Transmission Dynamics of the Global COVID-19 Epidemic: Analytical Modeling and Future Prediction

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Abstract

Background: The outbreak and epidemic of COVID-19 has created worldwide impact and attracted global attention. Considerable effort has been devoted to the study of the transmission dynamics of COVID-19. However, there are lack of simple and straightforward expressions of the growing curves of important indicators, such as the cumulative number of confirmed cases and the cumulative number of dead cases.

Methods: We adopt two methods. The first method is based on regression analysis. We fit the available data into a curve by the method of least squares. The best curve is obtained by solving a multivariable minimization problem. The second method is based on differential equations. We establish an analytical model of transmission dynamics based on the susceptible-exposed-infectious-recovered-dead (SEIRD) process using a linear system of ordinary differential equations, which characterize the daily change in each compartment. The size of each compartment (i.e., the number of people in each stage of the SEIRD process) is readily available based on the solution to these differential equations.

Results: Both methods are applied to the COVID-19 epidemic data in the world as a case study. Furthermore, predictions of the cumulative number of confirmed cases and the cumulative number of dead cases in April 2020 using our models and methods are also provided. From a global perspective, unless powerful and effective social and medical impacts are made, by the end April of 2020, the cumulative number of confirmed cases is 23.333 and 36.068 millions respectively using regression analysis and analytical model, and the cumulative number of dead cases is 1.148 and 2.528 millions respectively using regression analysis and analytical model, based on the current situation.

Conclusions: In this paper, we make some progress towards analytical expressions of the daily growth of the cumulative number of confirmed cases and the cumulative number of dead cases, two most important and daily reported figures.

Keywords: Analytical model, COVID-19, linear system of ordinary differential equations, prediction, regression analysis, SEIRD process, transmission dynamics.
1 Introduction

1.1 Motivation

The outbreak and epidemic of COVID-19 has created worldwide impact and attracted global attention \[6, 13\]. The total number of coronavirus cases and the total number of coronavirus deaths in the world are growing exponentially and explosively \[2, 4, 5\]. Such pandemic has generated significant impact socially, economically, and politically.

Considerable effort has been devoted to the study of the transmission dynamics of COVID-19 \[9, 12\]. Li \textit{et al.} reported that the mean incubation period was 5.2 days, the mean serial interval was 7.5 days, the basic reproductive number $R_0$ was 2.2, and the epidemic was doubled in size every 7.4 days \[15\]. Liu \textit{et al.} reported that the average incubation duration of 2019-nCoV infection was 4.8 days, and estimated the $R_0$ values of 2019-nCoV as 2.90 and 2.92 using the methods of exponential growth (EG) and maximum likelihood (ML) respectively \[16\]. Nishiura \textit{et al.} estimated the median serial interval at 4.0 days \[19\]. Zhang \textit{et al.} mentioned that the ML value of $R_0$ was 2.28 for COVID-19 outbreak at the early stage on the Diamond Princess cruise ship \[21\]. Zhao \textit{et al.} pointed out that the early outbreak data largely follow the exponential growth, and estimated that the mean $R_0$ for the 2019-nCoV ranges from 2.24 to 3.58 \[22\]. Zhou \textit{et al.} found that the median duration of viral RNA shedding from oropharyngeal specimens was 20 days (range of 8–37 days) \[23\].

It is clear that all existing studies have only focused on individual quantities such as the basic reproductive number, the mean serial interval, the mean incubation period, and the mean recovery time. However, based on these quantities, it is still not possible to predict how the number of infected people, the number of recovered individuals, and the number of dead cases change daily. There are lack of simple and straightforward expressions of the growing curves of important indicators, such as the cumulative number of confirmed cases and the cumulative number of dead cases.

1.2 Contributions

In this paper, we make some progress towards analytical expressions of the daily growth of the cumulative number of confirmed cases and the cumulative number of dead cases,
two most important and daily reported figures. We adopt two methods.

The first method is based on regression analysis. We fit the available data into a curve by the method of least squares. The best curve is obtained by solving a multivariable minimization problem. The details are presented in Section 2.

