Research Article

The Impact of Media Coverage and Curfew on the Outbreak of Coronavirus Disease 2019 Model: Stability and Bifurcation

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In this study, the spreading of the pandemic coronavirus disease (COVID-19) is formulated mathematically. The objective of this study is to stop or slow the spread of COVID-19. In fact, to stop the spread of COVID-19, the vaccine of the disease is needed. However, in the absence of the vaccine, people must have to obey curfew and social distancing and follow the media alert coverage rule. In order to maintain these alternative factors, we must obey the modeling rule. Therefore, the impact of curfew, media alert coverage, and social distance between the individuals on the outbreak of disease is considered. Five ordinary differential equations of the first-order are used to represent the model. The solution properties of the system are discussed. The equilibria and the basic reproduction number are computed. The local and global stabilities are studied. The occurrence of local bifurcation near the disease-free equilibrium point is investigated. Numerical simulation is carried out in applying the model to the sample of the Iraqi population through solving the model using the Runge–Kutta fourth-order method with the help of Matlab. It is observed that the complete application of the curfew and social distance makes the basic reproduction number less than one and hence prevents the outbreak of disease. However, increasing the media alert coverage does not prevent the outbreak of disease completely, instead of that it reduces the spread, which means the disease is under control, by reducing the basic reproduction number and making it an approachable one.

1. Introduction

The most considerable species in habitat are viruses. Since they cannot reproduce themselves, they are known as parasites. Some of these viruses, when they are replicated, cause dangerous diseases in the host, whether humans or animals. One of these species is the coronavirus (COVID-19). It is a disease caused by a new coronavirus called SARS-CoV-2. An outbreak of pneumonia caused by a novel coronavirus (COVID-19) began on 31 December 2019, in Wuhan, China. This pandemic that rapidly spread over 215 countries as of March 2020 continues to hit heavy public health and socio-economic burden in many parts of the world, including in Iraq. From January 22, 2020, up to March 3, 2021, it was reported that over 115 million confirmed cases and about 2,565,310 deaths around the world [1].

The experience in China showed that the use of complete curfew and the speed diagnosis of the infected individuals have a strong impact on the dynamics of the epidemic [2]. It is well known that there is a long history of applying mathematical models in epidemiology to understand and control the outbreak of disease. Carcione et al. [3] implemented an SEIR model to compute the COVID-19 infected population and the number of casualties of this epidemic. Eikenberry et al. [4] developed a compartmental model for assessing the community-wide impact of masks used in the US states of New York and Washington by the general, asymptomatic public, a portion of which may be asymptptomatically infectious. Maji [5] formulated a mathematical model for the COVID-19 outbreak by introducing a quarantine class with media-induced fear in the disease transmission rate to analyze the dynamic behavior of this epidemic. Asamoah et al. [6] developed a deterministic
model to study the transmission dynamics of the disease with two categories of the susceptibles: Immigrant Susceptibles and Local Susceptible. Peter et al. [7], proposed a new mathematical model to investigate the recent outbreak of coronavirus disease (COVID-19) in Pakistan. Das et al. [8] formulated a mathematical model with comorbidity to study the transmission dynamics as well as an optimal control-based framework to diminish COVID-19. Musa et al. [9] employed an SEIR model to study the transmission dynamics of SARS-CoV-2 outbreaks in Nigeria; the model incorporates a different group of populations, namely, high and moderate risk populations. They used the model to investigate the influence of each population on the overall transmission dynamics. Recently, Paul et al. [10] proposed a simple population dynamics solution based on the incidence-fitness relationship in predicting that a plateau or steady-state of SARS-CoV-2 will be reached using the basic concept of geometry. Many other researchers have been proposed and studied mathematical models to describe the dynamic of COVID-19 (see [11–14] and the references therein).

In view of the above, in this paper, however, a mathematical model that describes the spreading of COVID-19 is proposed and studied. It is assumed that the population is divided into five compartments including susceptible, exposed, infected, hospitalized infected, and recovered. The impact of curfew, media alert coverage, and social distance as the infected individuals increase, the susceptible individuals reduce their contact with other individuals to avoid the disease from spreading completely, it is assumed that \( b_1 \geq b_2 \).

Recall that the interaction functions between the compartments given by the right-hand side of the system (1) are continuous and have continuous partial derivatives. Therefore, they are Lipschitz functions, and hence system (1) has a unique solution.

