



A transmission dynamics model of COVID-19: Case of Cameroon



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ABSTRACT

In this work, we propose and investigate an ordinary differential equations model describing the spread of COVID-19 in Cameroon. The model takes into account the asymptomatic, unreported symptomatic, quarantine, hospitalized individuals and the amount of virus in the environment, for evaluating their impact on the transmission of the disease. After establishing the basic properties of the model, we compute the control reproduction number \mathcal{R}_c and show that the disease dies out whenever $\mathcal{R}_c \leq 1$ and is endemic whenever $\mathcal{R}_c > 1$. Furthermore, an optimal control problem is derived and investigated theoretically by mainly relying on Pontryagin's maximum principle. We illustrate the theoretical analysis by presenting some graphical results.

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1. Introduction

Many countries around the world are facing a new pandemic disease that destroys their populations daily. This is Coronavirus Disease 2019 (COVID-19) caused by Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2). Once the virus is in contact with a healthy person, the infection is contracted and the virus, once within the host, moves to the surface of the lungs, creating an inflammation of the lungs called pneumonia. This causes the blockade of the respiratory system and alters the immune system. This situation can degenerate and lead to the death of the patient. This phenomenon occurs over a small period of time estimated approximately to seven days (Gandhi et al., 2020). The COVID-19 symptoms are highly variable and are associated with severe illnesses such as fever, severe cold, shortness of breath or dyspnoea, chills, cough, lymphopenia, expectoration, fatigue, headache, acute pneumonia, sputum production, diarrhoea, hemoptysis most often followed by renal failure (Carlos et al., 2020; CDC 2020; Huang et al., 2020; Ren et al., 2020; WHO, 2020a, 2020b). The virus spreads mainly through the environment whenever people are close to each other, or through contaminated surfaces. This occurs when an

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infected person in this environment breathes, coughs, sneezes or speaks and then virus-containing particles exhaled comes into contact with another person either through the mouth, nose or eyes (CDC 2020). The longer the people interact, the more likely they are to transmit COVID-19. This disease has resulted in prolonged population containment, paralyzing economies in several countries. The total number of deaths worldwide due to this pandemic has exceeded 1.65 million and the cumulative number of confirmed cases topped 74.7 million (Wikipedia, 2020a, 2020b). In Africa for example, the number of confirmed cases amounted to 2,404,414 representing approximately 3.3% global infection and the overall deaths attributable to COVID-19 was around 56.74 thousand (Galal, 2020a, 2020b). Note that as of May 13, 2020, every country in Africa has recorded a COVID-19 case. South Africa was the most drastically affected country, with more than 866.1 thousand confirmed COVID-19 cases and 23,451 deaths (Galal, 2020a, 2020b). COVID-19 was confirmed to have reached Cameroon on 6 March 2020, through an infected person from France; this French citizen has been quarantined in the Yaounde Central Hospital (Kouagheu, 2020). The Cameroonian Government has implemented a nationwide series of measures in order to curtail the spread of COVID-19. The steps and dates of deaths and confirmed COVID-19 cases as well as recovered cases can be seen in (Wikipedia, 2020a, 2020b). Especially, a national state of disaster was declared on April 17, 2020. Schools, training institutions and many other activities were closed on the same date. As of December 2020, more than 441 deaths were reported and the number of COVID-19 confirmed cases was approximately 24,560. It is worth noting that many infected people due to COVID-19 in Cameroon were unreported, since the scheduled door-to-door screening campaign was not totally operated. These statistics make Cameroon the epicenter of COVID-19 in Central Africa. The causes of the rapid spread of COVID-19 in Cameroon are given in (Ojong, 2020). This mainly includes: negligence of quarantine, refusal of isolation and lack of financial means to hospitalize all symptomatic people.

The COVID-19 pandemic that continues to be a threat, resulting in increasing suffering of population, deserves a rigorous study to eradicate it within the community. Several mathematical models have been proposed and studied in order to understand the transmission dynamics of this pandemic. In (Ivorra et al., 2020), the authors developed a θ -SEIHRD mathematical model which takes into account the known special characteristics of COVID-19 pandemic such as the existence of unreported symptomatic infectious individuals and the different sanitary and infectiousness conditions of hospitalized individuals. This θ -SEIHRD model was also used to estimate a significant number of beds needed in hospitals. Mohsin and co-workers (Mohsin et al., 2020) formulated a mathematical model that included asymptomatic, quarantine and isolation compartments, and showed that the high levels of quarantine and isolation need to be maintained for controlling the disease. They also proposed an optimal control problem applied to the dynamics described by the obtained model. Based on reported data from December 31, 2019 to January 28, 2020, Wu and co-authors (Wu et al., 2020) used a SEIR model to predict the national and global spread of COVID-19 in China. In (Yang et al., 2020), the authors proposed a modified SEIR model that investigated the epidemic development of COVID-19 in China; the authors foretold the timing and magnitude of the epidemic peak as well as the ultimate epidemic size. This model has been recommended as a practical example of mathematical modeling techniques to investigate the spread of the pandemic (Krishna, 2020). In (Tang et al., 2020), the mathematical model developed includes individual epidemiological status, intervention measures and clinical progression of COVID-19. The authors found that mediation strategies such as intensive contact tracing followed by quarantine and isolation can effectively curtail the transmission risk and the control reproduction number. In (Zhang et al., 2020), Zhang and his collaborators thought that the increase in new cases of COVID-19 is due to crowding factor. They developed a mathematical model by using a nonlinear incidence rate and taking into account the aforementioned factors. They applied a nonstandard finite difference (NSFD) scheme and the fourth order Runge-Kutta (RK4) scheme to obtain the graphical results.

The main purpose of the present work is to propose and investigate an ordinary differential equations (ODE) model describing the spread of COVID-19 in Cameroon, and use it to evaluate the impact of control measures, such as quarantine and hospitalization strategies, on the spread of the pandemic in this country which occupies a strategic position in Central Africa.

This paper is organized as follows. We formulate the ODE model in Section 2. Section 3 is devoted to the mathematical analysis of the proposed model. Specifically, we prove the existence, uniqueness, positivity and boundedness of the solution. In Section 4, we compute the control reproduction number and study the existence and stability properties of equilibria. Moreover, we analyze the control reproduction number around the quarantine of exposed individuals and the isolation of hospitalized individuals. In Section 5, we propose and investigate an optimal control problem associated to the model studied in Section 3. In section 6, we provide numerical simulations to illustrate the theoretical results obtained. We conclude the work in section 7.

2. Model formulation

The model considered in this study consists of the total number of individuals in a human population at time t , denoted by $N(t)$, and sub-divided into eight distinct epidemiological subclasses of individuals, namely susceptible $S(t)$, exposed $E(t)$, asymptomatic infectious $A(t)$, symptomatic infectious $I(t)$, unreported symptomatic infectious $U(t)$, quarantined $Q(t)$, hospitalized $H(t)$, recovered $R(t)$, and the concentration of virus in the environment at time t , denoted by $V(t)$.

The dynamics description of each compartment is as follows.

Susceptible individuals, S , are recruited at a rate s , and decreased by natural death at a rate μ . Furthermore, as in (Safi and Gumel, 2010), we assume that the exposed class, E , and quarantined class, Q , do not transmit infection (i.e., exposed and quarantined individuals have a negligible number of contacts with members of the overall population; they play no role in the transmission process). So, it is assumed that only infected people presenting clinical symptoms can transmit the disease to

others. Thus, the susceptible population S may acquire infection, following effective contact with infectious individuals in the I , A , U , H or V classes at a rate λ_s , where

$$\lambda_s = \beta(I + A + U + \eta_1 H) + a_0 V. \quad (2.1)$$

In equation (2.1), the parameter β is the average number of effective contacts between susceptible and infected individuals (symptomatic, asymptomatic, unreported symptomatic and hospitalized individuals), while $0 = \eta_1 < 1$ is the modification parameter which accounts for the assumed reduction in disease transmission by hospitalized individuals in comparison to non-hospitalized infectious individuals in the I , A and U classes. η_1 measures the effectiveness of hospitalization; more precisely hospitalization is excellent if $\eta_1 = 0$, leaky if $0 < \eta_1 < 1$ and completely ineffective if $\eta_1 = 1$. Furthermore, we assume that the rate of transmissibility of the virus to the susceptible individuals is proportional to the free virus particles in the environment, and choose the force of infection as $a_0 V$. Thus, the rate of change of the susceptible population is expressed by the following equation:

$$\frac{dS}{dt} = s - \beta S(I + A + U + \eta_1 H) - a_0 SV - \mu S.$$

The population of exposed individuals, E , is generated by the infection of susceptible individuals at the rate λ_s . This class is decreased due to reported clinical symptoms at the rate η , unreported clinical symptoms at the rate b , asymptomatic infectious at a rate k , quarantine at the rate ϵ and natural death at the rate μ , so that

$$\frac{dE}{dt} = \beta S(I + A + U + \eta_1 H) + a_0 SV - (\mu + k + \epsilon + b + \eta)E.$$

The population of asymptomatic infectious individuals, A , is generated at the rate k . It is decreased due to natural recovery at the rate γ , unreported clinical symptoms at the rate θ , natural death at the rate μ and disease-induced death at the rate δ_4 . This gives

$$\frac{dA}{dt} = kE - (\mu + \delta_4 + \gamma + \theta)A.$$

The population of symptomatic infectious individuals, I , is generated at the rate η . This population is decreased due to natural recovery at the rate ρ , hospitalization at the rate d_0 , natural death at the rate μ and disease-induced death at the rate δ_1 . This is expressed as

$$\frac{dI}{dt} = \eta E - (\mu + \delta_1 + \rho + d_0)I.$$

The population of unreported symptomatic infectious individuals, U , is generated by the exposed individuals at the rate b and the asymptomatic infectious individuals at the rate θ . This class is decreased due to natural recovery at the rate ν , natural death at the rate μ and disease-induced death at the rate δ_3 . So we have

$$\frac{dU}{dt} = bE + \theta A - (\mu + \delta_3 + \nu)U.$$

Exposed individuals are quarantined at the rate ϵ . The population of quarantined individuals is decreased due to natural recovery at the rate α , hospitalization at the rate d_1 and natural death at the rate μ . Thus, one has

$$\frac{dQ}{dt} = \epsilon E - (\mu + \alpha + d_1)Q.$$

The population of hospitalized individuals, H , is generated by the hospitalization of quarantined individuals at the rate d_1 , and symptomatic infectious individuals at the rate d_0 . This population is decreased due to recovery at the rate r , natural death at the rate μ and disease-induced death at the rate δ_2 . We can assume that $\delta_2 < \delta_1$, $\delta_2 < \delta_3$ and $\delta_2 < \delta_4$. This means that hospitalized individuals have reduced disease-induced mortality rate in comparison to non-hospitalized infectious individuals because of care given in hospitals. Thus, the rate of change of the population of hospitalized individuals is expressed by the following equation:

$$\frac{dH}{dt} = d_0 I + d_1 Q - (\mu + \delta_2 + r)H.$$

The population of recovered individuals is generated by the recovery of asymptomatic infectious individuals at the rate γ , symptomatic infectious individuals at the rate ρ , unreported symptomatic infectious individuals at the rate ν , hospitalized infectious individuals at the rate r and quarantined individuals at the rate α . This population is decreased due to natural death at the rate μ . Therefore, we have the following equation:

$$\frac{dR}{dt} = \rho I + \nu U + \gamma A + \alpha Q + rH - \mu R.$$

Finally, the concentration of virus in the environment, V , is generated by the asymptomatic infectious individuals at the rate ω_1 , symptomatic infectious individuals at the rate σ , unreported symptomatic infectious individuals at the rate ω_0 and hospitalized infectious individuals at the rates a_1 . It is decreased by inactivation at the rate δ_5 . Thus,

$$\frac{dV}{dt} = \sigma I + a_1 H + \omega_1 A + \omega_0 U - \delta_5 V.$$

The flow diagram of the transmission dynamics of the COVID-19 is given in Fig. 1 below.

From the flow diagram in Fig. 1, we derive and propose the following nonlinear ODE system to describe the transmission dynamics of COVID-19 in Cameroon:

$$\left\{ \begin{array}{l} \frac{dS}{dt} = s - \beta S(I + A + U + \eta_1 H) - a_0 S V - \mu S, \\ \frac{dE}{dt} = \beta S(I + A + U + \eta_1 H) + a_0 S V - (\mu + k + \epsilon + b + \eta) E, \\ \frac{dA}{dt} = k E - (\mu + \delta_4 + \gamma + \theta) A, \\ \frac{dI}{dt} = \eta E - (\mu + \delta_1 + \rho + d_0) I, \\ \frac{dU}{dt} = b E + \theta A - (\mu + \delta_3 + \nu) U, \\ \frac{dQ}{dt} = \epsilon E - (\mu + \alpha + d_1) Q, \\ \frac{dH}{dt} = d_0 I + d_1 Q - (\mu + \delta_2 + r) H, \\ \frac{dV}{dt} = \sigma I + a_1 H + \omega_1 A + \omega_0 U - \delta_5 V, \\ \frac{dR}{dt} = \rho I + \nu U + \gamma A + \alpha Q + r H - \mu R, \end{array} \right. \quad (2.2)$$

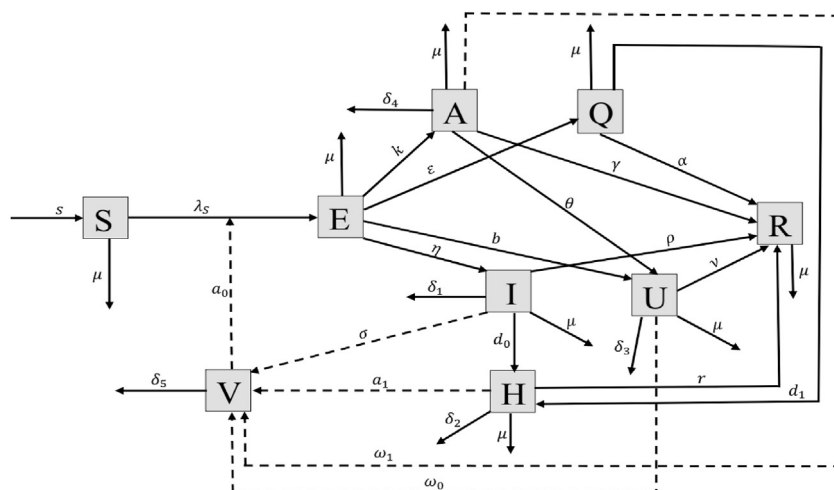


Fig. 1. Flow diagram of the COVID-19 transmission model in Cameroon.

with initial conditions:

$$S(0) > 0, E(0) > 0, A(0) > 0, I(0) > 0, U(0) > 0, Q(0) > 0, H(0) > 0, V(0) > 0, R(0) > 0. \tag{2.3}$$

The biological description of the parameters as well as their values and units are summed up in Table 1 below.

3. Basic properties of the full model

In this section, we explore the basic dynamical features of system (2.2). Since the COVID-19 model (2.2) monitors human populations, it will be epidemiologically meaningful if all its state variables are positive.

Theorem 3.1. *The solution $(S(t), E(t), A(t), I(t), U(t), Q(t), H(t), V(t), R(t))$ of system (2.2) starting from positive initial conditions (2.3) exists for all $t > 0$ and is unique. Furthermore,*

- a) $S(t) > 0, E(t) > 0, A(t) > 0, I(t) > 0, U(t) > 0, Q(t) > 0, H(t) > 0, V(t) > 0,$ and $R(t) > 0,$ for all time $t > 0.$
- b) *The biologically-feasible region $\Omega,$ defined by*

$$\Omega = \left\{ (S(t), E(t), A(t), I(t), U(t), Q(t), H(t), V(t), R(t)) \in \mathbb{R}_+^9 : \right. \\ \left. 0 < N(t) \leq \frac{s}{\mu}, \quad 0 \leq V(t) \leq \frac{s(\sigma + a_1 + \omega_1 + \omega_0)}{\mu\delta_5} \right\}, \tag{3.1}$$

is positively invariant for model (2.2).

proof. *The proof uses classical arguments from the theory of ODEs (Hale and Verduyn Lunel, 1993; Nkwayep et al., 2020). □* From Theorem 3.1, it follows that in Ω the system (2.2) is well-posed mathematically and epidemiologically. Accordingly, it is sufficient to study the dynamics of the flow generated by system (2.2) in $\Omega.$

Table 1
Biological description, values and units of the parameters of model (2.2).

