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Analysis, predicting, and controlling the COVID-19 pandemic in Iraq through SIR model

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A R T I C L E I N F O

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A B S T R A C T

Using the standard SIR model with three unknown biological parameters, the COVID-19 pandemic in Iraq has been studied. The least squares method and real data on confirmed infections, deaths, and recoveries over a long time (455 days) were used to estimate these parameters. In this regards, first, we find the basic reproductive number R_0 is 0.9422661124 which indicates and predicts that the COVID-19 pandemic in Iraq will gradually subside until it is eradicated permanently with time. Additionally, we develop an optimal vaccination strategy with the goal of reducing COVID-19 infections and preventing their spread in Iraq, thereby putting a clear picture of control this pandemic.

1. Introduction

Since its emergence over two years ago, the COVID-19 pandemic has overtaxed the ability of healthcare organizations throughout much of the world, affecting virtually every aspect of daily life. In fact, the last days of 2019 brought an unwelcome surprise: the first pandemic of the century. Wuhan, the capital of the Chinese province of Hubei, was the first place that COVID-19 appeared, and then it spread from Wuhan to around the world. This necessitated pooling all of the world's resources to combat the looming pandemic and halt its spread in any way possible, as most countries closed their borders and airports and prohibited citizens from roaming freely for fear of the disease spreading further. During this time, major pharmaceutical companies worldwide rushed to produce vaccines, and the FDA approved some vaccines for emergency use, including those manufactured by Pfizer, Moderna, and AstraZeneca [\[1\]](#page-13-0).

In recent years, the mathematical modeling of infectious diseases has become an active and important area of research being carried out. Because infectious diseases have dynamic behaviors, mathematical epidemiology can help scientists to better understand how they behave and what they can expect in the future. In fact, mathematical models are usually implemented to compare and evaluate various detection, prevention, therapy, and control programs, as well as to plan, implement, evaluate, and optimize these programmers throughout their lifecycle. In this vein, many studies have been published in recent years that have been adopted for the creation and evaluation of epidemiological models, and many of these contain significant findings [[2–](#page-13-1)[4\]](#page-13-2). Since the new pandemic's emergence, a slew of mathematical models has given health officials in many countries some useful insights into the most effective ways to stop the disease from spreading [[5–](#page-13-3)[9\]](#page-13-4).

Recently, the mathematical models provide future insight into the qualitative behavior of COVID-19 and give a prediction of how this disease will behave [\[5,](#page-13-3)[10–](#page-13-5)[15\]](#page-13-6). Some scholars in Ontario, Canada, offered a mathematical model to examine the impact of a variety of public interventions on COVID-19 behavior and how to reduce it [\[16\]](#page-13-7). Depending on nonlinear differential equations, Fanelli and Piazza have discussed the temporal models for COVID-19 infection in three countries: China, Italy, and France, based on the real dates of certain days [[17\]](#page-13-8). While Khan and Atangana used the COVID-19 confirmed infection reports in Wuhan to develop

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Fig. 1. SIR model flowchart.

and analyze a fractional model by using fractional derivatives [[6\]](#page-13-9). In the absence of any available vaccine or treatment at that time, Atangana studied effects of curfew on population to mitigate the infection of COVID-19 [[18\]](#page-13-10). Also, Sahin and Sahin introduced a fractional model depending on the cumulative daily report of COVID-19 in three countries: Italy, UK, and USA, to study the transmission of this disease [[19\]](#page-13-11). In Mexico, some authors used artificial neural networks to perform and predict the transmission model of COVID-19 [\[20](#page-13-12)]. Also, Based on real data from Ghana, many models for COVID-19 have been investigated, as the reader can see in [[21–](#page-13-13)[24\]](#page-13-14). Some scholar investigate the impact of self-isolation, quarantine, and contact tracing on the stopping and control of the COVID-19 pandemic using data from the New York population [[25\]](#page-13-15). On the other hand, fractional differential equations play a fundamental role in the distribution of the COVID-19 pandemic [[26,](#page-13-16)[27\]](#page-13-17), so there are numerous methods to solve them [[28–](#page-14-0)[31\]](#page-14-1).

In this study, we use the standard SIR to describe COVID-19 transmission in Iraq in order to help the Iraqi Ministry of Health develop a vaccination strategy. To begin, we used real data from confirmed infections over a period of 455 days to develop the model. After that, a sensitivity analysis is performed to see how parameters affect R_0 . To reduce the number of infected people and develop an optimal vaccination strategy, we constructed a quadratic optimal control problem. We hope that the strategy outlined in this paper will help predict the long-term trend of COVID-19 and reduce the number of infected people.

2. The SIR model and parameters fitting results

Now, the following standard SIR model will be use to study COVID-19 pandemic in Iraq.

$$
\frac{dX_S}{dt} = A - \frac{\beta X_I(t)X_S(t)}{(X_S(t) + X_I(t) + X_R(t))} - dX_S(t),
$$
\n
$$
\frac{dX_I}{dt} = \frac{\beta X_I(t)X_S(t)}{(X_S(t) + X_I(t) + X_R(t))} - (\gamma + \epsilon + d)X_I(t),
$$
\n
$$
\frac{dX_R}{dt} = \gamma X_I(t) - dX_R(t),
$$
\n(2.1)

where the biological parameters Λ , β , d , γ , and ϵ are represent the birth rate, the disease transmission rate, the natural death rate, the recovery rate, and the disease death rate respectively (see [Fig.](#page-1-0) [1\)](#page-1-0).