**Theorem 1.** All solutions of system (1) which initiate in \( \mathbb{R}^5_+ \) are uniformly bounded.

**Proof.** Let \( (S(t), E(t), I(t), I_h(t), R(t)) \) be any solution of system (1) that initiates at the nonnegative value \( (S(0), E(0), I(0), I_h(0), R(0)) \).

Consider the function \( N(t) = S(t) + E(t) + I(t) + I_h(t) + R(t) \).

By taking the derivative of \( N(t) \) with respect to the time along with the solution of system (1), it is obtained that

\[
\frac{dN}{dt} = \frac{dS}{dt} + \frac{dE}{dt} + \frac{dI}{dt} + \frac{dI_h}{dt} + \frac{dR}{dt}.
\]
of system (1): solution values of system instead of system (1) and then substitute the so-
equations of system (1), hence one can solve the following
covered individuals, does not appear in the first four
Note that since the variable
3. Existence of the Equilibrium Points and Basic Reproduction Number

Table 1: The parameters description.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>$A$</td>
<td>The recruitment rate</td>
</tr>
<tr>
<td>$c$</td>
<td>The probability of individuals responding to the curfew</td>
</tr>
<tr>
<td>$b_1$</td>
<td>The direct contact rate between $S$ and $I$ in the absence of media alerts</td>
</tr>
<tr>
<td>$b_2$</td>
<td>The direct contact rate between $S$ and $I$ in the existence of media alerts</td>
</tr>
<tr>
<td>$m$</td>
<td>Nonresponse rate of the individuals to the media alerts</td>
</tr>
<tr>
<td>$p$</td>
<td>The rate of incubation termination and transition from uninfected to infectious individuals</td>
</tr>
<tr>
<td>$q$</td>
<td>The fraction of the exposed individuals move to the infected compartment, while the rest $(1-q)$ moves to the hospitalized infected population</td>
</tr>
<tr>
<td>$r_1$</td>
<td>The recovery rate due to natural immune response</td>
</tr>
<tr>
<td>$r_2$</td>
<td>The recovery rate due to treatment</td>
</tr>
<tr>
<td>$v$</td>
<td>The fraction of the nonrecovered individuals who start suffering severe symptoms move to the hospitalized infected population</td>
</tr>
<tr>
<td>$d_0$</td>
<td>The natural death rate</td>
</tr>
<tr>
<td>$d_1$</td>
<td>The death rate of disease in the infectious compartment</td>
</tr>
<tr>
<td>$d_2$</td>
<td>The death rate due to disease in the hospitalized infected population</td>
</tr>
</tbody>
</table>

Then, using direct computation gives that

\[
\frac{dN}{dt} = A - d_0 S - d_0 E - (d_0 + d_1) I - (d_0 + d_2) I_h - d_0 R. \tag{3}
\]

Accordingly, it is obtained that

\[
\frac{dN}{dt} \leq A - d_0 N, \tag{4}
\]

which implies that $N(t) \leq A/d_0$, as $t \rightarrow \infty$. Hence, all the solutions of system (1) are uniformly bounded, and therefore we finished the proof.

According to system (1), the feasible region of it can be written as follows:

\[
\Omega = \{ (S,E,I,I_h,R) \in \mathbb{R}^5_+ : 0 \leq S + E + I + I_h + R \leq \frac{A}{d_0} \}. \tag{5}
\]

Clearly, system (7) has two equilibrium points: the disease-free equilibrium point and the endemic equilibrium point. The disease-free equilibrium point is given by $\rho_0 = (S_0, 0, 0, 0)$, where

\[
S_0 = \frac{A}{d_0}. \tag{8}
\]

Since the basic reproduction number is computed at this equilibrium point, hence in the following the computation of this number is carried out, and then the endemic equilibrium point is determined.

Now, in order to determine the basic reproduction number, the "next-generation method" or "Spectral Radius method" is used (see [16, 17]). Consider an epidemic model having $n$ different compartments from which $m$ compartments contained infected individuals with the disease, then the next-generation matrix (operator) is given by $(FV)^{-1}$, where

\[
F = \left[ \frac{\partial F_i(\rho_0)}{\partial x_j} \right] \quad \text{and} \quad V = \left[ \frac{\partial V_i(\rho_0)}{\partial x_j} \right], \tag{9}
\]