The second method is based on differential equations. We establish an analytical model of transmission dynamics based on the susceptible-exposed-infectious-recovered-dead (SEIRD) process using a linear system of ordinary differential equations, which characterize the daily change in each compartment. The size of each compartment (i.e., the number of people in each stage of the SEIRD process) is readily available based on the solution to these differential equations. The details are presented in Section 3.

Both methods are applied to the COVID-19 epidemic data in the world as a case study. Furthermore, predictions of the cumulative number of confirmed cases and the cumulative number of dead cases in April 2020 using our models and methods are also provided. The details are presented in Section 4.

2 Regression Analysis

In this section, we develop a regression analysis method.

2.1 The Method

Assume that a group of \( n \) available data points \( (x_1,y_1),(x_2,y_2),\ldots,(x_n,y_n) \) are to be fit into a function \( y = f(a_1,a_2,\ldots,a_k,x) \), where \( a_1,a_2,\ldots,a_k \) are parameters of \( f \) to be decided. The method of least squares [1] is used to find \( a_1,a_2,\ldots,a_k \). The sum of squared residuals is

\[
E(a_1,a_2,\ldots,a_k) = \sum_{i=1}^{n} (y_i - f(a_1,a_2,\ldots,a_k,x_i))^2,
\]

where \( E \) is viewed as a function of \( a_1,a_2,\ldots,a_k \). To minimize \( E(a_1,a_2,\ldots,a_k) \), we only need to find \( a_1,a_2,\ldots,a_k \) such that \( \nabla E(a_1,a_2,\ldots,a_k) = 0 \), i.e., \( \partial E/\partial a_j = 0 \), for all \( 1 \leq j \leq k \). This is a multivariable minimization problem.

For transmission dynamics of coronavirus, we consider an exponential function in the form of \( f(x) = ab^x + c \), where \( a \) and \( b \) are parameters and \( c \) is a given constant, such that \( y_i \approx ab^{x_i} + c \), for all \( 1 \leq i \leq n \). To use the method of least squares to find \( a \) and \( b \), the
sum of squared residuals is
\[ E(a, b) = \sum_{i=1}^{n} (y_i - (ab^{x_i} + c))^2, \]
where \( E \) is viewed as a function of \( a \) and \( b \). To minimize \( E(a, b) \), we need
\[ \frac{\partial E}{\partial a} = \sum_{i=1}^{n} 2(y_i - (ab^{x_i} + c))(-b^{x_i}) = 0, \]
and
\[ \frac{\partial E}{\partial b} = \sum_{i=1}^{n} 2(y_i - (ab^{x_i} + c))(-ax_i b^{x_i - 1}) = 0. \]

To solve the above equations, we notice that the first equation implies that
\[ a = \frac{s_1}{s_2} = \frac{\sum_{i=1}^{n} (y_i - c)b^{x_i}}{\sum_{i=1}^{n} b^{2x_i}}, \]
and the second equation implies that
\[ a = \frac{s_3}{s_4} = \frac{\sum_{i=1}^{n} (y_i - c)x_i b^{x_i - 1}}{\sum_{i=1}^{n} x_i b^{2x_i - 1}}. \]

To find \( b \), we need \( s_1/s_2 = s_3/s_4 \). Notice that \( F(b) = s_1 s_4 - s_2 s_3 \) is an increasing function of \( b \). Hence, \( b \) can be found by using the standard bisection method ([8], pp. 22), such that \( F(b) = 0. \)

The quality of the above regression analysis can be evaluated by the adjusted relative error (ARE), defined for \( y_i \) as
\[ \text{ARE}_i = \left( \frac{y_i}{\max_{1 \leq i \leq n}(y_i)} \right) \times \left( \frac{ab^{x_i} + c - y_i}{y_i} \right) \times 100\%. \]