Now, we adopt the use of model parameter estimation to examine the model's validity and reliability by comparing and fitting the SIR model with actual data. This will show the accuracy and the ability of the considered model to predict real results. The country of Iraq will be used as an example in this section. Also, the daily reports of COVID-19 by the WHO and the Iraqi Ministry of Health will be considered for this purpose. To give a complete picture of this subject, we collected data on cases of confirmed infection, death, and recovery for a long period of 455 days starting from January 1, 2021, to March 31, 2022.

According to Iraq demographics reported by U.N. [[32\]](#page-14-2) the life expectancy at birth of Iraqi people, both sexes, is 71.08 year, and the estimated number of population of Iraqi people is approximately $N(0)$ = 41828817. Therefore the natural death rate parameter d and the birth rate Λ of Iraqi population in the disease absence are $d = 1/(71.08 \times 365)$ per day and $\Lambda = d \times N(0) = 1635$ per day.

It is well known that the key to designing any epidemiological study is to figure out how many biological parameters to use in it and how to estimate them. In fact, in this study, we try to implement the stander SIR model with three unknown parameters to

Fig. 2. Fitted diagram of susceptible individuals in Iraq and the SIR model.

study the behavior of COVID-19 pandemic in Iraq. Thus, the main effort of this work is to focus on how to estimate these parameters to ensure the validity of the suggested SIR model. For this purpose, the least squares method will be used.

As a practical matter, we will use days as the time unit, and the observed collected data on the confirmed infected cases *̃*, death cases \tilde{D} , and recovered cases \tilde{R} for a long period of 455 days. The first step is to use these observed collected data to compute the number of susceptible people according to the following relationship:

$$
\tilde{X}_S(t) = N(t) - \tilde{X}_I(t) - \tilde{X}_R(t) - dN(t), \quad t = 0, 1, 2, \dots,
$$
\n(2.2)

where $N(t)$ is defined by

$$
N(t) = (1 - d) N(t - 1) + \Lambda, \quad t = 1, 2, 3, \dots
$$
\n
$$
(2.3)
$$

Now, we are ready to apply the least squares method to estimate the value of β , γ and ϵ . Therefore, we will construct the objective function, which is the sum of square errors, as follows:

$$
E(\beta, \gamma, \epsilon) = \sum_{\kappa=0}^{M} (X_S(\kappa) - \tilde{X}_{S\kappa})^2 + (X_I(\kappa) - \tilde{X}_{I\kappa})^2 + (X_R(\kappa) - \tilde{X}_{R\kappa})^2,
$$
\n(2.4)

where M is the total number of days with available observed data in our study, which is 455. Also, $X_S(\kappa)$, $X_I(\kappa)$, and $X_R(\kappa)$ are the estimated values at the day κ , which satisfied the Eq. ([2.1\)](#page-1-1). In order to determine the unknown biological parameters, we must first solve the optimization problems listed below:

Minimize
$$
E(\beta, \gamma, \epsilon)
$$

\nS.T. $Eq.(2.1)$ (2.5)

In fact, we will use two Matlab packages, ode45 (Runge–Kutta methods) and lsqcurvefit (least-square curve fitting), to perform the above task. The optimal fit to available collected data of susceptible individuals via $X_S(t)$ of our SIR model is illustrated in [Fig.](#page-2-0) [2](#page-2-0) and the corresponding estimated parameters are $\beta = 0.0953$, $\gamma = 0.0162$, and $\epsilon = 0.0849$.

3. Stability analysis of the SIR model

In order to make predictions and have a clear image about the future behavior of the Covid-19 pandemic in Iraq, we will focus in this section on studying the behavior of the model (2.1) (2.1) near the equilibrium points as time increases. First, we solve the next equations

$$
A - \frac{\beta X_I X_S}{(X_S + X_I + X_R)} - dX_S = 0,
$$

\n
$$
\frac{\beta X_I X_S}{(X_S + X_I + X_R)} - (\gamma + \epsilon + d)X_I = 0,
$$

\n
$$
\gamma X_I - dX_R = 0.
$$
\n(3.1)

Now, we can deduce that the model (2.1) (2.1) possesses two equilibrium points as follows:

1. The equilibrium point free of disease E_0 which given by:

$$
E_0 = (X_{S_0}, X_{I_0}, X_{R_0}) = (\frac{A}{d}, 0, 0). \tag{3.2}
$$

2. The endemic equilibrium point E_e , which given by:

$$
E_e = (X_{S_e}, X_{I_e}, X_{R_e}) = (\frac{A - (\gamma + d + \varepsilon)X_I}{d}, X_I, \frac{\gamma X_I}{d}).
$$
\n(3.3)

Here we emphasize that the compounds of point E_0 are free of X_I , unlike the compounds of point E_e that contain X_I which they were named.