where $i,j = 1,2, \ldots, m$, $\rho_0$ is the disease-free equilibrium point, while $x_i$ denotes the number of individuals in the $i^{th}$ infected compartment. However, $F_i(x_i)$ is the rate of
The reproduction number, denoted by $R_0$, is determined as $V_1$, which represents the rate of the shift of individuals into the $i$th compartment by all other means and $V_1^*$ denotes the rate of the shift of individuals out of the $i$th compartment. The difference $F_i(x_i) - V_i(x_i)$ gives the rate of change of $x_i$. Note that $F_i$ should include only infections that are newly arising but does not include terms that describe the shift of infectious individuals from one infected compartment to another. Finally, the basic reproduction number, that is denoted by $R_0$, is given by the spectral radius (dominant eigenvalue) of the matrix $FV^{-1}$. It is well known that the basic reproduction number $R_0$ is one of the most crucial quantities in infectious diseases as $R_0$ measures how contagious a disease is. For $R_0 < 1$, the disease is expected to stop spreading, but for $R_0 = 1$, an infected individual can infect on average 1 person; that is, the spread of the disease is stable. The disease can spread and become epidemic if $R_0 > 1$.

Accordingly, regarding system (7), it is obtained that
\[
F = \begin{bmatrix}
0 & (1 - c)b_1S_0 & 0 \\
0 & 0 & 0 \\
0 & 0 & 0
\end{bmatrix},
\]
\[
V = \begin{bmatrix}
B_1 & 0 & 0 \\
-qp & B_2 & 0 \\
-(1 - q)p & -c & (1 - r_1)B_3
\end{bmatrix},
\]

where $B_1 = p + d_0$, $B_2 = r_1 + v(1 - r_1) + (d_0 + d_1)$, and $B_3 = r_2 + (d_0 + d_2)$. Then, we get
\[
V^{-1} = \frac{1}{B_1B_2B_3} \begin{bmatrix}
B_2B_3 & 0 & 0 \\
qp(1 - r_1) + (1 - q) B_2 & B_1B_3 & 0 \\
qp & -(1 - r_1)B_1 & B_1B_2
\end{bmatrix}
\]

Therefore,
\[
FV^{-1} = \frac{1}{B_1B_2B_3} \begin{bmatrix}
qp(1 - c)b_1S_0B_3 & (1 - c)b_1S_0B_1B_3 & 0 \\
0 & 0 & 0 \\
0 & 0 & 0
\end{bmatrix}.
\]

Consequently, the basic reproduction number of system (7) is determined as
\[
R_0 = \frac{qp(1 - c)b_1S_0B_1}{B_1B_2B_3}.
\]

Now, the endemic equilibrium point of system (7) can be determined by $\rho_1 = (S^*, E^*, I^*, I_b^*)$, where
\[
S^* = \frac{A - B_1E^*}{d_0},
\]
\[
E^* = \frac{qpE^*}{B_2},
\]
\[
I^* = \frac{(pB_2 - qpB_1)E^*}{B_2B_3},
\]
\[
I_b^* = \frac{(pB_2 - qpB_1)E^*}{B_2B_3},
\]

where $B_1 = r_1 + d_0 + d_1$, while $E^*$ is a positive root of the second-order polynomial equation:
\[
\alpha_1E^2 + \alpha_2E + \alpha_3 = 0,
\]
where $\alpha_1 = - (1 - c)(b_1 - b_2)B_1 (qp)^2 < 0$, $\alpha_2 = qp[(1 - c)(b_1 - b_2)Aqp - (1 - c)b_1mB_1B_3 - B_1B_2]$, and $\alpha_3 = MB_2 [(1 - c)b_1qpA - B_1B_2d_0] = B_1B_2d_0mB_2 [R_0 - 1]$.

Therefore, equation (11) has a unique positive root, given by $E^* = \alpha_3/2\alpha_1 - 1/2\alpha_1 \sqrt{\alpha_2^2 - 4\alpha_1\alpha_3}$, if and only if the following condition holds:
\[
R_0 > 1.
\]

Moreover, in order to obtain a positive endemic equilibrium point $\rho_1$ for system (7), we should have
\[
E^* < \frac{A}{B_1}.
\]

4. Local Stability Analysis

This section treats the local stability of system (7) using the linearization technique. The Jacobian matrix for system (7) at the point $(S, E, I, I_b)$ can be written as follows:
where the symbols $b_i, i = 1, 2, 3,$ are given in equation (10). Therefore, the local stability analysis for the disease-free equilibrium point can be studied in the following theorem.