Notice that
\[ \left( \frac{ab^{x_i} + c - y_i}{y_i} \right) \times 100\% \]
is the relative error for \( y_i \). Since the magnitudes of \( y_1, y_2, \ldots, y_n \) can differ dramatically, simply taking the maximum relative error does not seem appropriate, since a large relative error for a very small \( y_i \) does not seem significant. Therefore, the relative error for \( y_i \) is adjusted by a factor of
\[ \frac{y_i}{\max_{1 \leq i \leq n}(y_i)}. \]
Table 1: Quality of Regression Analysis

|   | $C_{\text{World}}(t)$ |   | $D_{\text{World}}(t)$ |   |
|---|------------------------|---|------------------------|---|
| $t$ | Analytical | Actual | ARE | Analytical | Actual | ARE |
| 21 | 304874 | 304979 | -0.012% | 12931 | 13011 | -0.189% |
| 22 | 336362 | 337459 | -0.128% | 14465 | 14640 | -0.414% |
| 23 | 372070 | 378830 | -0.787% | 16225 | 16513 | -0.682% |
| 24 | 412565 | 422574 | -1.166% | 18244 | 18894 | -1.541% |
| 25 | 458490 | 471035 | -1.461% | 20560 | 21282 | -1.712% |
| 26 | 510571 | 531865 | -2.481% | 23217 | 24073 | -2.031% |
| 27 | 569634 | 596366 | -3.114% | 26264 | 27343 | -2.559% |
| 28 | 636614 | 663127 | -3.089% | 29759 | 30861 | -2.612% |
| 29 | 712575 | 723390 | -1.260% | 33769 | 34065 | -0.701% |
| 30 | 798718 | 784738 | 1.629% | 38369 | 37773 | 1.415% |
| 31 | 896410 | 858377 | 4.431% | 43645 | 42146 | 3.557% |

2.2 A Case Study

In this section, we apply our method to a case study.

Let $C_{\text{World}}(t)$ be the cumulative number of confirmed cases in the world by the $t$th day (i.e., $x_t = t$ and $y_t = C_{\text{World}}(t)$). Based on the data published by Worldometer [5] during March 1–31, 2020 (with March 1, 2020 as the first day), we get

$$C_{\text{World}}(t) = 16729.4550655 \times 1.1340597^t + 70000.$$  

Let $D_{\text{World}}(t)$ be the cumulative number of dead cases in the world by the $t$th day (i.e., $x_t = t$ and $y_t = D_{\text{World}}(t)$). Based on the data published by Worldometer [5] during March 1–31, 2020 (with March 1, 2020 as the first day), we get

$$D_{\text{World}}(t) = 584.4834711 \times 1.1470922^t + 2500.$$  

In Table 1, we display the analytical data, the actual data, and the ARE for $C_{\text{World}}(t)$ and $D_{\text{World}}(t)$ during March 21–31, 2020, with ARE$_i$ in the range $[-3.114\%, 4.431\%]$ for $C_{\text{World}}(t)$, and ARE$_i$ in the range $[-2.612\%, 3.557\%]$ for $D_{\text{World}}(t)$.  


3 An Analytical Model

In this section, we establish an analytical model of transmission dynamics.

3.1 The SEIRD Process

In this section, we describe the SEIRD process (see Figure 1).

The susceptible-exposed-infectious-recovered-dead (SEIRD) model belongs to the class of compartmental disease models [7]. “Susceptible”, “Exposed”, “Infectious”, “Recovered”, and “Dead” are compartments, and each individual in the population is assigned to one of these compartments. The model is extended from the SIR model [10] and the SEIR model [11, 14, 20] (other variations include SZR, SIZR, and SIZRQ models [18]).

- Susceptible individuals have no immunity to the disease, who may be exposed to the disease and move into the “Exposed" compartment through contact with an exposed or infectious person.

- Exposed individuals have been exposed to the disease, but have not shown sign or illness and thus behave like normal persons; however, they can transmit disease to others, and will move into the “Infectious" compartment.

- Infectious individuals show clear symptom of sickness, and have been tested and confirmed to be patients, who are typically quarantined or hospitalized, can still transmit disease to family members, medical staff, and other people, and eventually move into the “Recovered" or the “Dead" compartments.