Parameter	Biological description of the parameters of model (2.2)	Value/range	Reference
s	Constant recruitment rate into the community	3539 individual.day ⁻¹	(Population Data)
β	Effective contact rate between susceptible and infected individuals	$[3.62 \times 10^{-7}, 2 \times 10^{-6}]$ day ⁻¹	Estimated
η_1	Modification parameter for reduction of infectiousness for hospitalized individuals	$(0, 1]$ day ⁻¹	variable
η	Progression rate from exposed to symptomatic infectious class	0.12405 day ⁻¹	Tang et al. (2020)
ϵ	Quarantine rate of exposed individuals	0.1 day ⁻¹	Assumed
d_1	Hospitalization rate of quarantined individuals	0.04227 day ⁻¹	Assumed
d_0	Hospitalization rate of symptomatic infectious individuals	0.20619 day ⁻¹	Assumed
ρ	Recovery rate of symptomatic infectious individuals	0.33029 day ⁻¹	Tang et al. (2020)
r	Recovery rate of hospitalized individuals	0.11624 day ⁻¹	Tang et al. (2020)
δ_1	Disease-induced death rate of symptomatic infectious individuals	0.04227 day ⁻¹	Assumed
δ_2	Disease-induced death rate of hospitalized individuals	0.027855 day ⁻¹	Assumed
δ_3	Disease-induced death rate of unreported symptomatic infectious individuals	0.027855 day ⁻¹	Assumed
δ_4	Disease-induced death rate of asymptomatic infectious individuals	51×10^{-4} day ⁻¹	Estimated
δ_5	Decay rate of the virus	1/7 day ⁻¹	Estimated
μ	Natural death rate	$[1/59, 1/57]$ day ⁻¹	WHO (2020a, 2020b)
a_0	Transmission rate of the free virus	$[10^{-12}, 10^{-7}]$ (day.individual) ⁻¹	Estimated
b	Progression rate from exposed to unreported symptomatic infectious class	$(1-0.3)/7$ day ⁻¹	Estimated
ν	Recovery rate of unreported symptomatic infectious individuals	1/7 day ⁻¹	Liu et al. (2020)
α	Recovery rate of quarantined individuals	0.25 day ⁻¹	Estimated
k	Progression rate from exposed to asymptomatic infectious class	$(1-1.8887 \times 10^{-7})/7$ day ⁻¹	Tang et al. (2020)
ω_0	Shedding rate of the virus in the environment from unreported symptomatic infectious individuals	4.65×10^{-3} virus.(day.individual) ⁻¹	Estimated
ω_1	Shedding rate of the virus in the environment from asymptomatic infectious individuals	10^{-6} virus.(day.individual) ⁻¹	Estimated
θ	Progression rate from asymptomatic infectious to unreported symptomatic infectious class	$(1-0.7)/7$ day ⁻¹	Estimated
γ	Recovery rate of asymptomatic infectious individuals	0.13978 day ⁻¹	Tang et al. (2020)
σ	Shedding rate of the virus in the environment from symptomatic infectious individuals	6.39×10^{-3} virus.(day.individual) ⁻¹	Estimated

3.1. Existence and stability of equilibria

In this section, system (2.2) is analyzed to gain insight into its dynamical features.

3.2. Basic reproduction number and stability of the disease-free equilibrium (DFE)

The DFE of model (2.2) is obtained by setting the right hand sides of the equations to zero; it is given by:

$$\mathcal{E}_0 = (S^*, E^*, A^*, I^*, U^*, Q^*, H^*, V^*, R^*) = \left(\frac{S}{\mu}, 0, 0, 0, 0, 0, 0 \right). \tag{4.1}$$

Now, to explore the local stability of \mathcal{E}_0 , we will use the next generation operator method developed in (Diekmann et al., 1990; van den Driessche and Watmough, 2002). More precisely, by using the matrix notation of Lemma 1 in (van den Driessche and Watmough, 2002), it follows that the matrix, F , of the new infection terms, and the non-singular M -matrix, V_1 , of the remaining transfer terms associated with model (2.2), are given, respectively, by

$$F = \begin{pmatrix} 0 & \frac{\beta s}{\mu} & \frac{\beta s}{\mu} & \frac{\beta s}{\mu} & 0 & \frac{\beta \eta_1 s}{\mu} & \frac{s a_0}{\mu} \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 & 0 \end{pmatrix} \quad \text{and} \quad V_1 = \begin{pmatrix} k_1 & 0 & 0 & 0 & 0 & 0 & 0 \\ -k & k_2 & 0 & 0 & 0 & 0 & 0 \\ -\eta & 0 & k_3 & 0 & 0 & 0 & 0 \\ -b & -\theta & 0 & k_4 & 0 & 0 & 0 \\ -\epsilon & 0 & 0 & 0 & k_5 & 0 & 0 \\ 0 & 0 & -d_0 & 0 & -d_1 & k_6 & 0 \\ 0 & -\omega_1 & -\sigma & -\omega_0 & 0 & -a_1 & \delta_5 \end{pmatrix}.$$

It follows that the control reproduction number (Anderson & May 1982; Hethcote, 2000), denoted by $\mathcal{R}_c = \rho(FV_1^{-1})$, where $\rho(FV_1^{-1})$ is the spectral radius of the next generation matrix FV_1^{-1} , is given by

$$\mathcal{R}_c = \frac{\beta ks}{\mu k_1 k_2} + \frac{\beta \eta s}{\mu k_1 k_3} + \frac{\beta bs}{\mu k_1 k_4} + \frac{\beta \theta ks}{\mu k_1 k_2 k_4} + \frac{\beta \eta_1 \eta d_0 s}{\mu k_1 k_3 k_6} + \frac{\beta \eta_1 \epsilon d_1 s}{\mu k_1 k_5 k_6} + \frac{k \theta a_0 \omega_0 s}{\mu k_1 k_2 k_4 \delta_5} + \frac{k a_0 \omega_1 s}{\mu k_1 k_2 \delta_5} + \frac{\eta d_0 a_0 a_1 s}{\mu k_1 k_3 k_6 \delta_5} + \frac{\eta \sigma a_0 s}{\mu k_1 k_3 \delta_5} + \frac{b a_0 \omega_0 s}{\mu k_1 k_4 \delta_5} + \frac{\epsilon d_1 a_0 a_1 s}{\mu k_1 k_5 k_6 \delta_5},$$

where

$$\begin{aligned} k_1 &= \mu + k + \epsilon + b + \eta, & k_2 &= \mu + \delta_4 + \gamma + \theta, & k_3 &= \mu + \delta_1 + \rho + d_0, \\ k_4 &= \mu + \delta_3 + \nu, & k_5 &= \mu + \alpha + d_1 & \text{and} & k_6 &= \mu + \delta_2 + r. \end{aligned} \tag{4.2}$$

The epidemiological meaning of the quantity \mathcal{R}_c (reproduction number of the full model with control measures) is that, it measures the average number of new COVID-19 positive cases generated by a single typical COVID-19-infected individual (living or dead) introduced into a completely-susceptible human population. This infers that, COVID-19 can be effectively controlled in the community if the threshold quantity \mathcal{R}_c is less than unity (i.e. $\mathcal{R}_c < 1$). Thus, COVID-19 cannot develop from a small influx of infected individuals if $\mathcal{R}_c < 1$, but COVID-19 will develop if $\mathcal{R}_c > 1$. Now, the epidemiological interpretation of each term of \mathcal{R}_c is as follows. First, the mean duration of an infective individual in class E is $1/k_1$. A fraction k/k_1 of infective individuals moves from class E into class A with effective contact rate β and mean duration $1/k_2$, offering a contribution of $\beta ks/\mu k_1 k_2$ to \mathcal{R}_c . Next, a fraction η/k_1 of infective individuals moves from class E into class I , with effective contact rate β and mean duration $1/k_3$, offering a contribution of $\beta \eta s/\mu k_1 k_3$ to \mathcal{R}_c . A fraction b/k_1 moves from class E into class U with effective contact rate β and mean duration $1/k_4$, giving a contribution of $\beta bs/\mu k_1 k_4$, and after a severity of infection, a fraction $\theta k/k_1 k_2$ moves from class A into class U , giving a contribution of $\beta \theta ks/\mu k_1 k_2 k_4$ to \mathcal{R}_c . A fraction ϵ/k_1 moves from E to Q and the mean duration of Q is $1/k_5$. A fraction $\eta d_0/k_1 k_3$ moves from E to I then to H with effective contact rate $\beta \eta_1$ and mean duration $1/k_6$, offering a contribution of $\beta \eta_1 \eta d_0 s/\mu k_1 k_3 k_6$ to \mathcal{R}_c . A fraction $\epsilon d_1/k_1 k_5$ moves from E to Q then to H with effective contact rate $\beta \eta_1$ and mean duration $1/k_6$, offering a contribution of $\beta \eta_1 \epsilon d_1 s/\mu k_1 k_5 k_6$ to \mathcal{R}_c . A fraction $k \theta a_0 \omega_0 s/\mu k_1 k_2 k_4$ moves from E to A then to U and to V with effective contact rate a_0 and mean duration $1/\delta_5$, giving a contribution of $k \theta a_0 \omega_0 s/\mu k_1 k_2 k_4 \delta_5$ to \mathcal{R}_c . A fraction $k a_0 \omega_1 s/\mu k_1 k_2$ moves from E to A then to V with effective contact rate a_0 and mean duration $1/\delta_5$, giving a contribution of $k a_0 \omega_1 s/\mu k_1 k_2 \delta_5$ to \mathcal{R}_c . A fraction $\eta d_0 a_0 a_1 s/\mu k_1 k_3 k_6$ moves from E to I then to H and to V with effective contact rate a_0 and mean duration $1/\delta_5$, giving a contribution of $\eta d_0 a_0 a_1 s/\mu k_1 k_3 k_6 \delta_5$ to \mathcal{R}_c . A fraction $\eta \sigma a_0 s/\mu k_1 k_3$ moves from E to I then to V with effective contact rate a_0 and mean duration $1/\delta_5$, giving a contribution of $\eta \sigma a_0 s/\mu k_1 k_3 \delta_5$ to \mathcal{R}_c . A fraction $b a_0 \omega_0 s/\mu k_1 k_4$ moves from E to U then to V with effective contact rate a_0 and mean duration $1/\delta_5$, giving a contribution of $b a_0 \omega_0 s/\mu k_1 k_4 \delta_5$ to \mathcal{R}_c . Finally, a fraction $\epsilon d_1 a_1 s/\mu k_1 k_5 k_6$ moves

from E to Q then to H and to V with effective contact rate a_0 and mean duration $1/\delta_5$, giving a contribution of $\epsilon d_1 a_0 a_1 s / \mu k_1 k_5 k_6 \delta_5$ to \mathcal{R}_c .

Note that the basic reproduction number \mathcal{R}_0 is defined in the absence of control measures such as quarantine, isolation and environmental spraying techniques to disinfect exposed surfaces. Thus \mathcal{R}_0 is \mathcal{R}_c with $\epsilon = d_0 = a_1 = \sigma = \omega_0 = \omega_1 = 0$. It then follows that

$$\mathcal{R}_0 = \frac{\beta ks}{\mu k_{01} k_{02}} + \frac{\beta \eta s}{\mu k_{01} k_{03}} + \frac{\beta bs}{\mu k_{01} k_{04}} + \frac{\beta \theta ks}{\mu k_{01} k_{02} k_{04}},$$

where

$$k_{01} = \mu + k + b + \eta, \quad k_{02} = \mu + \delta_4 + \gamma + \theta, \quad k_{03} = \mu + \delta_1 + \rho, \quad \text{and} \quad k_{04} = \mu + \delta_3 + \nu.$$

The following result is obtained by using similar arguments as in the proof of Theorem 2 in (van den Driessche and Watmough, 2002).

Lemma 4.1. *The DFE, \mathcal{E}_0 , of system (2.2), given by (4.1), is locally asymptotically stable in Ω whenever $\mathcal{R}_c < 1$, and unstable if $\mathcal{R}_c > 1$.*

Proof. Linearizing (2.2) at the DFE \mathcal{E}_0 , we obtain the linearized system

$$\frac{dW}{dt} = J(\mathcal{E}_0)W,$$

where $W = (S, E, A, I, U, Q, H, V, R)$ and

$$J(\mathcal{E}_0) = \begin{pmatrix} -\mu & 0 & \frac{\beta s}{\mu} & \frac{\beta s}{\mu} & \frac{\beta s}{\mu} & 0 & \frac{\beta \eta_1 s}{\mu} & \frac{sa_0}{\mu} & 0 \\ 0 & -k_1 & \frac{\beta s}{\mu} & \frac{\beta s}{\mu} & \frac{\beta s}{\mu} & 0 & \frac{\beta \eta_1 s}{\mu} & \frac{sa_0}{\mu} & 0 \\ 0 & k & -k_2 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & \eta & 0 & -k_3 & 0 & 0 & 0 & 0 & 0 \\ 0 & b & \theta & 0 & -k_4 & 0 & 0 & 0 & 0 \\ 0 & \epsilon & 0 & 0 & 0 & -k_5 & 0 & 0 & 0 \\ 0 & 0 & 0 & d_0 & 0 & d_1 & -k_6 & 0 & 0 \\ 0 & 0 & \omega_1 & \sigma & \omega_0 & 0 & a_1 & -\delta_5 & 0 \\ 0 & 0 & \gamma & \rho & \nu & \alpha & r & 0 & -\mu \end{pmatrix}.$$

Now, to end the proof, it is necessary to prove that all eigenvalues of the Jacobian matrix, $J(\mathcal{E}_0)$, have negative real parts. So, writing the Jacobian matrix, $J(\mathcal{E}_0)$, under the distributed matrix form, we obtain

$$\begin{pmatrix} -\mu & c_1 & c_2 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & c_3 & c_4 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & k & -k_2 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & \eta & 0 & -k_3 & 0 & 0 & 0 & 0 & 0 \\ 0 & b & \theta & 0 & -k_4 & 0 & 0 & 0 & 0 \\ 0 & \epsilon & 0 & 0 & 0 & -k_5 & 0 & 0 & 0 \\ 0 & 0 & 0 & d_0 & 0 & d_1 & -k_6 & 0 & 0 \\ 0 & 0 & \omega_1 & \sigma & \omega_0 & 0 & a_1 & -\delta_5 & 0 \\ 0 & 0 & \gamma & \rho & \nu & \alpha & r & 0 & -\mu \end{pmatrix},$$

where

$$c_1 = -\frac{sa_0 a_1 d_1 \epsilon}{\mu k_5 k_6 \delta_5} - \frac{\beta \eta_1 d_1 s \epsilon}{\mu k_5 k_6} - \frac{sa_0 \omega_0 b}{\mu k_4 \delta_5} - \frac{\beta bs}{\mu k_4} - \frac{sa_0 \sigma \eta}{\mu k_3 \delta_5} - \frac{\beta \eta s}{\mu k_3} - \frac{sa_0 a_1 d_0 \eta}{\mu k_3 k_6 \delta_5} - \frac{\beta \eta_1 d_0 s \eta}{\mu k_3 k_6}, \quad c_2 = -\frac{\beta s}{\mu} - \frac{sa_0 \omega_1}{\mu \delta_5} - \frac{sa_0 \omega_0 \theta}{\mu k_4 \delta_5} - \frac{\beta \theta s}{\mu k_4},$$

$$c_3 = -\frac{1}{k_2} + \frac{sa_0 a_1 d_1 \epsilon}{\mu k_1 k_2 k_3 k_5 k_6 \delta_5} + \frac{\beta \eta_1 d_1 s \epsilon}{\mu k_1 k_2 k_3 k_5 k_6} + \frac{sa_0 \omega_0 b}{\mu k_1 k_2 k_4 \delta_5} + \frac{\beta bs}{\mu k_1 k_2 k_4} + \frac{sa_0 \sigma \eta}{\mu k_1 k_2 k_3 \delta_5} + \frac{\beta \eta s}{\mu k_1 k_2 k_3} + \frac{sa_0 a_1 d_0 \eta}{\mu k_1 k_2 k_3 k_6 \delta_5} + \frac{\beta \eta_1 d_0 s \eta}{\mu k_1 k_2 k_3 k_6},$$

$$c_4 = \frac{\beta s}{\mu k_1 k_2} + \frac{sa_0 \omega_1}{\mu k_1 k_2 \delta_5} + \frac{sa_0 \omega_0 \theta}{\mu k_1 k_2 k_4 \delta_5} + \frac{\beta \theta s}{\mu k_1 k_2 k_4}.$$