3.1. The basic reproductive number R_0

In this subsection, we will look into the vital threshold and chief quantity, well-known as the basic reproduction number, which is commonly abbreviated as R_0 . In fact, R_0 performs a critical function in the disease-free local stability as documented by [[33\]](#page-14-3).

Now, calculate R_0 , where R_0 is the eigenvalue of the matrix $G = FV^{-1}$, where F indicates new infections, while V indicates the transmission of infection from one place to another. Both are calculated in an equilibrium-free equilibrium state and are thus derived as follows.

Infectious compartments are found in the system ([2.1\)](#page-1-1) as follows.

$$
\frac{dX_S}{dt} = A - \frac{\beta X_I(t)X_S(t)}{(X_S(t) + X_I(t) + X_R(t))} - dX_S(t),
$$
\n
$$
\frac{dX_I}{dt} = \frac{\beta X_I(t)X_S(t)}{(X_S(t) + X_I(t) + X_R(t))} - (\gamma + \epsilon + d)X_I(t).
$$
\n(3.4)

Let $\Omega = [X_I, X_S]^T$. Then the system [\(3.4\)](#page-3-0) can be written as

$$
{}_{0}^{C}D_{t}^{\alpha}\Omega=\tilde{F}(\Omega)-\tilde{V}(\Omega),
$$
\n(3.5)

where

$$
\tilde{F}(\Omega) = \begin{bmatrix} \frac{\beta X_I(t)X_S(t)}{(X_S(t) + X_I(t) + X_R(t))} \\ 0 \end{bmatrix},\tag{3.6}
$$

and

$$
\tilde{V}(\Omega) = \begin{bmatrix} (\gamma + \epsilon + d)X_I(t) \\ -A + \frac{\beta X_I(t)X_S(t)}{(X_S(t) + X_I(t) + X_R(t))} + dX_S(t) \end{bmatrix},
$$
\n(3.7)

Therefore, we perform the calculation of the matrices F and V at the point E_0 in the following forms:

$$
F = \begin{bmatrix} 0 & \beta \\ 0 & 0 \end{bmatrix},\tag{3.8}
$$

and

$$
V = \begin{bmatrix} 0 & \gamma + d + \epsilon \\ d & \beta \end{bmatrix},\tag{3.9}
$$

Matrix V is inversed as follows

$$
V^{-1} = \begin{bmatrix} -\frac{\beta}{(\gamma + d + \epsilon)} & \frac{1}{d} \\ \frac{1}{\gamma + d + \epsilon} & 0 \end{bmatrix} . \tag{3.10}
$$

Now, by multiplying matrices ([3.8\)](#page-3-1) and [\(3.10\)](#page-3-2) we will get

$$
G = \begin{bmatrix} \frac{\beta}{\gamma + d + \epsilon} & 0 \\ 0 & 0 \end{bmatrix} . \tag{3.11}
$$

Since, the eigenvalues of the matrix ([3.11](#page-3-3)) are $\lambda_1 = 0$, and $\lambda_2 = \frac{\beta}{\gamma + d + \epsilon}$, we have

$$
R_0 = \frac{\beta}{\gamma + d + \epsilon}.\tag{3.12}
$$

It is important to note that when R_0 less than one, disease in the population is eradicated and infection is eliminated. Furthermore, when R_0 is greater than one, the disease will continue to exist in the population. Therefore, based on the estimated COVID-19 pandemic parameters in Iraq for the available time period starting from January 1, 2021, to March 31, 2022, we find that $R_0 = 0.9422661124$, indicating that the COVID-19 pandemic in Iraq will gradually reduce until it is permanently ended with the passage of time.

Table 1

3.2. A sensitivity analysis of R_0

Due to the fact that R_0 is an extremely biologically significant quantity that plays a chief role in the spread of any pandemic, investigating the sensitivity of R_0 is very interesting and crucial to the elimination and effective control of the disease. R_0 sensitivity to changes in parameter Y is represented by this index. So, we can calculate the changes in all parameters in the formula of R_0 by using the partial derivatives as follows:

Now, we give the following relationship that describes the R_0 forward sensitivity index with respect to the parameter Y :

$$
\Theta_Y^{R_0} = \left(\frac{\partial R_0}{\partial Y}\right) \left(\frac{Y}{R_0}\right),\tag{3.13}
$$

where Y is a parameter to describe the basic reproductive number R_0 . It is well known that a negative (positive) index means that any increase in the parameter Y leads to a decrease (increase) in R_0 [\[34](#page-14-4)]. The sensitivity indices with respect to the parameters can be given β , ϵ , and γ respectively, by the basic reproductive number mentioned in the Eq. [\(3.12\)](#page-3-4), as follows:

$$
\frac{\partial R_0}{\partial \beta} \frac{\beta}{R_0} = \frac{1}{\gamma + d + \epsilon} \frac{\beta}{R_0},
$$

\n
$$
= \frac{1}{\gamma + d + \epsilon} \frac{\beta}{\frac{\beta}{\gamma + d + \epsilon}} = 1.
$$