- Recovered individuals can no longer become infected, typically because they have immunity from a prior exposure, which is often appropriate if immunity is long-lasting or the disease is being modeled over a relatively short time period.

- Dead individuals remain in the “Dead" compartment forever.

3.2 A System of Differential Equations

In this section, we establish an analytical model of transmission dynamics using a system of ordinary differential equations.
Let $S(t), E(t), I(t), V(t), D(t)$ be respectively the numbers of individuals in the susceptible, exposed, infectious, recovered, and dead compartments at time $t$ (measured in days). $S(t), E(t), I(t), V(t), D(t)$ are nonnegative functions of $t \geq 1$, which satisfy

$$S(t) + E(t) + I(t) + V(t) + D(t) = N,$$

and

$$dS(t)/dt + dE(t)/dt + dI(t)/dt + dV(t)/dt + dD(t)/dt = 0.$$

The daily reported cumulative number of confirmed cases is $C(t) = I(t) + V(t) + D(t)$. The daily reported cumulative number of dead cases is $D(t)$.

Let $N, R_E, R_I, T_E, T_I, \alpha$ be positive constants defined below.

- $N$ is the size of (i.e., the number of people in) the population.
- $R_E$ is the reproductive number of an exposed individual.
- $R_I$ is the reproductive number of an infectious individual.
- $T_E$ is the average number of days in the “Exposed” compartment.
- $T_I$ is the average number of days in the “Infectious” compartment.
- $\alpha$ is the percentage of infectious individuals who eventually die.

Notice that the basic reproductive number is actually $R_0 = R_E + R_I$. 

Figure 1: The SEIRD process
We have the following linear system of ordinary differential equations:

\[
\frac{dS(t)}{dt} = -\left(\frac{R_E}{T_E} \cdot E(t) + \frac{R_I}{T_I} \cdot I(t)\right),
\]
\[
\frac{dE(t)}{dt} = \frac{R_E - 1}{T_E} \cdot E(t) + \frac{R_I}{T_I} \cdot I(t),
\]
\[
\frac{dI(t)}{dt} = \frac{E(t)}{T_E} - \frac{I(t)}{T_I},
\]
\[
\frac{dV(t)}{dt} = (1 - \alpha) \frac{I(t)}{T_I},
\]
\[
\frac{dD(t)}{dt} = \alpha \frac{I(t)}{T_I},
\]

with given initial values \(S(1), E(1), I(1), V(1), D(1)\). These equations are elaborated as follows.

- \(dS(t)/dt\): \(R_E/T_E\) is the number of newly exposed people contacted by an exposed individual every day, and \((R_E/T_E)E(t)\) is the total number of such people on the \(t\)th day. Similarly, \(R_I/T_I\) is the number of newly exposed people contacted by an infectious individual every day, and \((R_I/T_I)I(t)\) is the total number of such people on the \(t\)th day. Therefore, the total number of newly exposed people (who leave the "Susceptible" compartment) is \((R_E/T_E)E(t) + (R_I/T_I)I(t)\) on the \(t\)th day.

- \(dE(t)/dt\): The number of newly exposed people (who move into the "Exposed" compartment) is \((R_E/T_E)E(t) + (R_I/T_I)I(t)\) on the \(t\)th day. On the other hand, the number of exposed people who move into the "Infectious" compartment is \(E(t)/T_E\) on the \(t\)th day. Therefore, the change in the "Exposed" compartment is \(((R_E - 1)/T_E)E(t) + (R_I/T_I)I(t)\) on the \(t\)th day.

- \(dI(t)/dt\): The number of exposed people who move into the "Infectious" compartment is \(E(t)/T_E\) on the \(t\)th day. On the other hand, the number of infectious people who leave the "Infectious" compartment is \(I(t)/T_I\) on the \(t\)th day. Therefore, the change in the "Infectious" compartment is \(E(t)/T_E - I(t)/T_I\) on the \(t\)th day.

- \(dV(t)/dt\): The number of infectious people who move into the "Recovered" compartment is \((1 - \alpha)(I(t)/T_I)\) on the \(t\)th day.