**Theorem 2.** The disease-free equilibrium point $(\rho_0)$ of system (7) is locally asymptotically stable provided the following condition holds:

$$R_0 < 1.$$  \hfill (19)

**Proof.** According to equation (18), the Jacobian matrix at the point $(\rho_0)$ is given by

$$J(\rho_0) = \begin{bmatrix} -d_0 & -(1 - c)b_1 S_0 & 0 \\ 0 & -B_1 & -(1 - c)b_1 S_0 \\ 0 & q \rho & -B_2 \\ 0 & (1 - q) p & \nu (1 - r_1) - B_3 \end{bmatrix}.$$  \hfill (20)

Direct computations show that this Jacobian matrix has the following characteristic equation:

$$(-d_0 - \lambda)(-B_1 - \lambda)(\lambda^2 - T \lambda + D) = 0,$$  \hfill (21)

where $T = -(B_1 + B_2) < 0$ and $D = B_1 B_2 - (1 - c)q \rho b_1 S_0 = B_1 B_2 |1 - R_0|$. Therefore, all the eigenvalues of the matrix $J(\rho_0)$ are real and negative if and only if condition (19) holds. Moreover, since the other two eigenvalues $\lambda_1 = -d_0$ and $\lambda_4 = -B_3$ are negative, hence the disease-free equilibrium point $(\rho_0)$ of system (7) is locally asymptotically stable under condition (19). However, if condition (19) is revised, then $\rho_0$ is an unstable point. Thus, this completes the proof.

According to theorem (6), the basic reproduction number represents the vital value for controlling the outbreak of COVID-19, so that the disease will disappear provided that the basic reproduction number is less than one while the COVID-19 will outbreak when the basic reproduction number is greater than one.

**Theorem 3.** The endemic equilibrium point $(\rho_1)$ of system (7) is locally asymptotically stable provided that

$$b_{22} b_{33} > b_{23} b_{32},$$  \hfill (22)

where $b_{ij}, i, j = 1, 2, 3,$ are given in the proof.

**Proof.** Substituting the endemic equilibrium point $(\rho_1)$ in the Jacobian matrix given by equation (18) gives that

$$J(\rho_1) = \begin{bmatrix} -(1 - c) b_1 (b_2 - (2m + 1)l) - d_0 & -b_1 (b_2 - (2m + 1)l) - d_0 \\ -(1 - c) b_1 (b_2 - (2m + 1)l) + b_1 (b_2 - (2m + 1)l) - d_0 & 0 \\ 0 & q \rho \nu (1 - r_1) - B_3 \\ 0 & (1 - q) p & \nu (1 - r_1) - B_3 \end{bmatrix}.$$  \hfill (23)

where $b_{11} = -(1 - c) b_1 (b_2 - (2m + 1)l) - d_0$, $b_{12} = -(1 - c) b_1 (b_2 - (2m + 1)l) + b_1 (b_2 - (2m + 1)l) - d_0$, $b_{13} = -(1 - c) b_1 (b_2 - (2m + 1)l) + b_1 (b_2 - (2m + 1)l) - d_0$, $b_{14} = -(1 - c) b_1 (b_2 - (2m + 1)l) - d_0$, $b_{22} = b_1 (b_2 - (2m + 1)l) + b_1 (b_2 - (2m + 1)l) - d_0$, $b_{23} = b_1 (b_2 - (2m + 1)l) + b_1 (b_2 - (2m + 1)l) - d_0$, $b_{24} = b_1 (b_2 - (2m + 1)l) - d_0$, $b_{33} = b_1 (b_2 - (2m + 1)l) + b_1 (b_2 - (2m + 1)l) - d_0$, $b_{34} = b_1 (b_2 - (2m + 1)l) - d_0$, $b_{44} = b_1 (b_2 - (2m + 1)l) - d_0$, and $b_{44} = b_1 (b_2 - (2m + 1)l) - d_0$.

Hence, the characteristic equation of this Jacobian matrix can be written as

$$(b_{44} - \lambda)[A_1 \lambda^2 + A_2 \lambda + A_3] = 0,$$  \hfill (24)

where $A_1 = -[b_{11} + b_{12} + b_{13}]$, $A_2 = b_{11} b_{12} + b_{11} b_{13} + b_{12} b_{13} - b_{11} b_{12} b_{13}$, and $A_3 = -[b_{11} b_{12} b_{13} + b_{12} b_{13} - b_{11} b_{12} b_{13}]$.

While $\Delta = A_1 A_2 - A_3 = A_1 [b_{11} + b_{12} + b_{13} - b_{11} b_{12} b_{13}]$, accordingly, by substituting the values of $b_{ij}$ and then using condition (22) along with the existence condition (16), we obtain that $A_1, A_2, and \Delta$ are positive. Therefore, all the roots (eigenvalues) of the third-order polynomial equation in (24) have negative real parts. Furthermore, since the fourth eigenvalue $\lambda_4 = b_{44} - \lambda < 0$, then the proof is complete.