Let M be the following three dimensional matrix defined by

$$M = \begin{pmatrix} -\mu & c_1 & c_2 \\ 0 & c_3 & c_4 \\ 0 & k & -k_2 \end{pmatrix}.$$

Note that, all eigenvalues of the Jacobian matrix, $J(\mathcal{E}_0)$, have negative real parts whenever $\det(M) < 0$. The computation of $\det(M)$, gives

$$\begin{aligned} \det(M) &= \mu(kc_4 + k_2c_3), \\ &= \mu \left(\frac{\beta ks}{\mu k_1 k_2} + \frac{\beta \eta s}{\mu k_1 k_3} + \frac{\beta bs}{\mu k_1 k_4} + \frac{\beta \theta ks}{\mu k_1 k_2 k_4} + \frac{\beta \eta_1 \eta d_0 s}{\mu k_1 k_3 k_6} + \frac{\beta \eta_1 \epsilon d_1 s}{\mu k_1 k_5 k_6} + \frac{k \theta a_0 \omega_0 s}{\mu k_1 k_2 k_4 \delta_5} \right. \\ &\quad \left. + \frac{k a_0 \omega_1 s}{\mu k_1 k_2 \delta_5} + \frac{\eta d_0 a_0 a_1 s}{\mu k_1 k_3 k_6 \delta_5} + \frac{\eta \sigma a_0 s}{\mu k_1 k_3 \delta_5} + \frac{b a_0 \omega_0 s}{\mu k_1 k_4 \delta_5} + \frac{\epsilon d_1 a_0 a_1 s}{\mu k_1 k_5 k_6 \delta_5} - 1 \right), \\ &= \mu(\mathcal{R}_c - 1). \end{aligned}$$

So if $\mathcal{R}_c < 1$, it follows that $\det(M) < 0$. In this case, all eigenvalues of the Jacobian matrix $J(\mathcal{E}_0)$ have negative real parts. Thus, if $\mathcal{R}_c < 1$, the DFE, \mathcal{E}_0 , of system (2.2), given by (4.1), is locally asymptotically stable. If $\mathcal{R}_c > 1$, then $\det(M) > 0$. This infers that, there exists an eigenvalue of the Jacobian matrix $J(\mathcal{E}_0)$ with positive real part. So, if $\mathcal{R}_c > 1$, then \mathcal{E}_0 is unstable. This completes the proof. \square

Remark 4.2. Lemma 4.1 communicates that COVID-19 is eliminated from the population (when $\mathcal{R}_c < 1$) if the initial sizes of the sub-populations of the obtained system are in the basin of attraction of the DFE \mathcal{E}_0 . In what follows, to ensure that COVID-19 is eliminated from the population regardless of the initial sizes of the sub-populations, we need to prove the global stability of \mathcal{E}_0 .

3.3. Global stability of the DFE \mathcal{E}_0

In this section, we investigate the global stability of the DFE, \mathcal{E}_0 , by constructing a suitable Lyapunov functional and using LaSalle's invariance principle. For this purpose, consider the following function defined for positive real numbers by

$$g(x) = x - 1 - \ln x. \tag{4.3}$$

It can be shown that $g(x) \geq 0$ for all $x > 0$, and that $\min_{x>0} g(x) = g(1) = 0$. We have the following result.

Theorem 4.3. The DFE, \mathcal{E}_0 , of system (2.2), given by (4.1), is globally asymptotically stable in Ω whenever $\mathcal{R}_c \leq 1$.

Proof. Let $(S(t), E(t), A(t), I(t), U(t), Q(t), H(t), V(t), R(t))$ be any positive solution of system (2.2) in Ω . Recall that $S^* = s/\mu$. Define the following Lyapunov function

$$\begin{aligned} L(t) &= \frac{\delta_5 S^* \mathcal{R}_c}{\beta \eta_1} \left(\frac{S}{S^*} - 1 - \ln \frac{S}{S^*} \right) + \frac{\delta_5 \mathcal{R}_c}{\beta \eta_1} E + \frac{k_4 \beta \delta_5 S^* + k_4 a_0 \omega_1 S^* + a_0 \theta \omega_0 S^* + \beta \theta \delta_5 S^*}{\beta \eta_1 k_2 k_4} A \\ &\quad + \frac{k_6 \beta \delta_5 S^* + a_0 a_1 d_0 S^* + a_0 \sigma k_6 S^* + \beta \eta_1 d_0 \delta_5 S^*}{\beta \eta_1 k_3 k_6} I + \frac{\beta \delta_5 S^* + a_0 \omega_0 S^*}{\beta \eta_1 k_4} U \\ &\quad + \frac{\beta \eta_1 d_1 \delta_5 S^* + a_0 a_1 d_1 S^*}{\beta \eta_1 k_5 k_6} Q + \frac{\beta \eta_1 \delta_5 S^* + a_0 a_1 S^*}{\beta \eta_1 k_6} H + \frac{a_0 S^*}{\beta \eta_1} V. \end{aligned} \tag{4.4}$$

Then, it is clear that, the function L is nonnegative definite in Ω with respect to \mathcal{E}_0 . Calculating the time derivative of the function L along the solution of system (2.2), after lengthy computations, we get

$$\frac{dL}{dt} = \frac{s \delta_5 \mathcal{R}_c}{\beta \eta_1} \left(2 - \frac{S}{S^*} - \frac{S^*}{S} \right) + \frac{S^* \delta_5}{\eta_1} \left(I + A + U + \eta_1 H + \frac{a_0}{\beta} V \right) (\mathcal{R}_c - 1).$$

Thus, it follows that condition $\mathcal{R}_c \leq 1$ ensures $\frac{dL(t)}{dt} \leq 0$, for all $S, E, I, A, U, H, V \geq 0$, with equality $\left(\frac{dL(t)}{dt} = 0 \right)$ if and only if $S = S^*$, $E = 0, I = 0, A = 0, U = 0, H = 0, V = 0$. Thus, L is a Lyapunov function on Ω . So, by LaSalle's invariance Principle [12, Theorem 5.3.1], it follows that

$$\lim_{t \rightarrow \infty} (S, E, A, I, U, Q, H, V) = \left(\frac{S}{\mu}, 0, 0, 0, 0, 0, 0, 0 \right). \tag{4.5}$$

Let $\mathbb{K} = (A, I, U, Q, H)$. Then from (4.5), one has $\limsup_{t \rightarrow \infty} \mathbb{K} = 0$. This implies that for a sufficiently small $\varepsilon > 0$ there exist constants $M_i > 0, i = 1, \dots, 5$ such that $\limsup_{t \rightarrow \infty} \mathbb{K} \leq \varepsilon$, for all $t > M_i, i = 1, \dots, 5$. Thus, from the eighth equation of system (2.2), it follows that, for $t > \max_{i \in \{1, \dots, 5\}} M_i$,

$$R^\infty = \limsup_{t \rightarrow \infty} R \leq \frac{\rho\varepsilon + \nu\varepsilon + \gamma\varepsilon + \alpha\varepsilon + r\varepsilon}{\mu}, \tag{4.6}$$

so that, by letting $\varepsilon \rightarrow 0$ in (4.6), we get

$$R^\infty = \limsup_{t \rightarrow \infty} R \leq 0. \tag{4.7}$$

Also from (4.5), one has $\liminf_{t \rightarrow \infty} \mathbb{K} = 0$. Thus, by using a similar argument as above, it can be shown that

$$R_\infty = \liminf_{t \rightarrow \infty} R \geq 0. \tag{4.8}$$

It then follows from (4.7) and (4.8) that

$$R_\infty \geq 0 \geq R^\infty.$$

This infers that

$$\lim_{t \rightarrow \infty} R(t) = 0. \tag{4.9}$$

Thus we have from (4.5) and (4.9) that,

$$\lim_{t \rightarrow \infty} (S, E, A, I, U, Q, H, V, R) = \left(\frac{S}{\mu}, 0, 0, 0, 0, 0, 0, 0, 0 \right)$$

Moreover, Ω is an invariant and attracting set of \mathbb{R}_+^9 . It follows that the largest compact invariant subset in $\{(S, E, A, I, U, Q, H, V, R) \in \Omega : \frac{dL}{dt} = 0\}$ is the singleton $\{\mathcal{E}_0\}$. So, by LaSalle’s invariance Principle [12, Theorem 5.3.1], it follows that every solution of system (2.2), with initial conditions in \mathbb{R}_+^9 , approaches the DFE, \mathcal{E}_0 , as $t \rightarrow \infty$ whenever $\mathcal{R}_c \leq 1$. This completes the proof. \square

Remark 4.4. We note that the following Lyapunov function could also be used to prove Theorem 4.3

$$\begin{aligned} \tilde{\mathcal{M}}(t) = & \frac{\delta_5 S^*}{\beta\eta_1} \left(\frac{S}{S^*} - 1 - \ln \frac{S}{S^*} \right) + \frac{\delta_5}{\beta\eta_1} E + \frac{k_4\beta\delta_5 S^* + k_4 a_0 \omega_1 S^* + a_0 \theta \omega_0 S^* + \beta\theta\delta_5 S^*}{\beta\eta_1 k_2 k_4} A \\ & + \frac{k_6\beta\delta_5 S^* + a_0 a_1 d_0 S^* + a_0 \sigma k_6 S^* + \beta\eta_1 d_0 \delta_5 S^*}{\beta\eta_1 k_3 k_6} I + \frac{\beta\delta_5 S^* + a_0 \omega_0 S^*}{\beta\eta_1 k_4} U \\ & + \frac{\beta\eta_1 d_1 \delta_5 S^* + a_0 a_1 d_1 S^*}{\beta\eta_1 k_5 k_6} Q + \frac{\beta\eta_1 \delta_5 S^* + a_0 a_1 S^*}{\beta\eta_1 k_6} H + \frac{a_0 S^*}{\beta\eta_1} V. \end{aligned} \tag{4.10}$$

In this case, its derivative gives

$$\frac{d\tilde{\mathcal{M}}(t)}{dt} = \frac{s\delta_5}{\beta\eta_1} \left(2 - \frac{S}{S^*} - \frac{S^*}{S} \right) + \frac{k_1 \delta_5}{\beta\eta_1} E (\mathcal{R}_c - 1). \tag{4.11}$$

Thus, combining (4.11) and (4.9) also leads to the global asymptotical stability of the DFE, \mathcal{E}_0 .

Theorem 4.3 implies that COVID-19 is eliminated from the population if the control reproduction number, \mathcal{R}_c , of the model (2.2) is less than or equal to one. Thus, **Theorem 4.3** means epidemiologically that the use of quarantine, hospitalization and the control of the amount of virus in the environment can lead to elimination of the COVID-19 if the mentioned controls can keep the threshold quantity, \mathcal{R}_c , to a value less than or equal to unity. This implies that the condition $\mathcal{R}_c \leq 1$ is necessary and sufficient for the elimination of COVID-19. Moreover, it follows from **Theorem 4.3** that the longer infected individuals abide in the exposed class, the higher the likelihood of COVID-19 eradication from the population.

4.3. Existence of the endemic equilibrium point (EEP)

Let $\mathcal{E}^* = (S^{**}, E^{**}, A^{**}, I^{**}, U^{**}, Q^{**}, H^{**}, V^{**}, R^{**})$ be any arbitrary equilibrium of system (2.2). In this section, we provide conditions for the existence of equilibria for which COVID-19 is endemic in the community, that is, at least one of the infected variables is non-zero. For this, consider the following associated force of infection for COVID-19 at endemic steady state

$$\lambda_s^{**} = \beta(I^{**} + A^{**} + U^{**} + \eta_1 H^{**}) + a_0 V^{**}. \tag{4.12}$$

The endemic equilibrium point (EEP) of system (2.2) is obtained by setting the right hand side of the equations to zero; it is given in terms of $\lambda_s^{**} S^{**}$ as follows:

$$\begin{aligned} E^{**} &= \frac{\lambda_s^{**} S^{**}}{k_1}, & A^{**} &= \frac{k \lambda_s^{**} S^{**}}{k_1 k_2}, & I^{**} &= \frac{\eta \lambda_s^{**} S^{**}}{k_1 k_3}, & U^{**} &= N_1 \lambda_s^{**} S^{**}, \\ Q^{**} &= \frac{\epsilon \lambda_s^{**} S^{**}}{k_1 k_5}, & H^{**} &= N_2 \lambda_s^{**} S^{**}, & V^{**} &= N_3 \lambda_s^{**} S^{**}, & R^{**} &= N_4 \lambda_s^{**} S^{**}, \end{aligned} \tag{4.13}$$

where,

$$\begin{aligned} S^{**} &= \frac{s}{\lambda_s^{**} + \mu}, & N_1 &= \frac{b}{k_1 k_4} + \frac{k\theta}{k_1 k_2 k_4}, & N_2 &= \frac{d_0 \eta}{k_1 k_3 k_6} + \frac{d_1 \epsilon}{k_1 k_5 k_6}, & N_3 &= \frac{\sigma \eta}{k_1 k_3 \delta_5} + \frac{a_1 d_0 \eta}{k_1 k_3 k_6 \delta_5} + \frac{a_1 d_1 \epsilon}{k_1 k_5 k_6 \delta_5} + \frac{k \omega_1}{k_1 k_2 \delta_5} + \frac{b \omega_0}{k_1 k_4 \delta_5} + \frac{k \theta \omega_0}{k_1 k_2 k_4 \delta_5}, \\ N_4 &= \frac{\rho \eta}{k_1 k_3 \mu} + \frac{b \nu}{k_1 k_4 \mu} + \frac{k \nu \theta}{k_2 k_2 k_4 \mu} + \frac{k \gamma}{k_1 k_2 \mu} + \frac{\epsilon \alpha}{k_1 k_5 \mu} + \frac{r d_0 \eta}{k_1 k_3 k_6 \mu} + \frac{r d_1 \epsilon}{k_1 k_5 k_6 \mu}. \end{aligned}$$

Inserting the expressions of (4.13), except R^{**} , into (4.12), gives

$$\lambda_s^{**} = \lambda_s^{**} S^{**} \left[\frac{\beta \eta}{k_1 k_3} + \frac{\beta k}{k_1 k_2} + \beta N_1 + \beta \eta_1 N_2 + a_0 N_3 \right]. \tag{4.14}$$

Using the expression of S^{**} , equation (4.14) becomes

$$\lambda_s^{**} \lambda_s^{**} + \mu \lambda_s^{**} = s \lambda_s^{**} \left[\frac{\beta \eta}{k_1 k_3} + \frac{\beta k}{k_1 k_2} + \beta N_1 + \beta \eta_1 N_2 + a_0 N_3 \right]. \tag{4.15}$$

As mentioned above, we have $\lambda_s^{**} \neq 0$. Dividing each term in (4.15) by λ_s^{**} , we obtain

$$1 + \frac{1}{\mu} \lambda_s^{**} = \frac{\beta \eta s}{\mu k_1 k_3} + \frac{\beta k s}{\mu k_1 k_2} + \frac{\beta s}{\mu} N_1 + \frac{\beta \eta_1 s}{\mu} N_2 + \frac{a_0 s}{\mu} N_3. \tag{4.16}$$

It is worth noting that

$$\begin{aligned} 1 + \frac{1}{\mu} \lambda_s^{**} &= \frac{\beta k s}{\mu k_1 k_2} + \frac{\beta \eta s}{\mu k_1 k_3} + \frac{\beta b s}{\mu k_1 k_4} + \frac{\beta \theta k s}{\mu k_1 k_2 k_4} + \frac{\beta \eta_1 \eta d_0 s}{\mu k_1 k_3 k_6} + \frac{\beta \eta_1 \epsilon d_1 s}{\mu k_1 k_5 k_6} + \frac{k \theta a_0 \omega_0 s}{\mu k_1 k_2 k_4 \delta_5} \\ &\quad + \frac{k a_0 \omega_1 s}{\mu k_1 k_2 \delta_5} + \frac{\eta d_0 a_0 a_1 s}{\mu k_1 k_3 k_6 \delta_5} + \frac{\eta \sigma a_0 s}{\mu k_1 k_3 \delta_5} + \frac{b a_0 \omega_0 s}{\mu k_1 k_4 \delta_5} + \frac{\epsilon d_1 a_0 a_1 s}{\mu k_1 k_5 k_6 \delta_5}, \\ &= \mathcal{R}_c. \end{aligned}$$

Thus,

$$\lambda_s^{**} = \mu(\mathcal{R}_c - 1) > 0, \quad \text{if } \mathcal{R}_c > 1. \tag{4.17}$$

Hence, each coordinate of the EEP \mathcal{E}^* is obtained by introducing the unique value of λ_s^{**} provided in (4.17) into the different expressions in (4.13). Summarizing the above discussion on the EEP \mathcal{E}^* , we obtain the following result.