\n
$$
\frac{\partial R_0}{\partial \epsilon} \frac{\epsilon}{R_0} = \frac{-\beta}{(\gamma + d + \epsilon)^2} \frac{\epsilon}{\frac{\beta}{\gamma + d + \epsilon}},
$$

\n
$$
= \frac{-\epsilon}{\gamma + d + \epsilon} = -0.8394374916.
$$

\n
$$
\frac{\partial R_0}{\partial \gamma} \frac{\gamma}{R_0} = \frac{-\beta}{(\gamma + d + \epsilon)^2} \frac{\gamma}{\frac{\beta}{\gamma + d + \epsilon}},
$$

\n
$$
= \frac{-\gamma}{\gamma + d + \epsilon} = -0.1601753518.
$$

In fact, [Table](#page-4-0) [1](#page-4-0) describes and explains the R_0 sensitivity indices to biological parameters for the considered model, as determined using the estimated parameter values computed in Section [2,](#page-1-2) where the parameters are listed in decreasing order of sensitivity. The result demonstrated that when the disease transmission rate parameter is increased while the other parameters are kept fixed, the value of R_0 is increased, which means that we will experience an endemicity of the disease more since it has positive indices. On the other hand, the increase of the parameters γ , and ϵ will lead to a decrease in the value of R_0 which means that we will probably reduce the chance of spreading the disease more since these parameters have negative indices.

4. Optimal control on the model

In order to design an optimal vaccination strategy to minimize the number of COVID-19 infections and prevent its spread in Iraq, we reformulate the SIR model in Eq. [\(2.1](#page-1-1)) by imposing a control variable that represents the effect of the vaccine. Also, we will take into consideration the number of days, $\tau = 14$ days, that the vaccine will start working actively as in [Fig.](#page-5-0) [3](#page-5-0) and the following time-delay differential equations:

$$
\frac{dX_S}{dt} = A - \frac{\beta X_I(t)X_S(t)}{(X_S(t) + X_I(t) + X_R(t))} - dX_S(t) - u(t - \tau)X_S(t - \tau),
$$
\n
$$
\frac{dX_I}{dt} = \frac{\beta X_I(t)X_S(t)}{(X_S(t) + X_I(t) + X_R(t))} - (\gamma + \epsilon + d)X_I(t),
$$
\n
$$
\frac{dX_R}{dt} = \gamma X_I(t) - dX_R(t) + u(t - \tau)X_S(t - \tau).
$$
\n(4.1)

In addition, the cost function is constructed as follows:

$$
\mathfrak{F}(u(t)) = \int_0^{t_f} (D_1 X_I(t) + \frac{D_2}{2} u^2(t)) dt,
$$
\n(4.2)

where D_1 and D_2 are the balancing cost factors and t_f is the specific time. From this point on, our primary goal is to optimally figure out the most effective vaccination controls so that we can limit the speed of the spread of this epidemic, reduce mortality, and

Fig. 3. Time delay SIR model flowchart.

avoid unnecessary complications. As a result, our task will be to find the control variable $u(t)$ between zero and one that minimizes $\Im(u(t))$ under the constraints of Eq. [\(4.1](#page-4-1)).

In other words, we want to find the optimal vaccination controls, $u^*(t)$ that satisfied:

 $\mathfrak{F}(u^*(t)) = min\{\mathfrak{F}(u(t)) : 0 \le u(t) \le 1, \forall t \in [0, t_f] \}$ such that the state equations are satisfied *}*.

Therefore, we will use the minimum principle of Pontryagin, which is modified in [[35\]](#page-14-5), to find the optimal control solution for our problem that have delays in control and state variables.

For this purpose, we define the following Hamiltonian function:

$$
H = D_1 X_I(t) + \frac{D_2}{2} u^2(t) + \sum_{\kappa=1}^3 p_1(t) f_\kappa(X_S(t), X_I(t), X_R(t), U(t-\tau), X_S(t-\tau)),\tag{4.3}
$$

where f_k denotes the right side of the κ^{th} equation's SIR model.

Now, according the minimum principle of Pontryagin in [[35\]](#page-14-5), we have the optimal control $u^*(t)$ by

$$
u^*(t) = \max\{0, \min\{1, \frac{p_1^+(t) - p_2^+(t)}{D_2}h_{[0,t_f - \tau]}(t)X_S^*(t)\}\}, \ \forall t \in [0, t_f],
$$
\n(4.4)

where

$$
\hbar_{[0,T_f-\tau]}(t) = \begin{cases} 1, & t \in [0,t_f-\tau] \\ 0, & \text{otherwise} \end{cases} \tag{4.5}
$$

Also, the co-state variables $p_1(t)$, $p_2(t)$ and $p_3(t)$ at the optimal solution $X_S^*(t)$, $X_I^*(t)$, $X_R^*(t)$ and $u^*(t)$ satisfy the following differential equations:

$$
\frac{dp_1(t)}{dt} = d p_1(t) + \frac{\beta(p_1(t) - p_2(t))}{N^*(t)} X_I^*(t) + \hbar_{[0,t_f - \tau]}(t)(p_1(t) - p_3(t))u^*(t),
$$
\n
$$
\frac{dp_2(t)}{dt} = -D_1 + \frac{\beta(p_1(t) - p_2(t))}{N^*(t)} X_S^*(t) + (\varepsilon + d + \gamma)p_2(t) - \gamma p_3(t),
$$
\n
$$
\frac{dp_2(t)}{dt} = \gamma p_3(t),
$$
\n(4.6)

with terminal transversality conditions, $p_1(t_f) = 0$, $p_2(t_f) = 0$, and $p_3(t_f) = 0$. Also, $p_k^+(t) = p_k(t + \tau)$, for $\kappa = 1, 2, 3$.