From theorem (7), it concluded that the COVID-19 will be endemic and the disease is out of control if in addition to the basic reproduction number greater than one, condition (22) is satisfied.

\[ \square \]

5. Global Stability

In this section, the stability analysis of system (7) is studied using the Lyapunov function in order to obtain the global stability conditions or the basin of attraction for each equilibrium point. Therefore, the following theorem discusses the global stability of the free-disease equilibrium point.

**Theorem 4.** Assume that free-disease equilibrium point $\rho_0 = (A/d_0, 0, 0, 0)$ is locally asymptotically stable, then it is globally asymptotically stable.

**Proof.** Consider the following positive definite real valued function:

$$V(\rho) = \frac{1}{2} (\rho_1 - \rho_0)^T J(\rho_0) (\rho_1 - \rho_0).$$

where $J(\rho_0)$ is the Jacobian matrix at the free-disease equilibrium point $\rho_0$. Then, the time derivative of $V(\rho)$ along the solutions of system (7) is given by

$$\dot{V}(\rho) = (\rho_1 - \rho_0)^T J(\rho_0) \dot{\rho}.$$
\[ V_0(S, E, I_h) = \left( S - S_0 - S_0 \ln \frac{S}{S_0} \right) + E + I + I_h. \]  

\[ \left( 1 - c \right) \left( b_1 - \frac{b_2 I}{m + I} \right) IS + \left( 1 - c \right) \left( b_1 - \frac{b_2 I^*}{m + I^*} \right) I' S^* , \]

\[ + \left( 1 - c \right) \left( b_1 - \frac{b_2 I^*}{m + I^*} \right) IS - B_1 \left( E - E^* \right) - \left( 1 - c \right) \left( b_1 - \frac{b_2 I^*}{m + I^*} \right) I' S^* , \]

\[ + \left( I - I^* \right) \left[ \rho \left( E - E^* \right) - B_2 \left( I - I^* \right) \right] , \]

\[ + \left( I - I^* \right) \left[ \rho \left( E - E^* \right) + \nu \left( 1 - r_2 \right) \left( I - I^* \right) \right] - B_5 \left( I_h - I_{h}^* \right) , \]

\[ = -\pi_{11} \left( S - S^* \right)^2 - \pi_{12} \left( S - S^* \right) \left( E - E^* \right) - B_1 \left( E - E^* \right)^2 , \]

\[ - \pi_{13} \left( S - S^* \right) \left( I - I^* \right) - B_2 \left( I - I^* \right)^2 , \]

\[ - \pi_{23} \left( E - E^* \right) \left( I - I^* \right) - \pi_{24} \left( E - E^* \right) \left( I_h - I_{h}^* \right) , \]

\[ + \pi_{34} \left( I - I^* \right) \left( I_h - I_{h}^* \right) - B_3 \left( I_h - I_{h}^* \right)^2 . \]
where \( \pi_{11} = d_0 + (1 - c)I^*/(m + I^*)(m + I^*) \) \( m b_1 (1 + I) + \) \( (b_1 - b_2)(m + I^*) \), \( \pi_{12} = (1 - c)I^*/(m + I^*)(m + I^*) \) \( m b_1 \) \( (1 + I) + (b_1 - b_2)(m + I^*) \), \( \pi_{13} = (1 - c)(m + I^*)(m + I^*) \) \( m \) \( (b_1 - b_2)(m + I^*) \), \( \pi_{23} = (1 - c) I^* \), \( \pi_{24} = (1 - q) p \), and \( \pi_{25} = \nu (1 - r_i) \).

Accordingly, by using conditions (27a)–(27e), we obtain that

\[
\frac{dV}{dt} > -\frac{1}{2} \left[ \sqrt{\pi_{11}} (S - S^*) + \sqrt{\pi_{12}} (E - E^*) \right]^2 - \frac{1}{2} \left[ \sqrt{\pi_{13}} (S - S^*) + \sqrt{\pi_{14}} (I - I^*) \right]^2,
\]

\[
\frac{dV}{dt} > -\frac{1}{2} \left[ \sqrt{\pi_{15}} (S - S^*) + \sqrt{\pi_{16}} (T - T^*) \right]^2 - \frac{1}{2} \left[ \sqrt{\pi_{17}} (S - S^*) + \sqrt{\pi_{18}} (T - T^*) \right]^2.
\]  

where the functions \( f_i, i = 1, 2, 3, 4 \), are the interaction functions given on the right-hand side of system (7). Accordingly, the second derivative of \( F \) with respect to \( X \) can be written as