Lemma 4.5. *If $\mathcal{R}_c > 1$, then system (2.2) admits in Ω a unique positive endemic equilibrium, \mathcal{E}^* .*

4.4. Local stability of the endemic equilibrium point

This section is devoted to the local stability of the unique endemic equilibrium point guaranteed by Lemma 4.2 whenever $\mathcal{R}_c > 1$. To do this, we follow the method developed in (Hethcote and Thieme, 1985) that takes its essence from the technique proposed by Krasnoselskii (Krasnoselskii, 1964).

We have the following result.

Theorem 4.6. *If $\mathcal{R}_c > 1$, then the unique endemic equilibrium point, \mathcal{E}^* , of system (2.2) is locally asymptotically stable.*

proof. First of all, note that the total population N is asymptotically constant, that is $N \rightarrow N^*$ as $t \rightarrow \infty$. Thus, the proof of Theorem 4.6 is established by using a reduced system of (2.2), which is obtained by considering only the components E, A, I, U, Q, H, V, R . Thus, we can set $N = N^*$, for large t , so that the unique endemic equilibrium point, \mathcal{E}^* , of the system (2.2) becomes $\mathcal{E}_1^* = \mathcal{E}^*|_{N=N^*}$. This eliminates the equation for S from this part of the analysis through the substitution $S = N^* - (E + A + I + U + Q + H + R)$, in which case, system (2.2) is reduced to

$$\left\{ \begin{aligned} \frac{dE}{dt} &= (\beta(I + A + U + \eta_1 H) + a_0 V)(N^* - E - A - I - U - Q - H - R) - (\mu + k + \epsilon + b + \eta)E, \\ \frac{dA}{dt} &= kE - (\mu + \delta_4 + \gamma + \theta)A, \\ \frac{dI}{dt} &= \eta E - (\mu + \delta_1 + \rho + d_0)I, \\ \frac{dU}{dt} &= bE + \theta A - (\mu + \delta_3 + \nu)U, \\ \frac{dQ}{dt} &= \epsilon E - (\mu + \alpha + d_1)Q, \\ \frac{dH}{dt} &= d_0 I + d_1 Q - (\mu + \delta_2 + r)H, \\ \frac{dV}{dt} &= \sigma I + a_1 H + \omega_1 A + \omega_0 U - \delta_5 V, \\ \frac{dR}{dt} &= \rho I + \nu U + \gamma A + \alpha Q + rH - \mu R. \end{aligned} \right. \tag{4.18}$$

Now, linearizing system (4.18) at the endemic equilibrium point, \mathcal{E}_1^* , yields

$$\left\{ \begin{aligned} \frac{dE}{dt} &= [-e_1 - (\mu + k + \epsilon + b + \eta)]E + (e_2 - e_1)A + (e_2 - e_1)I + (e_2 - e_1)U - e_1 Q + (\eta_1 e_2 - e_1)H + \frac{a_0 e_2}{\beta} V - e_1 R, \\ \frac{dA}{dt} &= kE - (\mu + \delta_4 + \gamma + \theta)A, \\ \frac{dI}{dt} &= \eta E - (\mu + \delta_1 + \rho + d_0)I, \\ \frac{dU}{dt} &= bE + \theta A - (\mu + \delta_3 + \nu)U, \\ \frac{dQ}{dt} &= \epsilon E - (\mu + \alpha + d_1)Q, \\ \frac{dH}{dt} &= d_0 I + d_1 Q - (\mu + \delta_2 + r)H, \\ \frac{dV}{dt} &= \sigma I + a_1 H + \omega_1 A + \omega_0 U - \delta_5 V, \\ \frac{dR}{dt} &= \rho I + \nu U + \gamma A + \alpha Q + rH - \mu R, \end{aligned} \right. \tag{4.19}$$

where

$$e_1 = \mu(\mathcal{R}_c - 1) \quad \text{and} \quad e_2 = \frac{\beta s}{\mu \mathcal{R}_c}.$$

Thus, the Jacobian matrix of this linearized system (4.19), evaluated at \mathcal{E}_1^* , is

$$J_1(\mathcal{E}_1^*) = \begin{pmatrix} -e_1 - k_1 & e_2 - e_1 & e_2 - e_1 & e_2 - e_1 & -e_1 & \eta_1 e_2 - e_1 & \frac{a_0 e_2}{\beta} & -e_1 \\ k & -k_2 & 0 & 0 & 0 & 0 & 0 & 0 \\ \eta & 0 & -k_3 & 0 & 0 & 0 & 0 & 0 \\ b & \theta & 0 & -k_4 & 0 & 0 & 0 & 0 \\ \epsilon & 0 & 0 & 0 & -k_5 & 0 & 0 & 0 \\ 0 & 0 & d_0 & 0 & d_1 & -k_6 & 0 & 0 \\ 0 & \omega_1 & \sigma & \omega_0 & 0 & a_1 & -\delta_5 & 0 \\ 0 & \gamma & \rho & \nu & \alpha & r & 0 & -\mu \end{pmatrix}.$$

Now, following the method developed in (Hethcote and Thieme, 1985), we assume that the linearized system (4.19) has solution of the form

$$\mathbf{Z}(t) = \mathbf{Z}_0 e^{wt}, \tag{4.20}$$

with w and the components of $\mathbf{Z}_0 = (Z_1, Z_2, Z_3, Z_4, Z_5, Z_6, Z_7, Z_8)$ in \mathbb{C} . Substituting a solution of the form (4.20) into the linearized system (4.19) of the endemic equilibrium \mathcal{E}_1^* yields the following system of linear equations

$$\begin{cases} wZ_1 = [-e_1 - k_1]Z_1 + (e_2 - e_1)Z_2 + (e_2 - e_1)Z_3 + (e_2 - e_1)Z_4 - e_1Z_5 + (\eta_1 e_2 - e_1)Z_6 + \frac{a_0 e_2}{\beta}Z_7 - e_1Z_8, \\ wZ_2 = kZ_1 - k_2Z_2, \\ wZ_3 = \eta Z_1 - k_3Z_3, \\ wZ_4 = bZ_1 + \theta Z_2 - k_4Z_4, \\ wZ_5 = \epsilon Z_1 - k_5Z_5, \\ wZ_6 = d_0Z_3 + d_1Z_5 - k_6Z_6, \\ wZ_7 = \omega_1Z_2 + \sigma Z_3 + \omega_0Z_4 + a_1Z_6 - \delta_5Z_7, \\ wZ_8 = \gamma Z_2 + \rho Z_3 + \nu Z_4 + \alpha Z_5 + rZ_7 - \mu Z_8, \end{cases} \tag{4.21}$$

where $k_i, i = 1, \dots, 6$, are given in (4.2).

Now, by solving the second, third and fifth equations of (4.21) for Z_2, Z_3 and Z_5 , and substituting the results into the other equations, we obtain the following system:

$$\begin{cases} l[1 + F_1(w)]Z_1 = (GZ)_1, & [1 + F_2(w)]Z_2 = (GZ)_2, & [1 + F_3(w)]Z_3 = (GZ)_3, & [1 + F_4(w)]Z_4 = (GZ)_4, \\ [1 + F_5(w)]Z_5 = (GZ)_5, & [1 + F_6(w)]Z_6 = (GZ)_6, & [1 + F_7(w)]Z_7 = (GZ)_7, & [1 + F_8(w)]Z_8 = (GZ)_8, \end{cases} \tag{4.22}$$

where

$$F_1(w) = \frac{w}{k_1} + \frac{e_1}{k_1} \left[1 + \frac{k}{w+k_2} + \frac{\eta}{w+k_3} + \frac{\epsilon}{w+k_5} \right] + \frac{e_1}{k_1(w+k_4)} \left[b + \frac{k\theta}{w+k_2} \right] + \frac{e_1}{k_1(w+k_6)} \left[\frac{d_0\eta}{w+k_3} + \frac{\epsilon d_1}{w+k_5} \right] + \frac{e_1}{k_1(w+\mu)} \left\{ \frac{k\gamma}{w+k_2} + \frac{\rho\eta}{w+k_3} + \frac{\nu}{w+k_4} \left[b + \frac{k\theta}{w+k_2} \right] \right\} + \frac{e_1}{k_1(w+\mu)} \left\{ \frac{\epsilon\alpha}{w+k_5} + \frac{r}{w+\delta_5} \left[\frac{k\omega_1}{w+k_2} + \frac{\sigma\eta}{w+k_3} + \frac{\epsilon a_1}{w+k_5} + \frac{\omega_0}{w+k_4} \left(b + \frac{k\theta}{w+k_2} \right) \right] \right\},$$

$$F_2(w) = \frac{w}{k_2}, \quad F_3(w) = \frac{w}{k_3}, \quad F_4(w) = \frac{w}{k_4}, \quad F_5(w) = \frac{w}{k_5},$$

$$F_6(w) = \frac{w}{k_6}, \quad F_7(w) = \frac{w}{\delta_5}, \quad F_8(w) = \frac{w}{\mu}$$

and

$$G = \begin{pmatrix} 0 & \frac{e_2}{k_1} & \frac{e_2}{k_1} & \frac{e_2}{k_1} & 0 & \frac{\eta_1 e_2}{k_1} & \frac{a_0 e_2}{\beta k_1} & 0 \\ \frac{k}{k_2} & 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ \frac{\eta}{k_3} & 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ \frac{b}{k_4} & \frac{\theta}{k_4} & 0 & 0 & 0 & 0 & 0 & 0 \\ \frac{\epsilon}{k_5} & 0 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & \frac{d_0}{k_6} & 0 & \frac{d_1}{k_6} & 0 & 0 & 0 \\ 0 & \frac{\omega_1}{\delta_5} & \frac{\sigma}{\delta_5} & \frac{\omega_0}{\delta_5} & 0 & \frac{a_1}{\delta_5} & 0 & 0 \\ 0 & \frac{\gamma}{\mu} & \frac{\rho}{\mu} & \frac{\nu}{\mu} & \frac{\alpha}{\mu} & \frac{r}{\mu} & 0 & 0 \end{pmatrix}.$$

Note that, the non-zero entries of the matrix G are positive, and the equilibrium $\mathcal{E}_1^* = (E^{**}, A^{**}, I^{**}, U^{**}, Q^{**}, H^{**}, V^{**}, R^{**})$ satisfies $\mathcal{E}_1^* = GZ_1^*$. Here $(GZ)_i, i = 1, \dots, 8$, denotes the i th component of the vector matrix GZ . Since the components of \mathcal{E}_1^* are all positive, then if Z represents any solution of system (4.22), there is a minimal positive real number c_0 (see (Esteva et al., 2009; Esteva and Vargas, 2000; Safi and Gumel, 2010) and the references therein) such that

$$|Z| \leq c_0 \mathcal{E}_1^*, \tag{4.23}$$

where $|Z| = (|Z_1|, |Z_2|, |Z_3|, |Z_4|, |Z_5|, |Z_6|, |Z_7|, |Z_8|)$, and $|z|$ denotes the modulus of the complex number z . In fact, the goal is to prove that $\text{Re } w < 0$. This is done by contradiction. To do so, we assume that $\text{Re } w \geq 0$.

First, we assume that $w = 0$.

This case directly implies that (4.21) is a homogeneous linear system in the variables $Z_i, i = 1, \dots, 5$.

So, computing the determinant of this system yields

$$\begin{aligned} \Delta &= B + \left[1 - \frac{\beta s}{\mu \mathcal{R}_c} \left(\frac{k}{k_1 k_2} + \frac{\eta}{k_1 k_3} + \frac{b}{k_1 k_4} + \frac{\theta k}{k_1 k_2 k_4} + \frac{\eta_1 \eta d_0}{k_1 k_3 k_6} + \frac{\eta_1 \epsilon d_1}{k_1 k_5 k_6} + \frac{k \theta a_0 \omega_0}{\beta k_1 k_2 k_4 \delta_5} \right. \right. \\ &\quad \left. \left. + \frac{k a_0 \omega_1}{\beta k_1 k_2 \delta_5} + \frac{\eta d_0 a_0 a_1}{\beta k_1 k_3 k_6 \delta_5} + \frac{\eta \sigma a_0}{\beta k_1 k_3 \delta_5} + \frac{b a_0 \omega_0}{\beta k_1 k_4 \delta_5} + \frac{\epsilon d_1 a_0 a_1}{\beta k_1 k_5 k_6 \delta_5} \right) \right] \mu k_1 k_2 k_3 k_4 k_5 k_6 \delta_5, \\ &= B + \left(1 - \frac{\mathcal{R}_c}{\mathcal{R}_c} \right) \mu k_1 k_2 k_3 k_4 k_5 k_6 \delta_5, \\ &= B, \end{aligned} \tag{4.24}$$

where

$$\begin{aligned} B &= \epsilon \mu d_1 k_2 k_3 k_4 \delta_5 e_1 + \epsilon r d_1 k_2 k_3 k_4 \delta_5 e_1 + \epsilon \mu k_2 k_3 k_4 k_6 \delta_5 e_1 + \epsilon \alpha k_2 k_3 k_4 k_6 \delta_5 e_1 \\ &\quad + \mu k_2 k_3 k_4 k_5 k_6 \delta_5 e_1 + \mu \theta k k_3 k_5 k_6 \delta_5 e_1 + k \theta \nu k_3 k_5 k_6 \delta_5 e_1 + \mu k k_3 k_4 k_5 k_6 \delta_5 e_1 \\ &\quad + \gamma k k_3 k_4 k_5 k_6 \delta_5 e_1 + \mu b k_2 k_3 k_5 k_6 \delta_5 e_1 + b \nu k_2 k_3 k_5 k_6 \delta_5 e_1 + \mu d_0 \eta k_2 k_4 k_5 \delta_5 e_1 \\ &\quad + r d_0 \eta k_2 k_4 k_5 \delta_5 e_1 + \mu \eta k_2 k_4 k_5 k_6 \delta_5 e_1 + \eta \rho k_2 k_4 k_5 k_6 \delta_5 e_1 > 0. \end{aligned}$$

Thus, one has $\Delta = B > 0$. Accordingly, system (4.21) has the vanishing solution $Z = 0$, which corresponds to the DFE, \mathcal{E}_0 , given in (4.1).

Now, we evaluate the second case $w \neq 0$.

Since we have assumed that $\text{Re } w > 0$, then, it follows clearly that $|1 + F_i(w)| > 1$, for all $i = 1, \dots, 8$. Define $F(w) = \min_{i \in \{1, \dots, 8\}} |1 + F_i(w)|$. Thus, $F(w) > 1$, and then $\frac{c_0}{F(w)} < c_0$. Note that c_0 is a minimal positive real number such that $|Z| \leq c_0 \mathcal{E}_1^*$. Hence, it follows from the minimality of c_0 that

$$|Z| > \frac{c_0}{F(w)} \mathcal{E}_1^*. \tag{4.25}$$

Now, by taking the norm on left and right sides of the third equation in (4.22), and using the fact that G is a non-negative matrix, we get

$$F(w)|Z_3| \leq |1 + F_3(w)||Z_3| = |(GZ)_3| \leq G|Z_3| \leq c_0 G(\mathcal{E}_1^*)_3 = c_0 I^{**}. \tag{4.26}$$

From (4.26), we obtain $|Z_3| \leq \frac{c_0}{F(w)} I^{**}$. This contradicts (4.25). Thus, $\text{Re } w < 0$, that is, all eigenvalues of the characteristic equation associated with the linearized system (4.19) around \mathcal{E}_1^* , have negative real parts. Thus the unique EEP, \mathcal{E}_1^* , is locally asymptotically stable whenever $\mathcal{R}_c > 1$. This completes the proof of Theorem 4.6. \square

Theorem 4.6 implies that, when $\mathcal{R}_c > 1$, COVID-19 will persist in the community if the initial sizes of the sub-populations, of the model, are in the basin of attraction of the EEP $\mathcal{E}_1^* = \mathcal{E}^*|_{N=N^*}$.