To find the optimal vaccination strategy, we construct the following algorithm based on applying the forward and backward Euler method to solve state Eq. [\(4.1](#page-4-1)) and co-state equations Eq. ([4.6](#page-5-1)), respectively, and on the optimal control law in Eq. ([4.4\)](#page-5-2).

Algorithm 4.1.

Step 1 *Insert the values of the biological parameters* Γ , *d*, β , γ , ϵ , and τ . Also, insert the initial conditions of $X_S(\kappa) = X_S(0), X_I(\kappa) = \kappa$ $X_I(0), X_R(\kappa) = X_R(0)$ for all $\kappa = -M, -M + 1, \ldots, 0$ and terminal conditions $p_1(\kappa), p_2(\kappa), p_3(\kappa) = 0$, for all $\kappa = N, N + 1, \ldots, N + M$.

Fig. 4. The susceptible $X_S(t)$ without and with control. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

Step 2 *Suppose the time interval is* $[0, t_f]$ *and compute the step size* $h = \frac{t_f}{N} = \frac{\tau}{M}$ *, where N and M* are positive integer numbers.

Step 3 *Set* $u(\kappa h) = 0$ *, for all* $\kappa = -M, -M + 1, \ldots, 0, 1, \ldots, N$.

Step 4 *For all* $\kappa = 1, 2, ..., N$, compute $X_S(\kappa h)$, $X_I(\kappa h)$, and $X_R(\kappa h)$ by applying forward Euler method as follows:

$$
X_{S}((\kappa+1)h) = X_{S}(\kappa h) + h(A - \frac{\beta X_{I}(\kappa h)X_{S}(kh)}{(X_{S}(\kappa h) + X_{I}(\kappa h) + X_{R}(\kappa h))} - d X_{S}(\kappa h)),
$$

\n
$$
- u(h(\kappa - M))X_{S}(h(\kappa - M))
$$

\n
$$
X_{I}((\kappa+1)h) = X_{I}(\kappa h) + h(\frac{\beta X_{I}(\kappa h)X_{S}(\kappa h)}{(X_{S}(\kappa h) + X_{I}(\kappa h) + X_{R}(\kappa h))} - (\gamma + \epsilon + d)X_{I}(\kappa h)),
$$

\n
$$
X_{R}((\kappa+1)h) = X_{R}(\kappa h) + h(\gamma X_{I}(\kappa h) - d X_{R}(\kappa h)) + u(h(\kappa - M))X_{S}(h(\kappa - M)).
$$

Step 5 *For all* $\kappa = N - 1, N - 2, ..., 0$, compute $p_1(\kappa h), p_2(\kappa h)$, and $p_3(\kappa h)$ by applying backward Euler method as follows:

$$
p_1(\kappa h) = p_1((\kappa + 1)h) - h(dp_1((\kappa + 1)h) + \frac{\beta(p_1((\kappa + 1)h) - p_2((\kappa + 1)h))}{(X_S((\kappa + 1)h) + X_I((\kappa + 1)h) + X_R((\kappa + 1)h))}
$$

\n
$$
\times X_I((\kappa + 1)h) + h_{[0,t_{f^{-T}}]}((\kappa + 1)h)(p_1((\kappa + 1)h) - p_3(t))u((\kappa + 1)h)),
$$

\n
$$
p_2(\kappa h) = p_2((\kappa + 1)h) - h(-D_1 + \frac{\beta(p_1((\kappa + 1)h) - p_2((\kappa + 1)h))}{(X_S((\kappa + 1)h) + X_I((\kappa + 1)h) + X_R((\kappa + 1)h))}
$$

\n
$$
\times X_S((\kappa + 1)h) + (\epsilon + d + \gamma)p_2((\kappa + 1)h) - \gamma p_3((\kappa + 1)h)),
$$

\n
$$
p_2(\kappa h) = p_2((\kappa + 1)h) - h\gamma p_3((\kappa + 1)h).
$$

Step 6 *Apply the optimal control law to compute* $u(kh)$ *for all* $k = 1, 2, ..., N$ *as follows:*

$$
u(\kappa h) = \max\{0, \min\{1, \frac{p_1(\kappa h + M) - p_2(\kappa h + M)}{D_2} h_{[0,t_f - \tau]}(\kappa h) X_S^*(\kappa h)\}\}.
$$

Step 7 *If the stopping criterion (the absolute value of optimal control of the current and the previous iterations) is held, then the algorithm ends, else return to Step 4.*