\[
D^2 F(X)(V, V) = \left( \begin{array}{c}
-2 (1 - c) \left( b_1 - b_2 \frac{(m + I^*)}{(m + I)^2} \right) v_1 v_3 + (1 - c) \frac{2 m^2 b_2}{(m + I)^2} S v_3^2 \\
2 (1 - c) \left( b_1 - b_2 \frac{(m + I^*)}{(m + I)^2} \right) v_1 v_3 - (1 - c) \frac{2 m^2 b_2}{(m + I)^2} S v_3^2 \\
0 \\
0
\end{array} \right),
\]  

where \( V = (v_1, v_2, v_3, v_4)^T \) be any nonzero vector. Therefore, the local bifurcation condition at the disease-free equilibrium point is established by the following theorem.

**Theorem 6.** System (7) exhibits a transcritical bifurcation near the disease-free equilibrium point \( \rho_0 \), where \( R_0 = 1 \) or \( b_1 = B_1 B_2 / q p (1 - c) S_0 (\equiv b_1^*) \).

**Proof.** Straightforward computation shows that the Jacobian matrix of system (7) at \( (\rho_0, \ b_1^*) \) can be written as

\[
J_0 = J(\rho_0, \ b_1^*) = \left( \begin{array}{cccc}
-d_0 & 0 & (1 - c)b_1^* S_0 & 0 \\
0 & -B_1 & (1 - c)b_1^* S_0 & 0 \\
0 & q p & -B_2 & 0 \\
0 & (1 - q) p & \nu (1 - r_i) & -B_3
\end{array} \right).
\]

Therefore, the eigenvalues of \( J_0 \) can be written as

\[
\begin{align*}
J_0 &= \left( \begin{array}{cccc}
-d_0 & 0 & (1 - c)b_1^* S_0 & 0 \\
0 & -B_1 & (1 - c)b_1^* S_0 & 0 \\
0 & q p & -B_2 & 0 \\
0 & (1 - q) p & \nu (1 - r_i) & -B_3
\end{array} \right).
\end{align*}
\]
Accordingly, the disease-free equilibrium point becomes the nonhyperbolic point. Now, define \( \lambda_i = -d_i \), where \( i = 1, 2, 3, 4 \). Define \( \xi \) as the eigenvector corresponding to the eigenvalue \( \lambda_3 = 0 \). Direct computation gives that
\[
U_1 = (\xi_1, x_1, x_2, x_3, x_4, x_5)^T,
\]
where \( \eta_i = -b_iB_i / q \delta d_i < 0 \), \( \eta_3 = B_i / q \delta > 0 \), and \( \eta_3 = 1/2(B_i [(1-q)B_i/q + v(1-r_1)] > 0 \), with \( x_3 \) be any nonzero real number.
\[
U_2 = (\eta_4, y_1, y_2, y_3, y_4, y_5)^T,
\]
where \( \eta_4 = q \delta / B_i > 0 \), with \( y_3 \) be any nonzero real number.

Define \( \xi \) as the eigenvector corresponding to the eigenvalue \( \lambda_3 = 0 \) of the matrix \( J_i \). Direct computation gives that
\[
U_1^T D^2 F(\rho_0, b^*_i)(U_1, U_1) = 2(1-c)\eta_4
\]
where \( \eta_i = \lambda_i x_i \), \( \eta_3 = 1/2(B_i [(1-q)B_i/q + v(1-r_1)] > 0 \), with \( x_3 \) be any nonzero real number.

It is observed that although system (7) becomes a nonhyperbolic point near the endemic equilibrium point \( \rho_1 \), as \( p = \xi_2 / (\xi_1 + d_0) \), \( \xi_2 = \xi_2 d_0 / (\xi_1 + d_0) (p^* + d_0) > 0 \), and \( \xi_3 = 1/2(B_i [(1-q)B_i/q + v(1-r_1)] > 0 \), with \( x_3 \) and \( x_4 \) are any nonzero real numbers. Moreover, it is obtained that
\[
H_1^T D^2 F(\rho_0, b^*_i)(H_1, H_1) = 0.
\]
Hence,
\[
\frac{\partial F}{\partial b_i} = F_b = \begin{pmatrix}
-(1-c)S_i \\
(1-c)S_i \\
0 \\
0 \\
0 \\
0
\end{pmatrix}, \quad \Rightarrow F_b(\rho_0, b^*_i) = \begin{pmatrix}
0 \\
0 \\
0 \\
0
\end{pmatrix}.
\]
Then, it is obtained that \( U_1^T D F(\rho_0, b^*_i) = 0 \). Furthermore, it is observed that
\[
D F(\rho_0, b^*_i) = \begin{pmatrix}
0 & 0 & -1-c & 0 & 0 \\
0 & 0 & 1-c & 0 & 0 \\
0 & 0 & 0 & 0 & 0 \\
0 & 0 & 0 & 0 & 0
\end{pmatrix}.
\]
Therefore, \( U_1^T D F(\rho_0, b^*_i)U_1 = (1-c)S_0 \eta_4 x_1 y_3 \neq 0 \). However, substituting the values of \( U_1 \) in equation (32) instead of \( V \) and then computing \( D^2 F(\rho_0, b^*_i) \) give that
\[
U_1^T D^2 F(\rho_0, b^*_i)(U_1, U_1) = 2(1-c)\eta_4
\]
Accordingly, any type of the local bifurcations in the sense of Sotomayor cannot occur. \( \Box \)