4.5. Global stability of the endemic equilibrium

The following Theorem provides the global stability result for the endemic equilibrium point, \mathcal{E}^* , of system (2.2).

Theorem 4.7. *The unique endemic equilibrium point of system (2.2) is globally asymptotically stable in $\Omega \setminus \Omega_0$ whenever $\mathcal{R}_c > 1$.*

Proof. Let $(S(t), E(t), A(t), I(t), U(t), Q(t), H(t), V(t), R(t))$ be any positive solution of system (2.2) in $\Omega \setminus \Omega_0$. Define the following Lyapunov function

$$\begin{aligned} \mathcal{M}_1(t) = & \frac{\delta_5 S^{**}}{\beta \eta_1} \left(\frac{S}{S^{**}} - 1 - \ln \frac{S}{S^{**}} \right) + \frac{\delta_5 E^{**}}{\beta \eta_1} \left(\frac{E}{E^{**}} - 1 - \ln \frac{E}{E^{**}} \right) \\ & + \frac{k_4 \beta \delta_5 S^{**} + k_4 a_0 \omega_1 S^{**} + a_0 \theta \omega_0 S^{**} + \beta \theta \delta_5 S^{**}}{\beta \eta_1 k_2 k_4} A^{**} \left(\frac{A}{A^{**}} - 1 - \ln \frac{A}{A^{**}} \right) \\ & + \frac{k_6 \beta \delta_5 S^{**} + a_0 a_1 d_0 S^{**} + a_0 \sigma k_6 S^{**} + \beta \eta_1 d_0 \delta_5 S^{**}}{\beta \eta_1 k_3 k_6} I^{**} \left(\frac{I}{I^{**}} - 1 - \ln \frac{I}{I^{**}} \right) \\ & + \frac{\beta \delta_5 S^{**} + a_0 \omega_0 S^{**}}{\beta \eta_1 k_4} U^{**} \left(\frac{U}{U^{**}} - 1 - \ln \frac{U}{U^{**}} \right) + \frac{\beta \eta_1 d_1 \delta_5 S^{**} + a_0 a_1 d_1 S^{**}}{\beta \eta_1 k_5 k_6} Q^{**} \left(\frac{Q}{Q^{**}} - 1 - \ln \frac{Q}{Q^{**}} \right) \\ & + \frac{\beta \eta_1 \delta_5 S^{**} + a_0 a_1 S^{**}}{\beta \eta_1 k_6} H^{**} \left(\frac{H}{H^{**}} - 1 - \ln \frac{H}{H^{**}} \right) + \frac{a_0 S^{**}}{\beta \eta_1} V^{**} \left(\frac{V}{V^{**}} - 1 - \ln \frac{V}{V^{**}} \right). \end{aligned}$$

Using the equilibrium conditions, after lengthy computations, the derivative of the above Lyapunov function computed along the solutions of system (2.2) is given below:

$$\begin{aligned} \frac{d\mathcal{M}_1(t)}{dt} = & \frac{s\delta_5}{\beta \eta_1} \left(2 - \frac{S}{S^{**}} - \frac{S^{**}}{S} \right) + \frac{S^{**} \delta_5}{\eta_1} \left(3 - \frac{S^{**}}{S} - \frac{SIE^{**}}{ES^{**}I^{**}} - \frac{EI^{**}}{IE^{**}} \right) + \frac{S^{**} \delta_5 A^{**}}{\eta_1} \left(3 - \frac{S^{**}}{S} - \frac{SAE^{**}}{ES^{**}A^{**}} - \frac{EA^{**}}{AE^{**}} \right) \\ & + \frac{S^{**} a_0 \omega_1 A^{**}}{\beta \eta_1} \left(4 - \frac{S^{**}}{S} - \frac{EA^{**}}{AE^{**}} - \frac{AV^{**}}{VA^{**}} - \frac{SVE^{**}}{ES^{**}V^{**}} \right) + \frac{S^{**} a_0 \omega_0 \theta A^{**}}{\beta \eta_1 k_4} \left(5 - \frac{S^{**}}{S} - \frac{EA^{**}}{AE^{**}} - \frac{AU^{**}}{UA^{**}} - \frac{UV^{**}}{VU^{**}} - \frac{SVE^{**}}{ES^{**}V^{**}} \right) \\ & + \frac{S^{**} \theta \delta_5 A^{**}}{\eta_1 k_4} \left(4 - \frac{S^{**}}{S} - \frac{EA^{**}}{AE^{**}} - \frac{AU^{**}}{UA^{**}} - \frac{SUE^{**}}{ES^{**}U^{**}} \right) + \frac{S^{**} b \delta_5 E^{**}}{\eta_1 k_4} \left(3 - \frac{S^{**}}{S} - \frac{EU^{**}}{UE^{**}} - \frac{SUE^{**}}{ES^{**}V^{**}} \right) \\ & + \frac{S^{**} a_0 \omega_0 b E^{**}}{\beta \eta_1 k_4} \left(4 - \frac{S^{**}}{S} - \frac{EU^{**}}{UE^{**}} - \frac{UV^{**}}{VU^{**}} - \frac{SVE^{**}}{ES^{**}V^{**}} \right) + \frac{S^{**} a_0 a_1 d_0 I^{**}}{\beta \eta_1 k_6} \left(5 - \frac{S^{**}}{S} - \frac{EI^{**}}{IE^{**}} - \frac{IH^{**}}{HI^{**}} - \frac{HV^{**}}{VH^{**}} - \frac{SVE^{**}}{ES^{**}V^{**}} \right) \\ & + \frac{S^{**} a_0 \sigma I^{**}}{\beta \eta_1} \left(4 - \frac{S^{**}}{S} - \frac{EI^{**}}{IE^{**}} - \frac{IV^{**}}{VI^{**}} - \frac{SVE^{**}}{ES^{**}V^{**}} \right) + \frac{S^{**} d_0 \delta_5 I^{**}}{k_6} \left(4 - \frac{S^{**}}{S} - \frac{EI^{**}}{IE^{**}} - \frac{IH^{**}}{HI^{**}} - \frac{SHE^{**}}{ES^{**}H^{**}} \right) \\ & + \frac{S^{**} d_1 \delta_5 Q^{**}}{k_6} \left(4 - \frac{S^{**}}{S} - \frac{EQ^{**}}{QE^{**}} - \frac{QH^{**}}{HQ^{**}} - \frac{SHE^{**}}{ES^{**}H^{**}} \right) + \frac{S^{**} a_0 a_1 d_1 Q^{**}}{\beta \eta_1 k_6} \left(5 - \frac{S^{**}}{S} - \frac{EQ^{**}}{QE^{**}} - \frac{QH^{**}}{HQ^{**}} - \frac{HV^{**}}{VH^{**}} - \frac{SVE^{**}}{ES^{**}V^{**}} \right). \end{aligned}$$

Thus, by using the arithmetic-geometric means inequality and condition $\mathcal{R}_c > 1$, it follows that $\frac{d\mathcal{M}_1(t)}{dt} \leq 0$. Moreover, $\frac{d\mathcal{M}_1(t)}{dt} = 0$, holds if and only if $S = S^{**}, E = E^{**}, A = A^{**}, I = I^{**}, U = U^{**}, Q = Q^{**}, H = H^{**}, V = V^{**}$. Consequently, \mathcal{M}_1 is a Lyapunov function on $\Omega \setminus \Omega_0$. So, by LaSalle's invariance principle [12, Theorem 5.3.1], it follows that

$$\lim_{t \rightarrow \infty} (S(t), E(t), A(t), I(t), U(t), Q(t), H(t), V(t)) = (S^{**}, E^{**}, A^{**}, I^{**}, U^{**}, Q^{**}, H^{**}, V^{**}). \tag{4.27}$$

Again, combining this with system (2.2), gives $\lim_{t \rightarrow \infty} R(t) = R^{**}$ as described in the proof of Theorem 4.1. Thus, every solution of the model, with initial condition in $\Omega \setminus \Omega_0$, approaches the unique endemic equilibrium point of system (2.2) when t tends to ∞ for $\mathcal{R}_c > 1$. This completes the proof. \square

In other words, Theorem 4.7 shows that COVID-19 will persist in the community whenever $\mathcal{R}_c > 1$. Furthermore, it follows from Theorem 4.7 that an imperfect follow-up of patients tested positive could lead to infection of many people in the community. Fig. 2 below shows a good fit for total actual symptomatic infectious individuals and those predicted by the model (2.2).

4.6. Sensitivity analysis with respect to quarantine an hospitalization

Here we analyze the threshold quantity \mathcal{R}_c , around the parameters associated to the quarantine of exposed individuals (ϵ) and the hospitalization of individuals with COVID-19 symptoms (d_0), in order to measure the effect of quarantine and hospitalization on the transmission dynamics of the disease. For this, we compute the partial derivative of \mathcal{R}_c with respect to the aforementioned parameters. First, computing the partial derivative of \mathcal{R}_c with respect to ϵ , we obtain

$$\frac{\partial \mathcal{R}_c}{\partial \epsilon} = \frac{\beta \delta_5 s (d_1 k_{01} k_3 - \eta d_0 k_5) \eta_1 - s (B_0 - d_1 a_0 a_1 k_{01} k_3)}{\mu k_1^2 k_3 k_5 k_6 \delta_5}, \tag{4.28}$$

where

$$B_0 = \beta \eta k_5 k_6 \delta_5 + \eta d_0 a_0 a_1 k_5 + \eta \sigma a_0 k_5 k_6 + \frac{\beta k_3 k_5 k_6 \delta_5 k}{k_2} + \frac{\beta b k_3 k_5 k_6 \delta_5}{k_4} + \frac{\beta \theta k k_3 k_5 k_6 \delta_5}{k_2 k_4} + \frac{k \theta a_0 \omega_0 k_3 k_5 k_6}{k_2 k_4} + \frac{k a_0 \omega_1 k_3 k_5 k_6}{k_2} + \frac{b a_0 \omega_0 k_3 k_5 k_6}{k_4}.$$

It follows from (4.28) that

$$\frac{\partial \mathcal{R}_c}{\partial \epsilon} < 0 \quad \text{if and only if} \quad \eta_1 < \eta_{1\epsilon},$$

and

$$\frac{\partial \mathcal{R}_c}{\partial \epsilon} > 0 \quad \text{if and only if} \quad \eta_1 > \eta_{1\epsilon},$$

with

$$0 < \eta_{1\epsilon} = \frac{B_0 - d_1 a_0 a_1 k_{01} k_3}{\beta \delta_5 (d_1 k_{01} k_3 - \eta d_0 k_5)}. \tag{4.29}$$

This first evaluation implies that the quarantine of exposed individuals can reduce the control reproduction number, and COVID-19 will reduce burden if the relative infectiousness of hospitalized individuals, η_1 , does not exceed the threshold quantity $\eta_{1\epsilon}$. If $\eta_1 > \eta_{1\epsilon}$, the use of quarantine of exposed individuals will increase the control reproduction number, and COVID-19 will increase burden. Thus, the use of quarantine is injurious to the population.

The above discussion is summed up in the following result.

Lemma 4.8. *The use of quarantine of the exposed individuals will have positive impact on the population if $\eta_1 < \eta_{1\epsilon}$, and negative impact on the population whenever $\eta_1 > \eta_{1\epsilon}$.*

Similarly, the computation of the partial derivative of \mathcal{R}_c with respect d_0 , gives

$$\frac{\partial \mathcal{R}_c}{\partial d_0} = \frac{\beta \eta k_{03} \delta_5 s \eta_1 - s \eta (k_6 (\sigma a_0 + \beta \delta_5) - a_0 a_1 k_{03})}{\mu k_1 k_3^2 k_6 \delta_5}, \tag{4.30}$$

It follows from (4.30) that

$$\frac{\partial \mathcal{R}_c}{\partial d_0} < 0 \quad \text{if and only if} \quad \eta_1 < \eta_{1d_0},$$

and

$$\frac{\partial \mathcal{R}_c}{\partial d_0} > 0 \quad \text{if and only if} \quad \eta_1 > \eta_{1d_0},$$

with

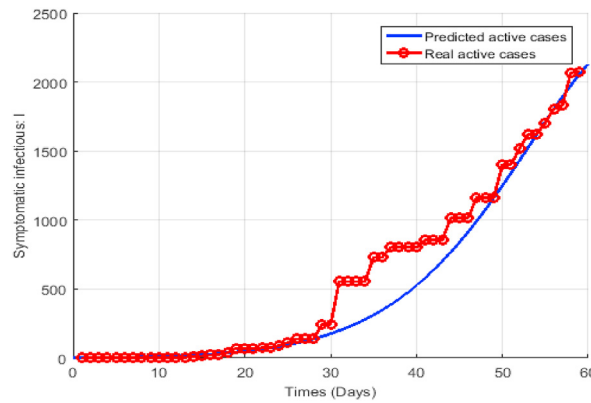


Fig. 2. Fitted results from the model (2.2) using the parameter values from Table 1 except the following parameters: $\beta = 1.55 \times 10^{-6}$, $\eta_1 = 0.49$, $a_0 = 10^{-7}$, $\mu = 1/59$, $d_1 = 0.156986$, and $\mathcal{R}_c = 1.2331 > 1$. Here, the red line indicates the real symptomatic infectious cases and the blue line indicates the predicted symptomatic infectious individuals.

$$0 < \eta_{1d_0} = \frac{k_6(\sigma a_0 + \beta \delta_5) - a_0 a_1 k_{03}}{\beta k_{03} \delta_5}. \tag{4.31}$$

This last evaluation implies that, the hospitalization of individuals with COVID-19 symptoms will be beneficial to the population if the relative infectiousness of hospitalized individuals does not exceed the threshold quantity η_{1d_0} , and is not beneficial if $\eta_1 > \eta_{1d_0}$. We have the following result.

Lemma 4.9. Hospitalization of individuals with COVID-19 symptoms will have positive impact on the population if $\eta_1 < \eta_{1d_0}$, and negative impact on the population if $\eta_1 > \eta_{1d_0}$.

Combining Lemma 4.3 and Lemma 4.4, we get the following result.

Theorem 4.10. The use of quarantine of exposed individual and hospitalization of individuals with COVID-19 symptoms will have

- a) positive impact on the population if $\eta_1 < \min\{\eta_{1\epsilon}, \eta_{1d_0}\}$;
- b) no impact on the population if $\eta_1 = \min\{\eta_{1\epsilon}, \eta_{1d_0}\}$;
- c) negative impact on the population if $\eta_1 > \max\{\eta_{1\epsilon}, \eta_{1d_0}\}$.

The first item of Theorem 4.4 means that the threshold quantity \mathcal{R}_c is a decreasing function of the quarantine and hospitalization parameters ϵ and d_0 , respectively; while the last item implies that \mathcal{R}_c is an increasing function of these parameters. The graph of Fig. 3 shows that the control reproduction number \mathcal{R}_c is a decreasing function of the quarantine rate ϵ and the hospitalization rate d_0 . This underscores the importance of the quarantine rate ϵ and the hospitalization rate d_0 in controlling the COVID-19 disease in Cameroon.

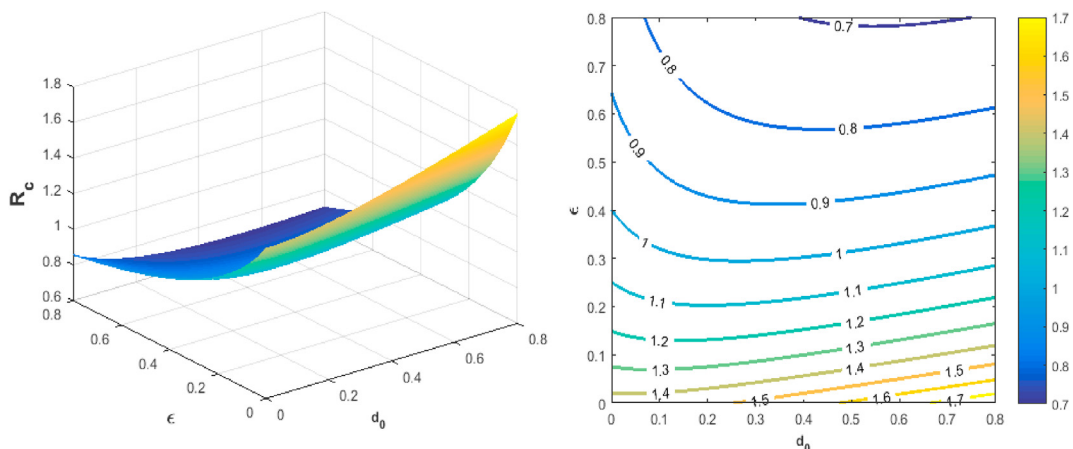


Fig. 3. Graph and contour plots of \mathcal{R}_c as a function of quarantine rate of exposed individuals ϵ and hospitalization rate of symptomatic infectious individuals d_0 .