5. Numerical simulation of optimal vaccination strategy

This section focuses on introducing a numerical simulation of an optimal vaccination strategy by solving the optimal control problem that was constructed in the previous section. Indeed, the optimal control solution was calculated using the estimated parameters, the [Algorithm](#page-5-3) [4.1,](#page-5-3) and Maple2020 software. The results appear to show the impact of the vaccination process in controlling and preventing the outbreak of this pandemic in Iraq. We show that if Iraq's healthcare system followed the vaccination process depicted in [Fig.](#page-7-0) [7](#page-8-0), the number of infected people would decrease as depicted in Fig. [6](#page-7-0), while the number of vaccinated and recovered people would increase as depicted in [Fig.](#page-7-1) [5.](#page-7-1)

Fig. 5. The recovery $X_R(t)$ without and with control. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

Fig. 6. The infections $X_I(t)$ without and with control. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

6. Conclusions

We use SIR model with three unknown parameters to study the behavior of COVID-19 pandemic in Iraq. In fact, there are two main parts of our work: The first is using the real data on cases of confirmed infection, death, and recovery for a long period of 455 days starting from January 1, 2021, to March 31, 2022 (see Appendix) to estimation these three unknown parameters as in Section [2.](#page-1-2) Also, we compute the basic reproductive number R_0 and study the sensitive analysis of each effective parameters. In fact, R_0 indicates and predicts that the COVID-19 pandemic in Iraq will gradually subside until it is eradicated permanently with time. The

Fig. 7. Optimal strategy of vaccination $u^*(t)$.

second is to design an optimal vaccination strategy to minimize the number of COVID-19 infections by reformulating the SIR model in Eq. ([2.1\)](#page-1-1) by imposing a control variable that represents the effect of the vaccine. Also, we will take into consideration the number of days, $\tau = 14$ days, that the vaccine will start working actively. Then the minimum principle of Pontryagin is implemented to find the optimal vaccination strategy and prevent its spread in Iraq. For this purpose, the results show the impact of the vaccination process on controlling and preventing the spread of COVID-19 in Iraq.

To make this analysis more reliable, we used the data from January 1, 2021, as the initial values. Also, to predict the behavior of the COVID-19 pandemic in Iraq, we documented and drew some figures, see [Figs.](#page-6-0) [4–](#page-6-0)[6.](#page-7-0) Indeed, the blue curve in these figures clarifies the number of susceptible, infected, and recovered individuals, which agree with the total number of recovered individuals until 15 May 2022, when it is 2299985. The red curve in [Figs.](#page-6-0) [4–](#page-6-0)[6](#page-7-0) shows that if the vaccination process depicted in [Fig.](#page-8-0) [7](#page-8-0) is followed, the number of susceptible, infected individuals will decrease while the cumulative number of recovered individuals will increase. Many changes can be made to improve this optimal vaccination strategy, such as selecting the balancing cost factors D_1 and D_2 based on the requirements of the Iraqi Ministry of Health or any other beneficiary. In addition, we can account for the effectiveness of any vaccine by assuming the new control variable, $u(t) = \rho v(t)$, where ρ is the vaccine efficacy. In this simulation, $D_1 = 0.001$ and $D_2 = 100$ are used. Also, we assume that the vaccine efficacy is 100%. We hope that this paper will assist the Iraqi Ministry of Health in predicting and controlling the COVID-19 pandemic. In this case, the number of infected people would decrease, as depicted in [Fig.](#page-7-0) [6,](#page-7-0) while the number of vaccinated and recovered people would increase, as depicted in [Fig.](#page-7-1) [5.](#page-7-1)

Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Data availability

No data was used for the research described in the article.