- \(dD(t)/dt\): The number of infectious people who move into the "Dead" compartment is \(\alpha(I(t)/T_I)\) on the \(t\)th day.
3.3 An Analytical Solution

In this section, we provide an analytical solution to the linear system of ordinary differential equations.

Notice that \( \frac{dE(t)}{dt} \) and \( \frac{dI(t)}{dt} \) form an autonomous linear system of differential equations:

\[
\frac{dE(t)}{dt} = \left( \frac{R}{T_E} - \frac{1}{T_E} \right) E(t) + \left( \frac{R}{T_I} \right) I(t),
\]
\[
\frac{dI(t)}{dt} = \left( \frac{1}{T_E} \right) E(t) - \left( \frac{1}{T_I} \right) I(t),
\]

It is well known that such differential equations accommodate analytical solutions [3].

Let

\[
A = \begin{bmatrix}
\frac{R}{T_E} - \frac{1}{T_E} & \frac{R}{T_I} \\
\frac{1}{T_E} & -\frac{1}{T_I}
\end{bmatrix}
\]

If matrix \( A \) has eigenvalues \( \lambda_1 \) and \( \lambda_2 \) with corresponding eigenvectors \( v_1 = (x_1, y_1) \) and \( v_2 = (x_2, y_2) \), then we have

\[
E(t) = c_1 x_1 e^{\lambda_1 t} + c_2 x_2 e^{\lambda_2 t},
\]
\[
I(t) = c_1 y_1 e^{\lambda_1 t} + c_2 y_2 e^{\lambda_2 t},
\]

for some constants \( c_1 \) and \( c_2 \), which can be decided based on the initial conditions \( E(1) = b_1 \), and \( I(1) = b_2 \), by solving the following equations:

\[
c_1 x_1 e^{\lambda_1} + c_2 x_2 e^{\lambda_2} = b_1,
\]
\[
c_1 y_1 e^{\lambda_1} + c_2 y_2 e^{\lambda_2} = b_2,
\]

whose solution is

\[
c_1 = \begin{vmatrix}
x_2 e^{\lambda_2} & b_1 \\
y_2 e^{\lambda_2} & b_2
\end{vmatrix} / \begin{vmatrix}
x_2 e^{\lambda_2} & x_1 e^{\lambda_1} \\
y_2 e^{\lambda_2} & y_1 e^{\lambda_1}
\end{vmatrix},
\]
\[
c_2 = \begin{vmatrix}
x_1 e^{\lambda_1} & b_1 \\
y_1 e^{\lambda_1} & b_2
\end{vmatrix} / \begin{vmatrix}
x_1 e^{\lambda_1} & x_2 e^{\lambda_2} \\
y_1 e^{\lambda_1} & y_2 e^{\lambda_2}
\end{vmatrix}.
\]
The eigenvalues of the matrix $A$ are values of $\lambda$ that satisfy the equation

\[
\begin{vmatrix}
\frac{R_E-1}{T_E} - \lambda & \frac{R_I}{T_I} \\
\frac{1}{T_E} & -\frac{1}{T_I} - \lambda
\end{vmatrix} = 0,
\]

or,

\[
\begin{vmatrix}
-\left(\lambda - \frac{R_E-1}{T_E}\right) & \frac{R_I}{T_I} \\
\frac{1}{T_E} & -(\lambda + \frac{1}{T_I})
\end{vmatrix} = 0,
\]

that is,

\[
\lambda^2 - \left(\frac{R_E - 1}{T_E} - \frac{1}{T_I}\right)\lambda - \frac{R_E + R_I - 1}{T_E T_I} = 0.
\]

The solutions are

\[
\lambda_1 = \frac{1}{2} \left(\left(\frac{R_E - 1}{T_E} - \frac{1}{T_I}\right) + \sqrt{\left(\frac{R_E - 1}{T_E} - \frac{1}{T_I}\right)^2 + 4 \left(\frac{R_E + R_I - 1}{T_E T_I}\right)}\right),
\]

and

\[
\lambda_2 = \frac{1}{2} \left(\left(\frac{R_E - 1}{T_E} - \frac{1}{T_I}\right) - \sqrt{\left(\frac{R_E - 1}{T_E} - \frac{1}{T_I}\right)^2 + 4 \left(\frac{R_E + R_I - 1}{T_E T_I}\right)}\right).
\]