5. Optimal control problem

COVID-19 has not yet been controlled and is still ongoing. Thus, to expect that the disease can stop, we need to comply with barrier measures (such as the regular washing of hands, the use of hydro-alcoholic gel, wearing face masks, social distancing rules). In this Section, we propose and investigate an optimal control problem applied to COVID-19 dynamics described by system (2.2) that we extend by adding three control functions u_1, u_2 and u_3 . The control u_1 denotes the quarantining rate of individuals who have been in contact with infected individuals and have accepted to be quarantined during a period of time (Yan et al., 2007). The term $\gamma_2 u_1$ denotes the rate of mandatory quarantine. In this case, the parameter ϵ becomes the natural quarantined rate. Next, the control function u_2 , which measures the rate of tracing, testing and hospitalization of people with clinical symptoms, moves infectious individuals from their symptomatic class to hospitalized class, under an hospitalization program for special medical treatment at rate γ_1 , with the natural hospitalization rate d_0 . Thus, u_2 decreases the evolution of symptomatic class to hospitalized class. The control u_3 represents the global effort of educational campaigns. The term $1 - u_3(t)$ is a decreasing factor that indicates the extent to which the production of unreported symptomatic individuals is blocked as a result of multiple educational campaigns. Furthermore, from the factor $1 - u_3(t)$, through the aforementioned barrier measures, people in the community can significantly reduce the concentration of virus in the environment. The flow diagram of the model with controls which elucidates the transmission phases of COVID-19 is presented in Fig. 4.

From the flow diagram in Fig. 4, we propose the following nonlinear system with control:

$$\left\{ \begin{aligned} \frac{dS}{dt} &= s - \beta S(I + A + U + \eta_1 H) - a_0 S V - \mu S, \\ \frac{dE}{dt} &= \beta S(I + A + U + \eta_1 H) + a_0 S V - (\mu + k + \epsilon + b + \eta + \gamma_2 u_1(t)) E, \\ \frac{dA}{dt} &= k E - (\mu + \delta_4 + \gamma + \theta(1 - u_3(t))) A, \\ \frac{dI}{dt} &= \eta E - (\mu + \delta_1 + \rho + d_0 + \gamma_1 u_2(t)) I, \\ \frac{dU}{dt} &= b E + \theta(1 - u_3(t)) A - (\mu + \delta_3 + \nu) U, \\ \frac{dQ}{dt} &= (\epsilon + \gamma_2 u_1(t)) E - (\mu + \alpha + d_1) Q, \\ \frac{dH}{dt} &= (d_0 + \gamma_1 u_2(t)) I + d_1 Q - (\mu + \delta_2 + r) H, \\ \frac{dV}{dt} &= \sigma(1 - u_3(t)) I + a_1(1 - u_3(t)) H + \omega_1(1 - u_3(t)) A + \omega_0(1 - u_3(t)) U - \delta_5 V, \\ \frac{dR}{dt} &= \rho I + \nu U + \gamma A + \alpha Q + r H - \mu R. \end{aligned} \right. \tag{5.1}$$

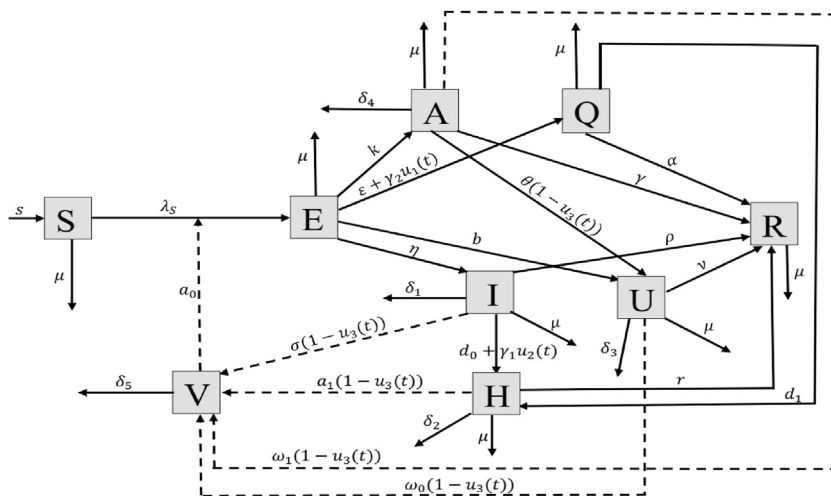


Fig. 4. Flow diagram of the model with controls.

All the parameters and classes of system (5.1) are the same as in system (2.2). The optimal control problem associated to model (5.1) requires the minimization of $E(t)$, $A(t)$, $I(t)$, $U(t)$, $Q(t)$, $H(t)$ and $V(t)$ as well as the cost of implementation of the interventions needed. Let T be a fixed terminal time. The objective functional which we seek to minimize is defined as in (Yan et al., 2007) as follows:

$$\mathcal{J}(u_1(t), u_2(t), u_3(t)) = \int_0^T [B_1E(t) + B_2A(t) + B_3I(t) + B_4U(t) + B_5Q(t) + B_6H(t) + B_7V(t) + \frac{1}{2}R_1u_1^2(t) + \frac{1}{2}R_2u_2^2(t) + \frac{1}{2}R_3u_3^2(t)] dt. \tag{5.2}$$

B_i , $i = 1, \dots, 7$ represent the cost coefficients for $E(t)$, $A(t)$, $I(t)$, $U(t)$, $Q(t)$, $H(t)$ and $V(t)$, respectively. R_1 , R_2 and R_3 are cost balancing coefficients associated with the hospitalized individuals in designated, susceptible quarantined individuals, and a strategy applied to the whole population.

The admissible controls set is defined as

$$\mathcal{F} = \{(u_1, u_2, u_3) : u_i \text{ is measurable, } 0 \leq u_i(t) \leq b_i, 0 < b_i \leq 1, t \in [0, T], i = 1, 2, 3\},$$

where b_i , $i = 1, 2, 3$, are fixed positive constant which depend on the amount of resources available for the implementation of the control strategies. We need to determine the optimal control (u_1^*, u_2^*, u_3^*) such that

$$\mathcal{J}(u_1^*, u_2^*, u_3^*) = \min\{\mathcal{J}(u_1, u_2, u_3) : (u_1, u_2, u_3) \in \mathcal{F}\}.$$

This is given in the following Theorem.

Theorem 5.1. Consider the control problem with objective functional (5.2) and system (5.1). Then, there exists an optimal control $u^* = (u_1^*, u_2^*, u_3^*) \in \mathcal{F}$ such that

$$\mathcal{J}(u_1^*, u_2^*, u_3^*) = \min_{(u_1, u_2, u_3) \in \mathcal{F}} \mathcal{J}(u_1, u_2, u_3),$$

provided the following conditions are satisfied:

- (a) The class of all initial conditions with controls $u = (u_1, u_2, u_3)$ in the set of admissible controls, with system (5.1) being satisfied, is not empty.
- (b) The set of admissible controls \mathcal{F} is convex and closed.
- (c) The right-hand side of system (5.1) is continuous, bounded from above by a sum of the bounded control and the state, and can be written as a linear function of controls (u_1, u_2, u_3) with coefficients depending on time and state.
- (d) The integrand of the objective functional (5.2) is convex on \mathcal{F} and bounded from below by $-e_0 + e_1(|u_1|^2 + |u_2|^2 + |u_3|^2)$, where $e_0 \geq 0$ and $e_1 > 0$.

proof. The proof is done by applying similar arguments as in the proof of Theorem 4.1 in (Fleming and Rishel, 1975). \square We now investigate the necessary conditions for the optimal control by using the Pontryagin's maximum principle (Pontryagin et al., 1962). Define the Lagrangian for this control problem as in (Joshi, 2002; Kirk, 2004):

$$\begin{aligned}
 &\mathcal{L}(S, E, A, I, U, Q, H, V, R, u_1, u_2, u_3, \lambda_1, \lambda_2, \lambda_3, \lambda_4, \\
 &\quad \lambda_5, \lambda_6, \lambda_7, \lambda_8, \lambda_9, w_{11}, w_{12}, w_{21}, w_{22}, w_{31}, w_{32}) \\
 &= [B_1E(t) + B_2A(t) + B_3I(t) + B_4U(t) + B_5Q(t) \\
 &\quad + B_6H(t) + B_7V(t) + \frac{1}{2}R_1u_1^2 + \frac{1}{2}R_2u_2^2 + \frac{1}{2}R_3u_3^2] \\
 &\quad + \lambda_1[s - \beta S(I(t) + A(t) + U(t) + \eta_1H(t)) - a_0S(t)V(t) - \mu S(t)] \\
 &\quad + \lambda_2[\beta S(I(t) + A(t) + U(t) + \eta_1H(t)) + a_0S(t)V(t) \\
 &\quad - (\mu + k + \epsilon + b + \eta + \gamma_2u_1(t))E(t)] \\
 &\quad + \lambda_3[kE(t) - (\mu + \delta_4 + \gamma + \theta(1 - u_3(t)))A(t)] \\
 &\quad + \lambda_4[\eta E(t) - (\mu + \delta_1 + \rho + d_0 + \gamma_1u_2(t))I(t)] \\
 &\quad + \lambda_5[bE(t) + \theta(1 - u_3(t))A(t) - (\mu + \delta_3 + \nu)U(t)] \\
 &\quad + \lambda_6[(\epsilon + \gamma_2u_1(t))E(t) - (\mu + \alpha + d_1)Q(t)] \\
 &\quad + \lambda_7[(d_0 + \gamma_1u_2(t))I(t) + d_1Q(t) - (\mu + \delta_2 + r)H(t)] \\
 &\quad + \lambda_8[\sigma(1 - u_3(t))I(t) + a_1(1 - u_3(t))H(t) + \omega_1(1 - u_3(t))A(t) \\
 &\quad + \omega_0(1 - u_3(t))U(t) - \delta_5V(t)] \\
 &\quad + \lambda_9[\rho I(t) + \nu U(t) + \gamma A(t) + \alpha Q(t) + rH(t) - \mu R(t)] \\
 &\quad - w_{11}(t)(b_1 - u_1(t)) - w_{12}(t)u_1(t) - w_{21}(t)(b_2 - u_2(t)) \\
 &\quad - w_{22}(t)u_2(t) - w_{31}(t)(b_3 - u_3(t)) - w_{32}(t)u_3(t),
 \end{aligned}$$

where $w_{11}(t), w_{12}(t), w_{21}(t), w_{22}(t), w_{31}(t), w_{32}(t) \geq 0$ are penalty multipliers satisfying the following equations at u^* :

$$\begin{aligned}
 w_{11}(t)(b_1 - u_1(t)) &= 0, & w_{12}(t)u_1(t) &= 0, \\
 w_{21}(t)(b_2 - u_2(t)) &= 0, & w_{22}(t)u_2(t) &= 0, \\
 w_{31}(t)(b_3 - u_3(t)) &= 0, & w_{32}(t)u_3(t) &= 0.
 \end{aligned}$$

Differentiating the above Lagrangian with respect to state variables $S, E, A, I, U, Q, H, V,$ and $R,$ respectively, and applying Pontryagin’s maximum principle, the adjoint system reads

$$\begin{aligned}
 \lambda'_1 &= -\frac{\partial \mathcal{L}}{\partial S}, & \lambda'_2 &= -\frac{\partial \mathcal{L}}{\partial E}, & \lambda'_3 &= -\frac{\partial \mathcal{L}}{\partial A}, & \lambda'_4 &= -\frac{\partial \mathcal{L}}{\partial I}, & \lambda'_5 &= -\frac{\partial \mathcal{L}}{\partial U}, \\
 \lambda'_6 &= -\frac{\partial \mathcal{L}}{\partial Q}, & \lambda'_7 &= -\frac{\partial \mathcal{L}}{\partial H}, & \lambda'_8 &= -\frac{\partial \mathcal{L}}{\partial V}, & \lambda'_9 &= -\frac{\partial \mathcal{L}}{\partial R},
 \end{aligned} \tag{5.3}$$

and the transversality conditions $\lambda_i(T) = 0, i = 1, \dots, 9,$ hold. Setting $\frac{\partial \mathcal{L}}{\partial u_i} = 0, i = 1, 2, 3,$ the optimality conditions are given by

$$\begin{aligned}
 \frac{\partial \mathcal{L}}{\partial u_1} &= R_1u_1 - \lambda_2\gamma_2E + \lambda_6\gamma_2E + w_{11}(t) - w_{12}(t) = 0, \\
 \frac{\partial \mathcal{L}}{\partial u_2} &= R_2u_2 - \lambda_4\gamma_1I + \lambda_7\gamma_1I + w_{21}(t) - w_{22}(t) = 0, \\
 \frac{\partial \mathcal{L}}{\partial u_3} &= R_3u_3 + \lambda_3\theta A - \lambda_5\theta A - \lambda_8\sigma I - \lambda_8a_1H - \lambda_8\omega_1A \\
 &\quad - \lambda_8\omega_0U + w_{31}(t) - w_{32}(t) = 0.
 \end{aligned}$$

The resolution of the above equations gives the following optimal controls

$$\begin{aligned}
 u_1^*(t) &= \frac{(\lambda_2 - \lambda_6)\gamma_2 E - w_{11}(t) + w_{12}(t)}{R_1}, \\
 u_2^*(t) &= \frac{(\lambda_4 - \lambda_7)\gamma_1 I - w_{21}(t) + w_{22}(t)}{R_2}, \\
 u_3^*(t) &= \frac{(\lambda_3 - \lambda_5)\theta A + \lambda_8(\sigma I + a_1 H + \omega_1 A + \omega_0 U) - w_{31}(t) + w_{32}(t)}{R_3}.
 \end{aligned}$$

For the explicit expression of the optimal control u_1^* on $[0, b_1]$, we consider three cases. First, when $u_1^*(t) = 0$, we have $w_{11}(t) = 0$. It then follows that

$$0 = u_1^*(t) = \frac{(\lambda_2 - \lambda_6)\gamma_2 E + w_{12}(t)}{R_1}.$$

Owing to $w_{12} \geq 0$, it follows that $\frac{(\lambda_2 - \lambda_6)\gamma_2 E}{R_1} \leq 0$.

Next, when $0 < u_1^* < b_1$, it follows that $w_{11}(t) = 0$ and $w_{12} = 0$. Consequently, one has

$$u_1^*(t) = \frac{(\lambda_2 - \lambda_6)\gamma_2 E}{R_1}.$$

Finally, when $u_1^*(t) = b_1$, one gets $w_{12}(t) = 0$. Thus,

$$b_1 = u_1^*(t) = \frac{(\lambda_2 - \lambda_6)\gamma_2 E - w_{11}(t)}{R_1}.$$

This means that $R_1 b_1 = (\lambda_2 - \lambda_6)\gamma_2 E - w_{11}(t)$, so that $w_{11}(t) = (\lambda_2 - \lambda_6)\gamma_2 E - R_1 \geq 0$. Thus $\frac{(\lambda_2 - \lambda_6)\gamma_2 E}{R_1} \geq b_1$. From the above discussion, we obtain

$$u_1^* = \begin{cases} \frac{(\lambda_2 - \lambda_6)\gamma_2 E}{R_1}, & \text{if } 0 < \frac{(\lambda_2 - \lambda_6)\gamma_2 E}{R_1} < b_1 \\ 0, & \text{if } \frac{(\lambda_2 - \lambda_6)\gamma_2 E}{R_1} \leq 0 \\ b_1, & \text{if } \frac{(\lambda_2 - \lambda_6)\gamma_2 E}{R_1} \geq b_1. \end{cases}$$

This can also be written under the following compact form

$$u_1^*(t) = \min \left\{ \max \left\{ 0, \frac{(\lambda_2 - \lambda_6)\gamma_2 E}{R_1} \right\}, b_1 \right\}.$$

Similarly, we get the following expressions for the second and third optimal control

$$\begin{aligned}
 u_2^*(t) &= \min \left\{ \max \left\{ 0, \frac{(\lambda_4 - \lambda_7)\gamma_1 I}{R_2} \right\}, b_2 \right\}, \\
 u_3^*(t) &= \min \left\{ \max \left\{ 0, \frac{(\lambda_3 - \lambda_5)\theta A + \lambda_8(\sigma I + a_1 H + \omega_1 A + \omega_0 U)}{R_3} \right\}, b_3 \right\}.
 \end{aligned}$$

Summarizing the above characterization we obtain the following result.