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Appendix

References

- [1] Craig AM, Hughes BL, Swamy GK. Coronavirus disease 2019 vaccines in pregnancy. Amer J Obstetrics Gynecol MFM 2021;3(2):100295. [http://dx.doi.](http://dx.doi.org/10.1016/j.ajogmf.2020.100295) [org/10.1016/j.ajogmf.2020.100295](http://dx.doi.org/10.1016/j.ajogmf.2020.100295).
- [2] Anderson RM, May RM. Population biology of infectious diseases: Part I. Nature 1979;280(5721):361–7. [http://dx.doi.org/10.1038/280361a0.](http://dx.doi.org/10.1038/280361a0)
- [3] Anderson RM, Lee PA. Infectious diseases of humans. OUP Oxford; 1992, URL [https://www.ebook.de/de/product/3246618/roy_m_anderson_pamela_](https://www.ebook.de/de/product/3246618/roy_m_anderson_pamela_anderson_lee_infectious_diseases_of_humans.html) [anderson_lee_infectious_diseases_of_humans.html](https://www.ebook.de/de/product/3246618/roy_m_anderson_pamela_anderson_lee_infectious_diseases_of_humans.html).
- [4] [Diekmann O, Heesterbeek J. Mathematical epidemiology of infectious diseases: Model building, analysis and interpretation. Wiley series in mathematical](http://refhub.elsevier.com/S2666-7207(23)00016-4/sb4) [and computational biology, United States: John Wiley and Sons; 2000.](http://refhub.elsevier.com/S2666-7207(23)00016-4/sb4)
- [5] [Kucharski AJ, Russell TW, Diamond C, Liu Y, Edmunds J, Funk S, et al. Early dynamics of transmission and control of COVID-19: A mathematical modelling](http://refhub.elsevier.com/S2666-7207(23)00016-4/sb5) [study. Lancet Infect Dis 2020;20\(5\):553–8.](http://refhub.elsevier.com/S2666-7207(23)00016-4/sb5)
- [6] Liu Y, Gayle AA, Wilder-Smith A, Rocklöv J. The reproductive number of COVID-19 is higher compared to SARS coronavirus. J Travel Med 2020;27(2). http://dx.doi.org/10.1093/itm/taaa021.
- [7] Ivorra B, Ferrández M, Vela-Pérez M, Ramos A. Mathematical modeling of the spread of the coronavirus disease 2019 (COVID-19) taking into account the undetected infections. The case of China. Commun Nonlinear Sci Numer Simul 2020;88:105303. <http://dx.doi.org/10.1016/j.cnsns.2020.105303>.
- [8] Tuan NH, Mohammadi H, Rezapour S. A mathematical model for COVID-19 transmission by using the Caputo fractional derivative. Chaos Solitons Fractals 2020;140:110107. [http://dx.doi.org/10.1016/j.chaos.2020.110107.](http://dx.doi.org/10.1016/j.chaos.2020.110107)
- [9] Al-Saedi HM, Hameed HH. Mathematical modeling for COVID-19 pandemic in Iraq. J Interdiscipl Math 2021;24(5):1407–27. [http://dx.doi.org/10.1080/](http://dx.doi.org/10.1080/09720502.2021.1923943) [09720502.2021.1923943.](http://dx.doi.org/10.1080/09720502.2021.1923943)
- [10] [Lenhart S, Workman JT. Optimal control applied to biological models. Chapman and Hall/CRC; 2007.](http://refhub.elsevier.com/S2666-7207(23)00016-4/sb10)
- [11] Asamoah JKK, Okyere E, Abidemi A, Moore SE, Sun G-Q, Jin Z, et al. Optimal control and comprehensive cost-effectiveness analysis for COVID-19. Results Phys 2022;33:105177. [http://dx.doi.org/10.1016/j.rinp.2022.105177.](http://dx.doi.org/10.1016/j.rinp.2022.105177)
- [12] Asamoah JKK, Bornaa C, Seidu B, Jin Z. Mathematical analysis of the effects of controls on transmission dynamics of SARS-CoV-2. Alex Eng J 2020;59(6):5069–78. [http://dx.doi.org/10.1016/j.aej.2020.09.033.](http://dx.doi.org/10.1016/j.aej.2020.09.033)
- [13] Asamoah JKK, Jin Z, Sun G-Q. Non-seasonal and seasonal relapse model for Q fever disease with comprehensive cost-effectiveness analysis. Results Phys 2021;22:103889. [http://dx.doi.org/10.1016/j.rinp.2021.103889.](http://dx.doi.org/10.1016/j.rinp.2021.103889)
- [14] Faniran TS, Ali A, Al-Hazmi NE, Asamoah JKK, Nofal TA, Adewole MO. New variant of SARS-CoV-2 dynamics with imperfect vaccine. In: Selişteanu D, editor. Complexity 2022;2022:1–17. <http://dx.doi.org/10.1155/2022/1062180>.
- [15] Khalaf SL, Kadhim MS, Khudair AR. Studying of COVID-19 fractional model: Stability analysis. Part Diff Equ Appl Math 2023;7:100470. [http://dx.doi.](http://dx.doi.org/10.1016/j.padiff.2022.100470) [org/10.1016/j.padiff.2022.100470](http://dx.doi.org/10.1016/j.padiff.2022.100470).
- [16] Wu J, Tang B, Bragazzi NL, Nah K, McCarthy Z. Quantifying the role of social distancing, personal protection and case detection in mitigating COVID-19 outbreak in Ontario, Canada. J Math Ind 2020;10(1). [http://dx.doi.org/10.1186/s13362-020-00083-3.](http://dx.doi.org/10.1186/s13362-020-00083-3)
- [17] Fanelli D, Piazza F. Analysis and forecast of COVID-19 spreading in China, Italy and France. Chaos Solitons Fractals 2020;134:109761. [http://dx.doi.org/](http://dx.doi.org/10.1016/j.chaos.2020.109761) [10.1016/j.chaos.2020.109761](http://dx.doi.org/10.1016/j.chaos.2020.109761).