7. Numerical Simulation

To complete the analysis of the proposed system, it should be simulated numerically with real and reasonable parameter values and then the obtained results are validated with those obtained in the previous sections. It is well known that the model parameters affect the dynamical behavior of the model greatly. This occurs due to either error in their measurement or missing the parameter values. Accordingly, the impact of varying the model’s parameters on the behavior of the solution of the model is important. Therefore, the parameters that are having vital roles in the dynamics of the model will be given clear attention to understand their effects on the control of the spread of disease. Most of the parameter values used in this section are taken from the World Health Organization reports and adopted by many references as stated in Table 2. Since the population of Iraq is nonhomogenous distributed, therefore, we will confine ourselves to apply the proposed model on a sample that exists in a specific region to get more accurate results.

For the parameter values used in Table 2, system (1) is solved, and the trajectories are obtained as shown in Figure 1.
parameter \( m \), that \( m = 0, 100, 200, 300 \), respectively, the trajectories of system (1) are drawn in Figure 2.

It is clear from Figure 2 that with the increase in the value of \( m \), the trajectory still settled at endemic but the populations of exposed, infected, and hospitalized infected increased too.

Now, for the data given in Table 2 with \( c = 0.9 \), it is observed that the basic reproduction number decreases to \( R_0 = 0.6193 \) and the trajectory of system (1) is shown in Figure 3. It is clear from Figure 3 that when \( R_0 < 1 \), system (1) approaches asymptotically to disease-free equilibrium point and that confirms the theoretical results in previous sections.

On the other hand, increasing the parameter \( b_1 \) from \( b_1 = 0.001 \) to \( b_1 = 0.01 \) with data used in Figure 3 makes the trajectory of system (7) return back to settle at the endemic equilibrium point as shown in Figure 4.

Clearly, from Figure 4, the bifurcation occurs in system (1) and the trajectory of system (1) transfers from disease-free point to endemic point as the bifurcation parameter \( b_1 \) increases, passing through the value \( b_1^* \) and then becomes \( b_1 = 0.01 \). It is noted that the basic reproduction number increases from the value \( R_0 = 0.6193 \) to the value \( R_0 = 6.192717 \) passing the bifurcation value \( R_0^* = 1 \). Now, for the data given in Table 2 with reducing the value of \( q \) from the value \( q = 0.5 \) to the value \( q = 0.2 \), the basic reproduction number decreases from the value \( R_0 = 2.4771 \) to the value \( R_0 = 0.9908 \), and the trajectory of system (1) is shown in Figure 5.

Again, Figure 5 ensures the obtained results regarding the stability analysis of the disease-free equilibrium point. Finally, other parameters are also varying to study their effects on the dynamical behavior of system (1), and it is

