, Baden LR, Chan BT, Walker BD, Lichterfeld M, Brigl M, Kwon DS, Kanjilal S, Richardson ET, Jonsson AH, Alter G, Barczak AK, Hanage WP, Yu XG, Gaiha GD, Seaman MS, Cernadas M, Li JZ. 2020. Persistence and evolution of SARS-Cov-2 in an immune-compromised host. *NEJM* 383:2291-2293.

15. Contreras S, Villavicencio HA, Medina-Ortiz D, Saavedra CP, Olivera-Nappa A. 2020. Real-time estimation of R_0 for supporting public-health policies against COVID-19. DOI: 10.1101/2020.04.23.20076984.

16. Cressler CE, McLeod D V., Rozins C, Van Den Hoogen J, Day T. 2016. The adaptive evolution of virulence: A review of theoretical predictions and empirical tests. *Parasitology* 143:915–930. DOI: 10.1017/S003118201500092X.

17. Ewald, P., *Evolution of Infectious Disease*, Oxford University Press, Oxford,1994.

18. GhaniAC, Donnelly CA, Cox DR, Griffin JT, Fraser C, Lam TH, Ho LM, Chan WS, Anderson RM, Hedley AJ, Leung GM. 2005. Methods for estimating the case fatality ratio for a novel, emerging infectious disease. *American Journal of Epidemiology*. DOI: 10.1093/aje/kwi230.

19. Havers FP, Reed C, Lim T, Montgomery JM, Klena JD, Hall AJ, Fry AM, Cannon DL, Chiang CF, Gibbons A, Krapiunaya I, Morales-Betouille M, Roguski K, Rasheed MAU, Freeman B, Lester S, Mills L, Carroll DS, Owen SM, Johnson JA, Semenova V, Blackmore C, Blog D, Chai SJ, Dunn A, Hand J, Jain S, Lindquist S, Lynfield R, Pritchard S, Sokol T, Sosa L, Turabelidze G, Watkins SM, Wiesman J, Williams RW,
Yendell S, Schiffer J, Thornburg NJ. 2020. Seroprevalence of Antibodies to SARS-CoV-2 in 10 Sites in the United States, March 23-May 12, 2020. *JAMA Internal Medicine.* DOI: 10.1001/jamainternmed.2020.4130.

Ioannidis J. 2020. The infection fatality rate of COVID-19 inferred from seroprevalence data. *Bulletin of the World Health Organization* Article ID: BLT.20.265892

Johansson MA, Quandelacy TM, Kada S, Prasad PV, Steele M, Brooks JT, Slayton RB, Biggerstaff M, Butler JC. 2020. SARS-CoV-2 Transmission from People Without COVID-19 Symptoms. *JAMA Netw Open.* 4(1):e2035057.

Kale AU, Chanda S Watve M. 2002. Evolution of Mycobacterium leprae towards reduced virulence. *Current Science* 83:1078–1080.

Kerr PJ, Ghedin E, DePasse J V, Fitch A, Cattadori IM, Hudson PJ, Tscharke DC, Read AF, Holmes EC. 2012. Evolutionary History and Attenuation of Myxoma Virus on Two Continents. *PLoS Pathogens* 8. DOI: 10.1371/journal.ppat.1002950.

Lennon NJ, Bhattacharyya RP, Mina MJ, Rehm HL, Hung DT, Smole S, Woolley A, Lander ES, Gabriel SB. 2020. Comparison of viral levels in individuals with or without symptoms at time of COVID-19 testing among 32,480 residents and staff of nursing homes and assisted living facilities in Massachusetts. *medRxiv*:2020.07.20.20157792. DOI: 10.1101/2020.07.20.20157792.

Levin BR, Bull JJ. 1994. Short-sighted evolution and the virulence of pathogenic microorganisms. *Trends in Microbiology.* DOI: 10.1016/0966-842X(94)90538-X.