- [18] Atangana A. Modelling the spread of COVID-19 with new fractal-fractional operators: Can the lockdown save mankind before vaccination? Chaos Solitons Fractals 2020;136:109860. <http://dx.doi.org/10.1016/j.chaos.2020.109860>.
- [19] Şahin U, Şahin T. Forecasting the cumulative number of confirmed cases of COVID-19 in Italy, UK and USA using fractional nonlinear grey Bernoulli model. Chaos Solitons Fractals 2020;138:109948. [http://dx.doi.org/10.1016/j.chaos.2020.109948.](http://dx.doi.org/10.1016/j.chaos.2020.109948)
- [20] Torrealba-Rodriguez O, Conde-Gutiérrez R, Hernández-Javier A. Modeling and prediction of COVID-19 in Mexico applying mathematical and computational models. Chaos Solitons Fractals 2020;138:109946. <http://dx.doi.org/10.1016/j.chaos.2020.109946>.
- [21] Asamoah JKK, Owusu MA, Jin Z, Oduro F, Abidemi A, Gyasi EO. Global stability and cost-effectiveness analysis of COVID-19 considering the impact of the environment: Using data from Ghana. Chaos Solitons Fractals 2020;140:110103. [http://dx.doi.org/10.1016/j.chaos.2020.110103.](http://dx.doi.org/10.1016/j.chaos.2020.110103)
- [22] Asamoah JKK, Jin Z, Sun G-Q, Seidu B, Yankson E, Abidemi A, et al. Sensitivity assessment and optimal economic evaluation of a new COVID-19 compartmental epidemic model with control interventions. Chaos Solitons Fractals 2021;146:110885. <http://dx.doi.org/10.1016/j.chaos.2021.110885>.
- [23] Acheampong E, Okyere E, Iddi S, Bonney JH, Asamoah JKK, Wattis JA, et al. Mathematical modelling of earlier stages of COVID-19 transmission dynamics in Ghana. Results Phys 2022;34:105193. [http://dx.doi.org/10.1016/j.rinp.2022.105193.](http://dx.doi.org/10.1016/j.rinp.2022.105193)
- [24] Moore SE, Nyandjo-Bamen HL, Menoukeu-Pamen O, Asamoah JKK, Jin Z. Global stability dynamics and sensitivity assessment of COVID-19 with timely-delayed diagnosis in Ghana. Comput Math Biophys 2022;10(1):87–104. [http://dx.doi.org/10.1515/cmb-2022-0134.](http://dx.doi.org/10.1515/cmb-2022-0134)
- [25] Ngonghala CN, Iboi E, Eikenberry S, Scotch M, MacIntyre CR, Bonds MH, et al. Mathematical assessment of the impact of non-pharmaceutical interventions on curtailing the 2019 novel coronavirus. Math Biosci 2020;325:108364. [http://dx.doi.org/10.1016/j.mbs.2020.108364.](http://dx.doi.org/10.1016/j.mbs.2020.108364)
- [26] Addai E, Zhang L, Asamoah JKK, Preko AK, Arthur YD. Fractal–fractional age-structure study of Omicron SARS-CoV-2 variant transmission dynamics. Partial Differ Equ Appl Math 2022;6:100455. [http://dx.doi.org/10.1016/j.padiff.2022.100455.](http://dx.doi.org/10.1016/j.padiff.2022.100455)
- [27] Addai E, Zhang L, Preko AK, Asamoah JKK. Fractional order epidemiological model of SARS-CoV-2 dynamism involving Alzheimer's disease. Healthc Anal 2022;2:100114. <http://dx.doi.org/10.1016/j.health.2022.100114>.
- [28] Khudair AR. On solving non-homogeneous fractional differential equations of Euler type. Comput Appl Math 2013;32(3):577–84. [http://dx.doi.org/10.](http://dx.doi.org/10.1007/s40314-013-0046-2) [1007/s40314-013-0046-2](http://dx.doi.org/10.1007/s40314-013-0046-2).
- [29] Khudair AR, Haddad S, khalaf SL. Restricted fractional differential transform for solving irrational order fractional differential equations. Chaos Solitons Fractals 2017;101:81–5. [http://dx.doi.org/10.1016/j.chaos.2017.05.026.](http://dx.doi.org/10.1016/j.chaos.2017.05.026)
- [30] Khalaf SL, Khudair AR. Particular solution of linear sequential fractional differential equation with constant coefficients by inverse fractional differential operators. Differ Equ Dyn Syst 2017;25(3):373–83. <http://dx.doi.org/10.1007/s12591-017-0364-8>.
- [31] Jalil AFA, Khudair AR. Toward solving fractional differential equations via solving ordinary differential equations. Comput Appl Math 2022;41(1). <http://dx.doi.org/10.1007/s40314-021-01744-8>.
- [32] Iraq Population (1950 - 2020). <https://www.worldometers.info/world-population/iraq-population/>.
- [33] [van den Driessche P, Watmough J. Reproduction numbers and sub-threshold endemic equilibria for compartmental models of disease transmission. Math](http://refhub.elsevier.com/S2666-7207(23)00016-4/sb33) [Biosci 2002;180\(1–2\):29–48.](http://refhub.elsevier.com/S2666-7207(23)00016-4/sb33)
- [34] Chitnis N, Hyman JM, Cushing JM. Determining important parameters in the spread of malaria through the sensitivity analysis of a mathematical model. Bull Math Biol 2008;70(5):1272–96. <http://dx.doi.org/10.1007/s11538-008-9299-0>.
- [35] Göllmann L, Kern D, Maurer H. Optimal control problems with delays in state and control variables subject to mixed control-state constraints. Optim Control Appl Methods 2009;30(4):341–65. [http://dx.doi.org/10.1002/oca.843.](http://dx.doi.org/10.1002/oca.843)