Theorem 5.2. Given the optimal controls (u_1^*, u_2^*, u_3^*) and the existence of solutions of system (5.1), there exist adjoint variables λ_i , $i = 1, \dots, 9$ satisfying the adjoint equations (5.3) together with the transversality conditions $\lambda_i(T) = 0$, for $i = 1, \dots, 9$. Furthermore, the optimal controls $u_1^*(t)$, $u_2^*(t)$ and $u_3^*(t)$ are characterized as

$$u_1^*(t) = \min \left\{ \max \left\{ 0, \frac{(\lambda_2 - \lambda_6)\gamma_2 E}{R_1} \right\}, b_1 \right\}, \tag{5.4}$$

$$u_2^*(t) = \min \left\{ \max \left\{ 0, \frac{(\lambda_4 - \lambda_7)\gamma_1 I}{R_2} \right\}, b_2 \right\}, \tag{5.5}$$

$$u_3^*(t) = \min \left\{ \max \left\{ 0, \frac{(\lambda_3 - \lambda_5)\theta A + \lambda_8(\sigma I + a_1 H + \omega_1 A + \omega_0 U)}{R_3} \right\}, b_3 \right\}. \tag{5.6}$$

We finally deal with the uniqueness for the optimality system including system (5.1) and adjoint equation (5.3). To do this, we need the following result.

Lemma 5.3. (Garira et al., 2005; Joshi, 2002) *The function $u_1^*(\varphi) = \min\{\max\{\varphi, a_2^1\}, b_2^1\}$ is Lipschitz continuous with respect to φ , where a_2^1 and b_2^1 are two arbitrary fixed positive constants, with $a_2^1 < b_2^1$.*

Now, from the fact that the state variables are uniformly bounded, it can easily be shown that the adjoint variables have finite upper bounds. The uniqueness result for the optimality system states as follows.

Theorem 5.4. *Bounded solutions of the optimality system are unique for a sufficiently small $T > 0$.*

Proof. Let $(S, E, A, I, U, Q, H, V, R, \lambda_1, \lambda_2, \lambda_3, \lambda_4, \lambda_5, \lambda_6, \lambda_7, \lambda_8, \lambda_9)$ and $(\bar{S}, \bar{E}, \bar{A}, \bar{I}, \bar{U}, \bar{Q}, \bar{H}, \bar{V}, \bar{R}, \bar{\lambda}_1, \bar{\lambda}_2, \bar{\lambda}_3, \bar{\lambda}_4, \bar{\lambda}_5, \bar{\lambda}_6, \bar{\lambda}_7, \bar{\lambda}_8, \bar{\lambda}_9)$ be two solutions of the optimality system. Set $S = e^{\sigma t} p_1, E = e^{\sigma t} p_2, A = e^{\sigma t} p_3, I = e^{\sigma t} p_4, U = e^{\sigma t} p_5, Q = e^{\sigma t} p_6, H = e^{\sigma t} p_7, V = e^{\sigma t} p_8, R = e^{\sigma t} p_9, \lambda_1 = e^{-\sigma t} q_1, \lambda_2 = e^{-\sigma t} q_2, \lambda_3 = e^{-\sigma t} q_3, \lambda_4 = e^{-\sigma t} q_4, \lambda_5 = e^{-\sigma t} q_5, \lambda_6 = e^{-\sigma t} q_6, \lambda_7 = e^{-\sigma t} q_7, \lambda_8 = e^{-\sigma t} q_8, \lambda_9 = e^{-\sigma t} q_9$. Analogously, let $\bar{S} = e^{\sigma t} \bar{p}_1, \bar{E} = e^{\sigma t} \bar{p}_2, \bar{A} = e^{\sigma t} \bar{p}_3, \bar{I} = e^{\sigma t} \bar{p}_4, \bar{U} = e^{\sigma t} \bar{p}_5, \bar{Q} = e^{\sigma t} \bar{p}_6, \bar{H} = e^{\sigma t} \bar{p}_7, \bar{V} = e^{\sigma t} \bar{p}_8, \bar{R} = e^{\sigma t} \bar{p}_9, \bar{\lambda}_1 = e^{-\sigma t} \bar{q}_1, \bar{\lambda}_2 = e^{-\sigma t} \bar{q}_2, \bar{\lambda}_3 = e^{-\sigma t} \bar{q}_3, \bar{\lambda}_4 = e^{-\sigma t} \bar{q}_4, \bar{\lambda}_5 = e^{-\sigma t} \bar{q}_5, \bar{\lambda}_6 = e^{-\sigma t} \bar{q}_6, \bar{\lambda}_7 = e^{-\sigma t} \bar{q}_7, \bar{\lambda}_8 = e^{-\sigma t} \bar{q}_8, \bar{\lambda}_9 = e^{-\sigma t} \bar{q}_9$, where σ is a positive constant. Define

$$u_1^*(t) = \min \left\{ \max \left\{ 0, \frac{(\lambda_2 - \lambda_6)\gamma_2 E}{R_1} \right\}, b_1 \right\},$$

$$u_2^*(t) = \min \left\{ \max \left\{ 0, \frac{(\lambda_4 - \lambda_7)\gamma_1 I}{R_2} \right\}, b_2 \right\},$$

$$u_3^*(t) = \min \left\{ \max \left\{ 0, \frac{(\lambda_3 - \lambda_5)\theta A + \lambda_8(\sigma I + a_1 H + \omega_1 A + \omega_0 U)}{R_3} \right\}, b_3 \right\},$$

and

$$\bar{u}_1^*(t) = \min \left\{ \max \left\{ 0, \frac{(\bar{\lambda}_2 - \bar{\lambda}_6)\gamma_2 \bar{E}}{R_1} \right\}, b_1 \right\},$$

$$\bar{u}_2^*(t) = \min \left\{ \max \left\{ 0, \frac{(\bar{\lambda}_4 - \bar{\lambda}_7)\gamma_1 \bar{I}}{R_2} \right\}, b_2 \right\},$$

$$\bar{u}_3^*(t) = \min \left\{ \max \left\{ 0, \frac{(\bar{\lambda}_3 - \bar{\lambda}_5)\theta \bar{A} + \bar{\lambda}_8(\sigma \bar{I} + a_1 \bar{H} + \omega_1 \bar{A} + \omega_0 \bar{U})}{R_3} \right\}, b_3 \right\}.$$

Then, from Lemma 5.3, it follows that

$$|u_1^*(t) - \bar{u}_1^*(t)| \leq \left| \frac{\gamma_2}{R_1} \{p_2(q_2 - q_6) - \bar{p}_2(\bar{q}_2 - \bar{q}_6)\} \right|,$$

$$= \frac{\gamma_2}{R_1} |p_2(q_2 - q_6) - \bar{p}_2(\bar{q}_2 - \bar{q}_6)|,$$

$$|u_2^*(t) - \bar{u}_2^*(t)| \leq \left| \frac{\gamma_1}{R_2} \{p_4(q_4 - q_7) - \bar{p}_4(\bar{q}_4 - \bar{q}_7)\} \right|,$$

$$= \frac{\gamma_1}{R_2} |p_4(q_4 - q_7) - \bar{p}_4(\bar{q}_4 - \bar{q}_7)|,$$

and

$$\begin{aligned}
 |u_3^*(t) - \bar{u}_3^*(t)| &\leq \frac{1}{R_3} \{ \theta p_3(q_3 - q_5) - \bar{p}_3(\bar{q}_3 - \bar{q}_5) + \sigma(p_4q_8 - \bar{p}_4\bar{q}_8) \\
 &\quad + a_1(p_7q_8 - \bar{p}_7\bar{q}_8) + \omega_1(p_3q_8 - \bar{p}_3\bar{q}_8) + \omega_0(p_5q_8 - \bar{p}_5\bar{q}_8) \}, \\
 &= \frac{1}{R_3} | \theta p_3(q_3 - q_5) - \bar{p}_3(\bar{q}_3 - \bar{q}_5) + \sigma(p_4q_8 - \bar{p}_4\bar{q}_8) \\
 &\quad + a_1(p_7q_8 - \bar{p}_7\bar{q}_8) + \omega_1(p_3q_8 - \bar{p}_3\bar{q}_8) + \omega_0(p_5q_8 - \bar{p}_5\bar{q}_8) |.
 \end{aligned}$$

Inserting $S = e^{\sigma t} p_1$ in the first equation of (5.1) yields

$$p_1' + \varpi p_1 = se^{-\sigma t} - \beta p_1(p_3 + p_4 + p_5 + \eta_1 p_7) e^{\sigma t} - a_0 p_1 p_8 e^{\sigma t} - \mu p_1. \tag{5.7}$$

In a similar way for the other state and adjoint variables, we obtain

$$p_2' + \varpi p_2 = \beta p_1(p_3 + p_4 + p_5 + \eta_1 p_7) e^{\sigma t} + a_0 p_1 p_8 e^{\sigma t} - (\mu + k + \epsilon + b + \eta + \gamma_2 u_1^*) p_2, \tag{5.8}$$

$$p_3' + \varpi p_3 = k p_2 - (\mu + \delta_4 + \gamma + \theta(1 - u_3^*)) p_3, \tag{5.9}$$

$$p_4' + \varpi p_4 = \eta p_2 - (\mu + \delta_1 + \rho + d_0 + \gamma_1 u_2^*) p_4, \tag{5.10}$$

$$p_5' + \varpi p_5 = b p_2 + \theta(1 - u_3^*) p_3 - (\mu + \delta_3 + \nu) p_5, \tag{5.11}$$

$$p_6' + \varpi p_6 = (\epsilon + \gamma_2 u_1^*) p_2 - (\mu + \alpha + d_1) p_6, \tag{5.12}$$

$$p_7' + \varpi p_7 = (d_0 + \gamma_1 u_2^*) p_4 + d_1 p_6 - (\mu + \delta_2 + r) p_7, \tag{5.13}$$

$$p_8' + \varpi p_8 = \sigma(1 - u_3^*) p_4 + a_1(1 - u_3^*) p_7 + \omega_1(1 - u_3^*) p_3 + \omega_0(1 - u_3^*) p_5 - \delta_5 p_8, \tag{5.14}$$

$$p_9' + \varpi p_9 = \rho p_4 + \nu p_5 + \gamma p_3 + \alpha p_6 + r p_7 - \mu p_9, \tag{5.15}$$

$$\begin{aligned}
 -q_1' + \varpi q_1 &= -\mu q_1 - \beta q_1(p_3 + p_4 + p_5 + \eta_1 p_7) e^{\sigma t} + a_0 q_1 p_8 e^{\sigma t} \\
 &\quad + \beta q_2(p_3 + p_4 + p_5 + \eta_1 p_7) e^{\sigma t} + a_0 q_2 p_8 e^{\sigma t},
 \end{aligned} \tag{5.16}$$

$$\begin{aligned}
 -q_2' + \varpi q_2 &= B_1 e^{\sigma t} - (\mu + k + \epsilon + b + \eta + \gamma_2 u_1^*) q_2 \\
 &\quad + k q_3 + \eta q_4 + b q_5 + (\epsilon + \gamma_2 u_1^*) q_6,
 \end{aligned} \tag{5.17}$$

$$\begin{aligned}
 -q_3' + \varpi q_3 &= B_2 e^{\sigma t} - \beta p_1 q_1 e^{\sigma t} - (\mu + \delta_4 + \gamma + \theta(1 - u_3^*)) q_3 \\
 &\quad + \beta p_1 q_2 e^{2\sigma t} + \theta(1 - u_3^*) q_5 + \omega_1(1 - u_3^*) q_8 + \gamma q_9,
 \end{aligned} \tag{5.18}$$

$$\begin{aligned}
 -q_4' + \varpi q_4 &= B_3 e^{\sigma t} - \beta p_1 q_1 e^{\sigma t} - (\mu + \delta_1 + \rho + d_0 + \gamma_1 u_2^*) q_4 \\
 &\quad + \beta p_1 q_2 e^{\sigma t} + (d_0 + \gamma_1 u_2^*) q_7 + \sigma(1 - u_3^*) q_8 + \rho q_9,
 \end{aligned} \tag{5.19}$$

$$\begin{aligned}
 -q_5' + \varpi q_5 &= B_4 e^{\sigma t} - \beta p_1 q_1 e^{\sigma t} - (\mu + \delta_3 + \nu) q_5 \\
 &\quad + \beta p_1 q_2 e^{\sigma t} + \omega_0(1 - u_3) q_8 + \nu q_9,
 \end{aligned} \tag{5.20}$$

$$-q_6' + \varpi q_6 = B_5 e^{\sigma t} - (\mu + \alpha + d_1) q_6 + d_1 q_7 + \alpha q_9 \tag{5.21}$$

$$\begin{aligned}
 -q_7' + \varpi q_7 &= B_6 e^{\sigma t} - \beta \eta_1 p_1 q_1 e^{\sigma t} - (\mu + \delta_2 + r) q_7 \\
 &\quad + \beta \eta_1 p_1 q_2 e^{\sigma t} + a_1(1 - u_3^*) q_8 - r q_9,
 \end{aligned} \tag{5.22}$$

$$-q'_8 + \varpi q_8 = B_7 e^{\varpi t} - a_0 p_1 q_1 e^{\varpi t} + a_0 p_1 q_2 e^{\varpi t} - \delta_5 q_8, \tag{5.23}$$

$$-q'_9 + \varpi q_9 = -\mu q_9. \tag{5.24}$$

Also, introducing $\bar{S} = e^{\varpi t} \bar{p}_1$ in the first equation of (5.1) yields

$$\bar{p}'_1 + \varpi \bar{p}_1 = se^{-\varpi t} - \beta \bar{p}_1 (\bar{p}_3 + \bar{p}_4 + \bar{p}_5 + \eta_1 \bar{p}_7) e^{\varpi t} - a_0 \bar{p}_1 \bar{p}_8 e^{\varpi t} - \mu \bar{p}_1. \tag{5.25}$$

Then, the result is obtained by subtracting the equations for S and \bar{S} , E and \bar{E} , A and \bar{A} , I and \bar{I} , U and \bar{U} , Q and \bar{Q} , H and \bar{H} , V and \bar{V} , R and \bar{R} , λ_1 and $\bar{\lambda}_1$, λ_2 and $\bar{\lambda}_2$, λ_3 and $\bar{\lambda}_3$, λ_4 and $\bar{\lambda}_4$, λ_5 and $\bar{\lambda}_5$, λ_6 and $\bar{\lambda}_6$, λ_7 and $\bar{\lambda}_7$, λ_8 and $\bar{\lambda}_8$, λ_9 and $\bar{\lambda}_9$, multiplying each resulting equation by an appropriate difference of functions, and integrating from 0 to T . For example, subtracting equation (5.7) from (5.25) leads to

$$(p_1 - \bar{p}_1)' + \varpi(p_1 - \bar{p}_1) = -\beta e^{\varpi t} [p_1(p_3 + p_4 + p_5 + \eta_1 p_7) - \bar{p}_1(\bar{p}_3 + \bar{p}_4 + \bar{p}_5 + \eta_1 \bar{p}_7)] - a_0 e^{\varpi t} [p_1 p_8 - \bar{p}_1 \bar{p}_8] - \mu(p_1 - \bar{p}_1). \tag{5.26}$$