Li L, Zhang W, Hu Y, Tong X, Zheng S, Yang J, Kong Y, Ren L, Wei Q, Mei H, Hu C, Tao C, Yang R, Wang J, Yu Y, Guo Y, Wu X, Xu Z, Zeng L, Xiong N, Chen L, Wang J, Man N, Liu Y, Xu H, Deng E, Zhang X, Li C, Wang C, Su S, Zhang L, Wang J, Wu Y, Liu Z. 2020. Effect of Convalescent Plasma Therapy on Time to Clinical Improvement in Patients with Severe and Life-threatening COVID-19: A Randomized Clinical Trial. *JAMA - Journal of the American Medical Association* 324:460–470. DOI: 10.1001/jama.2020.10044.

Long SW, Olsen RJ, Christensen PA, Bernard DW, Davis JJ, Shukla M, Nguyen M, Saavedra MO, Yerramilli P, Pruitt L, Subedi S, Kuo HC, Hendrickson H, eskandari G, Nguyen HAT, Long JH, Kumaraswami M, Goike J, Boutz D, Gollihar J, McLellan JS, Chou CW, Javanmardi K, Finkelstein JJ, Musser JM. 2020. Molecular Architecture of Early Dissemination and Massive Second Wave of the SARS-CoV-2 Virus in a Major
Metropolitan Area. *medRxiv*. DOI: 10.1101/2020.09.22.20199125.

28. Majiya H, Aliyu-Paiko M, Balogu VT, Musa DA, Salihu IM, Kawu AA, Bashir YI, Sani RA, Baba J, Muhammad AT, Jibril FL, Bala E, Obaje NG, Aliyu BY, Muhammad RG, Mohammed H, Gimba NU, Uthman A, Liman HM, Sule AA, Joseph KJ, Makusidi MM, Isah, Abdullahi I, Ndagi U, Waziri B, Bisallah CI, Dadi-Mamud NJ, Ibrahim K, Adamu AK. 2020. Seroprevalence of COVID-19 in Niger State. *medRxiv*:2020.08.04.20168112. DOI: 10.1101/2020.08.04.20168112.

29. Mercatelli D, Giorgi FM. 2020. Geographic and Genomic Distribution of SARS-CoV-2 Mutations. *Frontiers in Microbiology* 11:1–13. DOI: 10.3389/fmicb.2020.01800.

30. Meyerowitz-Katz G, Merone L. 2020. A systematic review and meta-analysis of published research data on COVID-19 infection-fatality rates. *medRxiv*:2020.05.03.20089854. DOI: 10.1101/2020.05.03.20089854.

31. Mitjà O, Corbacho-Monné M, Ubals M, Alemany A, Suñer C, Tebé C, Tobias A, Peñafiel J, Ballana E, Pérez CA, Admella P, Riera-Martí N, Laporte P, Mitjà J, Clua M, Bertran L, Sarquella M, Gavilán S, Ara J, Argimon JM, Cuatrecasas G, Cañadas P, Elizalde-Torrent A, Fabregat R, Farré M, Forcada A, Flores-Mateo G, López C, Muntada E, Nadal N, Narejos S, Nieto A, Prat N, Puig J, Quiñones C, Ramírez-Viaplana F, Reyes-Urueña J, Riveira-Muñoz E, Ruiz L, Sanz S, Sentís A, Sierra A, Velasco C, Vivanco-Hidalgo RM, Zamora J, Casabona J, Vall-Mayans M, González-Beiras C, Clotet B; BCN-PEP-CoV2 Research Group. 2020. A Cluster-Randomized Trial of Hydroxychloroquine for Prevention of Covid-19. *NEJM*e. DOI: 10.1056/NEJMoa2021801

32. Murhekar MV, Selvaraju TB, Kiran Rade VS, Thangaraj JWV, Kumar MS, , Naman Shah RS, Turuk. 2020. Prevalence of SARS-CoV-2 infection in India: Findings from the national serosurvey, May-June 2020. *Indian J Med Res* 152:48–60. DOI: 10.4103/ijmr.IJMR_3290_20.