The eigenvector $v_i = (x_i, y_i)$ satisfies

\[
\begin{bmatrix}
-\left(\lambda_i - \frac{R_E - 1}{T_E}\right) & \frac{R_I}{T_I} \\
\frac{1}{T_E} & -(\lambda_i + \frac{1}{T_I})
\end{bmatrix}
\begin{bmatrix}
x_i \\
y_i
\end{bmatrix} = 0.
\]

We can set $x_i = 1$, and $y_i = (\lambda_i - \frac{R_E - 1}{T_E})/(\frac{R_I}{T_I})$, for $i = 1, 2$.

Based on $E(t)$ and $I(t)$, we can get

\[
V(t) = \left(1 - \frac{\alpha}{\lambda_1 T_I}\right) c_1 y_1 (e^{\lambda_1 t} - e^{\lambda_1}) + \left(1 - \frac{\alpha}{\lambda_2 T_I}\right) c_2 y_2 (e^{\lambda_2 t} - e^{\lambda_2}) + V(1),
\]

and

\[
D(t) = \left(\frac{\alpha}{\lambda_1 T_I}\right) c_1 y_1 (e^{\lambda_1 t} - e^{\lambda_1}) + \left(\frac{\alpha}{\lambda_2 T_I}\right) c_2 y_2 (e^{\lambda_2 t} - e^{\lambda_2}) + D(1),
\]

and

\[
C(t) = I(t) + V(t) + D(t)
= \left(1 + \frac{1}{\lambda_1 T_I}\right) c_1 y_1 e^{\lambda_1 t} + \left(1 + \frac{1}{\lambda_2 T_I}\right) c_2 y_2 e^{\lambda_2 t} - \frac{c_1 y_1 e^{\lambda_1}}{\lambda_1 T_I} - \frac{c_2 y_2 e^{\lambda_2}}{\lambda_2 T_I} + V(1) + D(1),
\]

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and

\[
S(t) = N - E(t) - I(t) - V(t) - D(t) \\
= N - c_1 x_1 e^{\lambda_1 t} - c_2 y_2 e^{\lambda_2 t} \\
- \left(1 + \frac{1}{\lambda_1 T_l}\right) c_1 y_1 e^{\lambda_1 t} - \left(1 + \frac{1}{\lambda_2 T_l}\right) c_2 y_2 e^{\lambda_2 t} + \frac{c_1 y_1 e^{\lambda_1 t}}{\lambda_1 T_l} + \frac{c_2 y_2 e^{\lambda_2 t}}{\lambda_2 T_l} - V(1) - D(1).
\]

Notice that since \(\lambda_2 < 0\), we have \(e^{\lambda_2} < 1\), and \(e^{\lambda_2 t} \to 0\) as \(t \to \infty\). Hence, we have the following approximations:

\[
C^*(t) = \left(1 + \frac{1}{\lambda_1 T_l}\right) c_1 y_1 e^{\lambda_1 t},
\]

and

\[
D^*(t) = \left(\frac{\alpha}{\lambda_1 T_l}\right) c_1 y_1 e^{\lambda_1 t},
\]

which are accurate for large \(t\).

Now, we can explain why \(R_0\) (or any other quantity) alone cannot predict how \(C(t)\) and \(D(t)\) grow. It is clear that \(C(t)\) is determined by \(R_E, R_I, T_E, T_I, E(1), I(1), V(1), D(1)\), collectively, and \(D(t)\) is determined by \(R_E, R_I, T_E, T_I, \alpha, E(1), I(1), D(1)\), collectively.

### 3.4 A Case Study

In this section, we apply our method to a case study.