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Range</th>
<th>Value used</th>
<th>Source</th>
</tr>
</thead>
<tbody>
<tr>
<td>( A )</td>
<td>30–1000 day(^{-1})</td>
<td>50</td>
<td>[6]</td>
</tr>
<tr>
<td>( c )</td>
<td>[0, 1]</td>
<td>0.5</td>
<td>Assumed</td>
</tr>
<tr>
<td>( b_1 )</td>
<td>( 6.038 \times 10^{-8} )–0.8158 day(^{-1})</td>
<td>0.001</td>
<td>[6, 11, 12]</td>
</tr>
<tr>
<td>( b_2 )</td>
<td>(0, ( b_1 )) day(^{-1})</td>
<td>0.00001</td>
<td>[15]</td>
</tr>
<tr>
<td>( m )</td>
<td>[0, ( 1/2 )]</td>
<td>10</td>
<td>[15]</td>
</tr>
<tr>
<td>( p )</td>
<td>[1/14–1/3] day(^{-1})</td>
<td>0.1</td>
<td>[4, 13, 14]</td>
</tr>
<tr>
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<td>(0, 1)</td>
<td>0.5</td>
<td>Assumed</td>
</tr>
<tr>
<td>( r_1 )</td>
<td>[1/14–1/3] day(^{-1})</td>
<td>0.08</td>
<td>[4, 13, 14]</td>
</tr>
<tr>
<td>( r_2 )</td>
<td>0.05–0.9 day(^{-1})</td>
<td>0.1</td>
<td>[6]</td>
</tr>
<tr>
<td>( \nu )</td>
<td>0.01–0.5 day(^{-1})</td>
<td>0.3</td>
<td>Assumed</td>
</tr>
<tr>
<td>( d_0 )</td>
<td>0.000042–0.011 day(^{-1})</td>
<td>0.01</td>
<td>[6, 11, 12]</td>
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<tr>
<td>( d_1 )</td>
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<td>0.001</td>
<td>[6, 11, 12]</td>
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<tr>
<td>( d_2 )</td>
<td>0.0044–0.1850 day(^{-1})</td>
<td>0.01</td>
<td>[6, 11, 12]</td>
</tr>
</tbody>
</table>

Figure 1: The trajectory of system (1) approaches to endemic equilibrium point.
Figure 2: The trajectory of system (1), for the data given in Table 1 with different values of $m$, approaches to endemic point. The blue color for $m = 0$, green color for $m = 100$, red color for $m = 200$, and cyan color for $m = 300$.

Figure 3: The trajectory of system (1) approaches to disease-free equilibrium point.
observed that quantitative effects exist rather than qualitative one so that the population size of the different compartments in the system changes slightly.

8. Discussion and Conclusion

In this paper, an epidemic model to describe the spread of COVID-19 is proposed using the first-order ordinary differential equations. The model takes into account the impact of the curfew and media coverage on the control of the spreading of disease. It is observed that model (1) has two equilibrium points: the disease-free point that is stable, which means the disease is under control from a biological point of view, when the basic reproduction number satisfies $R_0 < 1$ and unstable for $R_0 > 1$, and the endemic equilibrium point that is stable, which means the disease out of control from a biological point of view, whenever the disease-free becomes unstable. All the properties of the solution of the model are studied. The local bifurcation near the disease-free point is investigated too. An application of the proposed model on the sample of Iraqi population of size 500 has been done using baseline values of the parameters of system

Figure 4: The trajectory of system (1) approaches to endemic equilibrium point for the data in Table 2 with $c = 0.9$ and $b_1 = 0.01$.

Figure 5: The trajectory of system (1) approaches to disease-free equilibrium point for the data in Table 2 with $q = 0.2$. 
(1) as suggested by World Health Organization reports and adopted by many researchers, and the following observations are obtained.

The model has only one type of attractor, approaches either to disease-free equilibrium point or to the endemic equilibrium point, which means either stop the outbreak of COVID-19 or spread the COVID-19, depending on $R_0 < 1$ and $R_0 > 1$, respectively. The model is solved numerically using the Runge-Kutta fourth-order method with the help of Matlab. The model is applied to a sample of the Iraqi population using the parameter value ranges as suggested by the World Health Organization reports and adopted by many references as stated in Table 2. It is observed that increasing the probability of an individual’s response to the curfew leads to reducing the basic reproduction number below one and then controlling the spread of COVID-19, and hence system (1) approaches asymptotically to the disease-free point. However, increasing the direct contact rate between $S$ and $I$ in the absence of media alerts or decreasing the social distance between the population individuals leads to an increase in the basic reproduction number above one, and then an increase in the outbreak of the disease occurred. Hence, the trajectory of system (1) approaches the endemic point. Although increasing the response rate of the individuals to the media alerts (that means decreasing $m$) cannot prevent the disease from spreading completely, it leads to reduce it up to a minimum rate. A similar result is observed by increasing the parameter $b_2$, which increases the term $(b_2/(m + I))$ that reduces the value of the transmission rate when infectious individuals appear and are reported. Finally, decreasing the fraction of the exposed individuals moving to the infected compartment, which leads to increasing the number of individuals moving to hospitalized infected populations, can prevent the disease from spreading and the system approaches a disease-free equilibrium point.

Data Availability

The data used to support the findings of this study are included within the article.

Conflicts of Interest

The authors declare that there are no conflicts of interest.

References