33. Nisar I, Ansari N, Amin M, Khalid F, Hotwani A, Rehman N, Rizvi A, Memon A, Ahmed Z, Ahmed A, Iqbal J, Saleem FA, Bashir U, Larremore DB, Fosdick B, Jehan F Serial population based serosurvey of antibodies to SARS-CoV-2 in a low and high transmission area of Karachi, Pakistan. *medRxivPreprint*, 2020,2020.07.28.20163451.
34. Paul D, Jani K, Kumar J, Chauhan R. 2020. Phylogenomic analysis of SARS-CoV-2 genomes from western India reveals unique linked mutations. *bioRxiv preprint* 2020.07.30.228460 (Available at [https://www.biorxiv.org/content/10.1101/2020.07.30.228460v2](https://www.biorxiv.org/content/10.1101/2020.07.30.228460v2)). (Assessed August 04, 2020).

35. Pujadas E, Chaudhry F, McBride R, Richter F, Zhao S, Wajnberg A, Nadkarni G, Glicksberg BS, Houldsworth J, Cordon-Cardo C. 2020. SARS-CoV-2 viral load predicts COVID-19 mortality. *The Lancet Respiratory Medicine* 2:2. DOI: 10.1016/s2213-2600(20)30354-4.

36. Pune Municipal Corporation. Technical report: Epidemiological and Serological Surveillance of COVID-19 in Pune City. 16th August 2020.

37. Read AF. 1994. The evolution of virulence. *Trends in Microbiology* 2:73–76. DOI: 10.1016/0966-842X(94)90537-1.

38. Reich NG, Lessler J, Cummings DAT, Brookmeyer R. 2012. Estimating Absolute and Relative Case Fatality Ratios from Infectious Disease Surveillance Data. *Biometrics*. DOI: 10.1111/j.1541-0420.2011.01709.x.

39. Silveira MF, Barros AJD, Horta BL, Pellanda LC, Victora GD, Dellagostin OA, Struchiner CJ, Burattini MN, Valim ARM, Berlezi EM, Mesa JM, Ikeda MLR, Mesenburg MA, Mantesso M, Dall’Agnol MM, Bittencourt RA, Hartwig FP, Menezes AMB, Barros FC, Hallal PC, Victora CG. 2020. Population-based surveys of antibodies against SARS-CoV-2 in Southern Brazil. *Nature Medicine* 26:1196–1199. DOI: 10.1038/s41591-020-0992-3.

40. Simonovich VA, Pratx LDB, Scibona P, Beruto MV, Vallone MG, Vázquez C, Savoy N, Giunta DH, Pérez LG, Sánchez ML, Gamarnik AV, Ojeda DS, Santoro DM, Camino PJ, Antelo S, Rainero K, Vidiella GP, Miyazaki EA, Cornstein W, Trabadelo OA, Ross FM, Spotti M, Funtowicz G, Scordo WE, Losso MH, Ferniot I, Pardo PE, Rodriguez E, Rucci P, Pasquali J, Fuentes NA, Esperatti M, Speroni GA, Nannini EC, Matteaccio A, Michelangelo HG, Follmann D, Lane HC, and Bellosio WH. 2020. A Randomized Trial of
Convalescent Plasma in Covid-19 Severe Pneumonia. NEJM. DOI: 10.1056/NEJMoa2031304.

41. Spinner CD, Gottlieb RL, Criner GJ, Arribas López JR, Cattelan AM, Soriano Viladomiu A, Ogbuagu O, Malhotra P, Mullane KM, Castagna A, Chai LYA, Roestenberg M, Tsang OTY, Bernasconi E, Le Turnier P, Chang SC, SenGupta D, Hyland RH, Osinusi AO, Cao H, Blair C, Wang H, Gaggar A, Brainard DM, McPhail MJ, Bhagani S, Ahn MY, Sanyal AJ, Huhn G, Marty FM. 2020. Effect of Remdesivir vs Standard Care on Clinical Status at 11 Days in Patients With Moderate COVID-19: A Randomized Clinical Trial. JAMA 324:1048–1057.

42. Spooner R, Yilmaz Ö. 2011. The role of reactive-oxygen-species in microbial persistence and inflammation. International Journal of Molecular Sciences. DOI: 10.3390/ijms12010334.