The parameters in our model are set as follows. The incubation period means the time between catching the virus and beginning to have symptoms of the disease. Most estimates of the incubation period for COVID-19 range from 1–14 days, most commonly around five days. We set \(D_E = 5\) [15, 16]. Using available preliminary data, the median time from onset to clinical recovery is approximately 2 weeks for mild cases and 3–6 weeks for severe or critical cases. We set \(D_I = 20\) [17, 23].

For the same data in Section 2.2 for the world, we get

\[
C_{World}^*(t) = 29539.4430255 \times 1.1155783^t,
\]

and

\[
D_{World}^*(t) = 1454.9810657 \times 1.1155783^t,
\]

where we set \(R_E = 1.39\), \(R_I = 0.50\), and \(\alpha = 0.157\). The initial values are \(E(1) = 20,000\), \(I(1) = 10,000\), \(V(1) = 20,000\), \(D(1) = 2,000\).
Table 2: Quality of Analytical Model

| $t$ | Analytical $C_{\text{World}}(t)$ | Actual $C_{\text{World}}(t)$ | ARE | Analytical $D_{\text{World}}(t)$ | Actual $D_{\text{World}}(t)$ | ARE |
|-----|---------------------------------|-------------------------------|-----|---------------------------------|-------------------------------|-----|
| 21  | 296654                          | 304979                        | -0.970% | 13865                          | 13011                        | 2.026% |
| 22  | 330674                          | 337459                        | -0.790% | 15518                          | 14640                        | 2.083% |
| 23  | 368612                          | 378830                        | -1.190% | 17366                          | 16513                        | 2.024% |
| 24  | 410921                          | 422574                        | -1.358% | 19431                          | 18894                        | 1.274% |
| 25  | 458107                          | 471035                        | -1.506% | 21738                          | 21282                        | 1.082% |
| 26  | 510736                          | 531865                        | -2.462% | 24314                          | 24073                        | 0.572% |
| 27  | 569438                          | 596366                        | -3.137% | 27190                          | 27343                        | -0.363% |
| 28  | 634914                          | 663127                        | -3.287% | 30402                          | 30861                        | -1.089% |
| 29  | 707949                          | 723390                        | -1.799% | 33986                          | 34065                        | -0.187% |
| 30  | 789417                          | 784738                        | 0.545%  | 37988                          | 37773                        | 0.510% |
| 31  | 880294                          | 858377                        | 2.553%  | 42453                          | 42146                        | 0.728% |

In Table 2, we display the analytical data, the actual data, and the ARE for $C_{\text{World}}(t)$ and $D_{\text{World}}(t)$ during March 21–31, 2020, with $\text{ARE}_i$ in the range $[-3.287\%, 2.553\%]$ for $C_{\text{World}}(t)$, and $\text{ARE}_i$ in the range $[-1.089\%, 2.083\%]$ for $D_{\text{World}}(t)$.

4 Future Prediction

Using our analytical results in Sections 2 and 3, we display predictions of $C_{\text{World}}(t)$ and $D_{\text{World}}(t)$ on April $t$, 2020 in Figures 2 and 3 respectively. The regression analysis method gives higher predictions than the analytical model method, since the analytical model results in a smaller base of exponentiation.

From a global perspective, unless powerful and effective social and medical impacts are made, by the end April of 2020, the cumulative number of confirmed cases is 23.333 and 36.068 millions respectively using regression analysis and analytical model, and the cumulative number of dead cases is 1.148 and 2.528 millions respectively using regression analysis and analytical model, based on the current situation.
Figure 2: Prediction of $C_{\text{World}}(t)$ on April $t$, 2020.
Figure 3: Prediction of $D_{\text{World}}(t)$ on April $t$, 2020.
5 Conclusions

We have made some progress towards analytical expressions of the daily growth of the cumulative number of confirmed cases and the cumulative number of dead cases, two most important and daily reported figures. Our analytical methods and results have been tested using the COVID-19 epidemic data in the world, and proven to be effective and accurate. We have also predicted the cumulative number of confirmed cases and the cumulative number of dead cases in April 2020 using our models and methods.

Declarations

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