43. Stringhini S, Wisniak A, Piumatti G, Azman AS, Lauer SA, Baysson H, De Ridder D, Petrovic D, Schrempft S, Marcus K, Yerly S, Arm Vernez I, Keiser O, Hurst S, Posfay-Barbe KM, Trono D, Pittet D, Gétaz L, Chappuis F, Eckerle I, Vuilleumier N, Meyer B, Flahault A, Kaiser L, Guessous I. 2020. Seroprevalence of anti-SARS-CoV-2 IgG antibodies in Geneva, Switzerland (SEROCoV-POP): a population-based study. The Lancet 396:313–319. DOI: 10.1016/S0140-6736(20)31304-0.

44. Sutton M, Cieslak P, Linder M. 2020. Notes from the Field: Seroprevalence Estimates of SARS-CoV-2 Infection in Convenience Sample — Oregon, May 11–June 15, 2020. MMWR. Morbidity and Mortality Weekly Report 69:1100–1101. DOI: 10.15585/mmwr.mm6932a4.

45. Tardy L, Giraudeau M, Hill GE, McGraw KJ, Bonneaud C. 2019. Contrasting evolution of virulence and replication rate in an emerging bacterial pathogen. Proceedings of the National Academy of Sciences of the United States of America. DOI: 10.1073/pnas.1901556116.

46. Van Dorp L, Acman M, Richard D, Shaw LP, Ford CE, Ormond L, Owen CJ, Pang J, Tan CCS, Boshier FAT, Ortiz AT, Balloux F. 2020. Emergence of genomic diversity and recurrent mutations in SARS-CoV-2. Infection, Genetics and Evolution. DOI: 10.1016/j.meegid.2020.104351.

47. WHO Solidarity Trial Consortium. Repurposed antiviral drugs for COVID-19;
interim WHO SOLIDARITY trial results. medRxiv 2020, 2020.10.15.20209817

48. Williamson EJ, Walker AJ, Bhaskaran K, Bacon S, Bates C, Morton CE, Curtis HJ, Mehrkar A, Evans D, Inglesby P, Cockburn J, McDonald HJ, MacKenna B, Tomlinson L, Douglas IJ, Rentsch CT, Mathur R, Wong AYS, Grieve R, Harrison D, Forbes H, Schultze A, Croker R, Parry J, Hester F, Harper S, Perera R, Evans SJW, Smeeth L, Goldacre B. 2020. OpenSAFELY: factors associated with COVID-19 death in 17 million patients. Nature. DOI: 10.1038/s41586-020-2521-4.
Footnotes to figures:

Figure 1: Consistent monotonic decline by a factor of 4 to 5 in the ND/NC (blue line) and ND/NR (green line) ratios expressed as percentages between mid-April to mid-November. A. On the linear scale the decline appears to be saturated after July, but the subtle continued decline is revealed by the log scale (B).

Figure 2: Time trends in the proportion of positives detected per day during the testing effort (blue lines), which inversely reflects the testing efforts and the ND/NC ratio (red lines). Since global data on testing are not available, we use three countries with maximum number of cases reported. This includes India (A), Russia (B) and United States (C). If increased testing efficiency was mainly responsible for the apparent downward trend in death rate, the time trends in both the curves should have matched at least qualitatively, i.e. a downward trend in the proportion of positives should be matched by a downward trend in ND/NC. Observed trends do not support this prediction consistently.

Figure 3: Correlation between positive tests and CFR: If the true death rate was more or less constant spatiotemporally but the apparent trends were caused by testing biases, we would expect a positive correlation between the proportion tested positive and the CFR. In data on 91 countries for which test data were available and which had at least 100 deaths, the expected correlation is not seen ($r^2 = 0.0004$, NS).

Figure 4: The time trend in the ratio of patients under critical care on a given day to the number of cases recorded on that day (blue line) and percentage deaths under critical care (green line). The decline in death rates appears to be more due to decline in the proportion of serious cases needing critical care than due to success rate in critical care.
Figure 1

Consistent monotonic decline by a factor of 4 to 5 in the ND/NC (blue line) and ND/NR (green line) ratios expressed as percentages between mid-April to mid-November.

A. On the linear scale the decline appears to be saturated after July, but the subtle continued decline is revealed by the log scale (B).
Figure 2

Time trends in the proportion of positives detected per day during the testing effort (blue lines), which inversely reflects the testing efforts and the ND/NC ratio (red lines).

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