Multiplying the left and right hand sides of (5.26) by $(p_1 - \bar{p}_1)$ and integrating from 0 to T gives

$$\begin{aligned} & \frac{1}{2}(p_1 - \bar{p}_1)^2 + \varpi \int_0^T (p_1 - \bar{p}_1)^2 dt \\ & \leq \mu \int_0^T (p_1 - \bar{p}_1)^2 dt + C_1 e^{\varpi T} \int_0^T [(p_1 - \bar{p}_1)^2 + (p_3 - \bar{p}_3)^2 \\ & \quad + (p_4 - \bar{p}_4)^2 + (p_5 - \bar{p}_5)^2 + (p_7 - \bar{p}_7)^2 + (p_8 - \bar{p}_8)^2] dt, \end{aligned}$$

where the constant C_1 depends on the coefficients and the bounds on state variables p_1, p_3, p_4, p_5 and p_7 . Noting that $e^{\varpi T} \leq e^{3\varpi T}$, we get

$$\begin{aligned} & \frac{1}{2}(p_1 - \bar{p}_1)^2 + \varpi \int_0^T (p_1 - \bar{p}_1)^2 dt \\ & \leq C'_1 \int_0^T [(p_1 - \bar{p}_1)^2 dt + C'_2 e^{3\varpi T} \int_0^T [(p_1 - \bar{p}_1)^2 + (p_3 - \bar{p}_3)^2 \\ & \quad + (p_4 - \bar{p}_4)^2 + (p_5 - \bar{p}_5)^2 + (p_7 - \bar{p}_7)^2 + (p_8 - \bar{p}_8)^2] dt. \end{aligned} \tag{5.27}$$

C'_1 and C'_2 depend on the coefficients and the upper bounds of state variables $p_1, p_3, p_4, p_5, p_7, p_7$. Now, introducing $\bar{\lambda}_2 = e^{-\varpi t} \bar{q}_2$ in the second equation of (5.3), we obtain

$$\begin{aligned} -\bar{q}'_2 + \varpi \bar{q}_2 &= B_1 e^{\varpi t} - (\mu + k + \epsilon + b + \eta + \gamma_2 \bar{u}_1^*) q_2 \\ & \quad + k \bar{q}_3 + \eta \bar{q}_4 + b \bar{q}_5 + (\epsilon + \gamma_2 \bar{u}_1^*) \bar{q}_6. \end{aligned} \tag{5.28}$$

Subtracting equation (5.17) from (5.27), yields

$$\begin{aligned} -(q_1 - \bar{q}_2)' + \varpi(q_2 - \bar{q}_2) &= -[(k_1 + \gamma_2 \bar{u}_1^*) q_2 - (k_1 + \gamma_2 \bar{u}_1^*) \bar{q}_2] \\ & \quad + k(q_3 - \bar{q}_3) + \eta(q_4 - \bar{q}_4) + b(q_5 - \bar{q}_5) + [(\epsilon + \gamma_2 \bar{u}_1^*) q_6 - (\epsilon + \gamma_2 \bar{u}_1^*) \bar{q}_6], \end{aligned} \tag{5.29}$$

where $k_1 = \mu + k + \epsilon + b + \eta$. Multiplying the left and right hand sides of (5.28) by $(q_2 - \bar{q}_2)$ and integrating from 0 to T gives

$$\begin{aligned} & \frac{1}{2}(q_2(0) - \bar{q}_2(0))^2 + \varpi \int_0^T (q_2 - \bar{q}_2)^2 dt \\ &= -k_1 \int_0^T (q_2 - \bar{q}_2)^2 dt - \gamma_2 \int_0^T (u_1^* q_2 - \bar{u}_1^* \bar{q}_2)(q_2 - \bar{q}_2) dt \\ & \quad + k \int_0^T (q_2 - \bar{q}_2)(q_3 - \bar{q}_3) dt + \eta \int_0^T (q_2 - \bar{q}_2)(q_4 - \bar{q}_4) dt \\ & \quad + b \int_0^T (q_2 - \bar{q}_2)(q_5 - \bar{q}_5) dt + \epsilon \int_0^T (q_2 - \bar{q}_2)(q_6 - \bar{q}_6) dt + \gamma_2 \int_0^T (u_1^* q_6 - \bar{u}_1^* \bar{q}_6)(q_2 - \bar{q}_2) dt. \end{aligned}$$

The last term of the above equation reads

$$\begin{aligned} & \gamma_2 \int_0^T (u_1^* q_6 - \bar{u}_1^* \bar{q}_6)(q_2 - \bar{q}_2) dt = \gamma_2 \int_0^T u_1^*(q_6 - \bar{q}_6)(q_2 - \bar{q}_2) + \bar{q}_2(u_1^* - \bar{u}_1^*)(q_2 - \bar{q}_2) dt, \\ & \leq C_{01} \int_0^T [(q_2 - \bar{q}_2)^2 + (q_6 - \bar{q}_6)^2 + (u_1^* - \bar{u}_1^*)^2] dt. \end{aligned} \tag{5.30}$$

Now

$$\begin{aligned} & \int_0^T (u_1^* - \bar{u}_1^*)^2 dt \leq \left(\frac{\gamma_2}{R_1}\right)^2 \int_0^T [p_2(q_2 - q_6) - \bar{p}_2(\bar{q}_2 - \bar{q}_6)]^2 dt, \\ & = \left(\frac{\gamma_2}{R_1}\right)^2 \int_0^T [p_2^2(q_2 - q_6)^2 - 2p_2(q_2 - q_6)\bar{p}_2(\bar{q}_2 - \bar{q}_6) + \bar{p}_2^2(\bar{q}_2 - \bar{q}_6)^2] dt, \\ & \leq C_{03} \left(\frac{\gamma_2}{R_1}\right)^2 \int_0^T [(q_2 - \bar{q}_2)^2 + (q_6 - \bar{q}_6)^2] dt. \end{aligned} \tag{5.30}$$

From this inequality, (5.29) becomes

$$\gamma_2 \int_0^T (\mu_1^* q_6 - \bar{\mu}_1^* \bar{q}_6)(q_2 - \bar{q}_2) dt \leq C_4 \int_0^T [(q_2 - \bar{q}_2)^2 + (q_6 - \bar{q}_6)^2] dt.$$

The constants C_{01} , C_{03} , C_{04} , C_3 and C_4 which appear in the preceding inequalities depend on the coefficients and the bounds on state and adjoint variables. Consequently, we get

$$\begin{aligned}
 & \frac{1}{2}(q_2(0) - \bar{q}_2(0))^2 + \varpi \int_0^T (q_2 - \bar{q}_2)^2 dt \\
 & \leq k \int_0^T [(q_2 - \bar{q}_2)^2 + (q_3 - \bar{q}_3)^2] dt + \eta \int_0^T [(q_2 - \bar{q}_2)^2 + (q_4 - \bar{q}_4)^2] dt \\
 & + b \int_0^T [(q_2 - \bar{q}_2)^2 + (q_5 - \bar{q}_5)^2] dt + \epsilon \int_0^T [(q_2 - \bar{q}_2)^2 + (q_6 - \bar{q}_6)^2] dt \\
 & + C'_3 \int_0^T [(q_2 - \bar{q}_2)^2 + (q_3 - \bar{q}_3)^2 + (q_4 - \bar{q}_4)^2 + (q_5 - \bar{q}_5)^2 \\
 & + (q_6 - \bar{q}_6)^2 + (q_7 - \bar{q}_7)^2 + (q_8 - \bar{q}_8)^2 + (q_9 - \bar{q}_9)^2] dt,
 \end{aligned} \tag{5.31}$$

where the constant C'_3 depends on the coefficients and the upper bounds of state and adjoint variables.

Using the same reasoning for the remaining eight state and adjoint variables, we obtain their integral equations and their estimates. The combination of these eighteen estimates yields

$$\begin{aligned}
 & \frac{1}{2}(p_1 - \bar{p}_1)^2(T) + \frac{1}{2}(p_2 - \bar{p}_2)^2(T) + \frac{1}{2}(p_3 - \bar{p}_3)^2(T) + \frac{1}{2}(p_4 - \bar{p}_4)^2(T) \\
 & + \frac{1}{2}(p_5 - \bar{p}_5)^2(T) + \frac{1}{2}(p_6 - \bar{p}_6)^2(T) + \frac{1}{2}(p_7 - \bar{p}_7)^2(T) + \frac{1}{2}(p_8 - \bar{p}_8)^2(T) \\
 & + \frac{1}{2}(p_9 - \bar{p}_9)^2(T) + \frac{1}{2}(q_1 - \bar{q}_1)^2(0) + \frac{1}{2}(q_2 - \bar{q}_2)^2(0) + \frac{1}{2}(q_3 - \bar{q}_3)^2(0) \\
 & + \frac{1}{2}(q_4 - \bar{q}_4)^2(0) + \frac{1}{2}(q_5 - \bar{q}_5)^2(0) + \frac{1}{2}(q_6 - \bar{q}_6)^2(0) \\
 & + \frac{1}{2}(q_7 - \bar{q}_7)^2(0) + \frac{1}{2}(q_8 - \bar{q}_8)^2(0) + \frac{1}{2}(q_9 - \bar{q}_9)^2(0) \\
 & + \varpi \int_0^T [(p_1 - \bar{p}_1)^2 + (p_2 - \bar{p}_2)^2 + (p_3 - \bar{p}_3)^2 + (p_4 - \bar{p}_4)^2 + (p_5 - \bar{p}_5)^2 + (p_6 - \bar{p}_6)^2 \\
 & + (p_7 - \bar{p}_7)^2 + (p_8 - \bar{p}_8)^2 + (p_9 - \bar{p}_9)^2 + (q_1 - \bar{q}_1)^2 + (q_2 - \bar{q}_2)^2 + (q_3 - \bar{q}_3)^2 \\
 & + (q_4 - \bar{q}_4)^2 + (q_5 - \bar{q}_5)^2 + (q_6 - \bar{q}_6)^2 + (q_7 - \bar{q}_7)^2 + (q_8 - \bar{q}_8)^2 + (q_9 - \bar{q}_9)^2] dt \\
 & \leq \left(\bar{D}_1 + \bar{D}_2 e^{3\varpi T} \right) \int_0^T [(p_1 - \bar{p}_1)^2 + (p_2 - \bar{p}_2)^2 + (p_3 - \bar{p}_3)^2 \\
 & + (p_4 - \bar{p}_4)^2 + (p_5 - \bar{p}_5)^2 + (p_6 - \bar{p}_6)^2 + (p_7 - \bar{p}_7)^2 + (p_8 - \bar{p}_8)^2 + (p_9 - \bar{p}_9)^2 \\
 & + (q_1 - \bar{q}_1)^2 + (q_2 - \bar{q}_2)^2 + (q_3 - \bar{q}_3)^2 + (q_4 - \bar{q}_4)^2 + (q_5 - \bar{q}_5)^2 + (q_6 - \bar{q}_6)^2 \\
 & + (q_7 - \bar{q}_7)^2 + (q_8 - \bar{q}_8)^2 + (q_9 - \bar{q}_9)^2] dt.
 \end{aligned}$$

From this, we deduce that

$$\begin{aligned}
 & \left(\varpi - \tilde{D}_1 - \tilde{D}_2 e^{3\varpi T} \right) \int_0^T \left[(p_1 - \bar{p}_1)^2 + (p_2 - \bar{p}_2)^2 + (p_3 - \bar{p}_3)^2 \right. \\
 & + (p_4 - \bar{p}_4)^2 + (p_5 - \bar{p}_5)^2 + (p_6 - \bar{p}_6)^2 + (p_7 - \bar{p}_7)^2 + (p_8 - \bar{p}_8)^2 + (p_9 - \bar{p}_9)^2 \\
 & + (q_1 - \bar{q}_1)^2 + (q_2 - \bar{q}_2)^2 + (q_3 - \bar{q}_3)^2 + (q_4 - \bar{q}_4)^2 + (q_5 - \bar{q}_5)^2 + (q_6 - \bar{q}_6)^2 \\
 & \left. + (q_7 - \bar{q}_7)^2 + (q_8 - \bar{q}_8)^2 + (q_9 - \bar{q}_9)^2 \right] dt \leq 0,
 \end{aligned} \tag{5.32}$$

where the constants \tilde{D}_1 and \tilde{D}_2 depend on the coefficients and the upper bounds on state and adjoint variables.

By choosing $\varpi > \tilde{D}_1 + \tilde{D}_2$ and $T < \frac{1}{3\varpi} \ln \left(\frac{\varpi - \tilde{D}_1}{\tilde{D}_2} \right)$, it follows that $p_i = \bar{p}_i$ and $q_i = \bar{q}_i$, for $i = 1, \dots, 9$. Thus, the solution of the optimality system is unique for T sufficiently small. \square

Theorem 5.3. *implies that the unique optimal controls u_1^* , u_2^* and u_3^* are characterized in terms of the unique solution of the optimality system.*

6. Numerical simulations

In this section, we simulate the COVID-19 model (2.2) as a function of time. Recall that COVID-19 is eliminated from the population if $\mathcal{R}_c < 1$ and persists whenever $\mathcal{R}_c > 1$. The parameter values used here are given in Table 1. Most of the parameters were obtained from (Tang et al., 2020), ν is from (Liu et al., 2020), μ is from (WHO, 2020a, 2020b) and some are chosen arbitrarily to satisfy the stability property of the disease-free equilibrium as well as the endemic equilibrium of the COVID-19 model (2.2). Taking the parameter values from Table 1, except: $\beta = 3.62 \times 10^{-7}$, $\eta_1 = 0.9$, $a_0 = 10^{-12}$, $\mu = 1/57$, $d_1 = 0.156986$, we obtain $\mathcal{R}_c = 0.3073 < 1$. Here we consider the following four sets of initial conditions:

- Initial-1: $(S(0), E(0), A(0), I(0), U(0), Q(0), H(0), V(0), R(0)) = (12081, 8, 0, 0, 0, 80, 10, 1, 0)$,
- Initial-2: $(S(0), E(0), A(0), I(0), U(0), Q(0), H(0), V(0), R(0)) = (20081, 16, 0.5, 0.2, 0.3, 100, 20, 1.4, 10)$,
- Initial-3: $(S(0), E(0), A(0), I(0), U(0), Q(0), H(0), V(0), R(0)) = (50000, 26, 1.3, 0.4, 0.7, 200, 28, 2, 20)$,
- Initial-4: $(S(0), E(0), A(0), I(0), U(0), Q(0), H(0), V(0), R(0)) = (100000, 33, 2.3, 0.55, 1.1, 320, 40, 4, 40)$.

In these cases, the disease-free equilibrium \mathcal{E}_0 is globally asymptotically stable. Figs. 5 and 6 clearly confirm this fact and we also observe that the COVID-19 system initiating with Initial-1, Initial-2, Initial-3 and Initial-4 approaches the disease-free equilibrium $\mathcal{E}_0 = (2 \times 10^5, 0, 0, 0, 0, 0, 0, 0, 0)$. Thus, the numerical findings support Theorem 4.1. This illustrates the fact that COVID-19 could be eliminated from the Cameroonian population.

Again considering the parameter values from Table 1 and taking $\beta = 2.08 \times 10^{-6}$, $\eta_1 = 0.49$, $a_0 = 10^{-7}$, $\mu = 1/59$, $d_1 = 0.156986$, we obtain $\mathcal{R}_c = 1.6543 > 1$. Here we also consider four sets of initial conditions:

- Initial-5: $(S(0), E(0), A(0), I(0), U(0), Q(0), H(0), V(0), R(0)) = (120810, 8, 0, 0, 0, 80, 10, 1, 0)$,
- Initial-6: $(S(0), E(0), A(0), I(0), U(0), Q(0), H(0), V(0), R(0)) = (170810, 16, 0.5, 0.2, 0.3, 100, 20, 1.4, 10)$,
- Initial-7: $(S(0), E(0), A(0), I(0), U(0), Q(0), H(0), V(0), R(0)) = (280050, 26, 1.3, 0.4, 0.7, 200, 28, 2, 20)$,
- Initial-8: $(S(0), E(0), A(0), I(0), U(0), Q(0), H(0), V(0), R(0)) = (550000, 33, 2.3, 0.55, 1.1, 320, 40, 4, 40)$.

It follows that the unique endemic equilibrium point is globally asymptotically stable as can be observed numerically from Figs. 7 and 8, where the state variables initiating with Initial-5, Initial-6, Initial-7 and Initial-8 approach the endemic equilibrium $\mathcal{E}^* = (10^5, 0.3 \times 10^4, 2800, 800, 2500, 900, 0.6 \times 10^4, 99, 1.3 \times 10^5)$, which agrees with Theorem 4.7. This means epidemiologically that COVID-19 could persist in Cameroon.

Fig. 9 shows a good fit for total actual recovered individuals and those predicted by the model (2.2).

Figs. 10 and 11 illustrate the magnitude of quarantine and hospitalization. From these Figures, we clearly see that if the quarantine and hospitalization are operated efficiently, the disease will reduce considerably.

Figs. 12 and 13 illustrate Theorem 4.10.

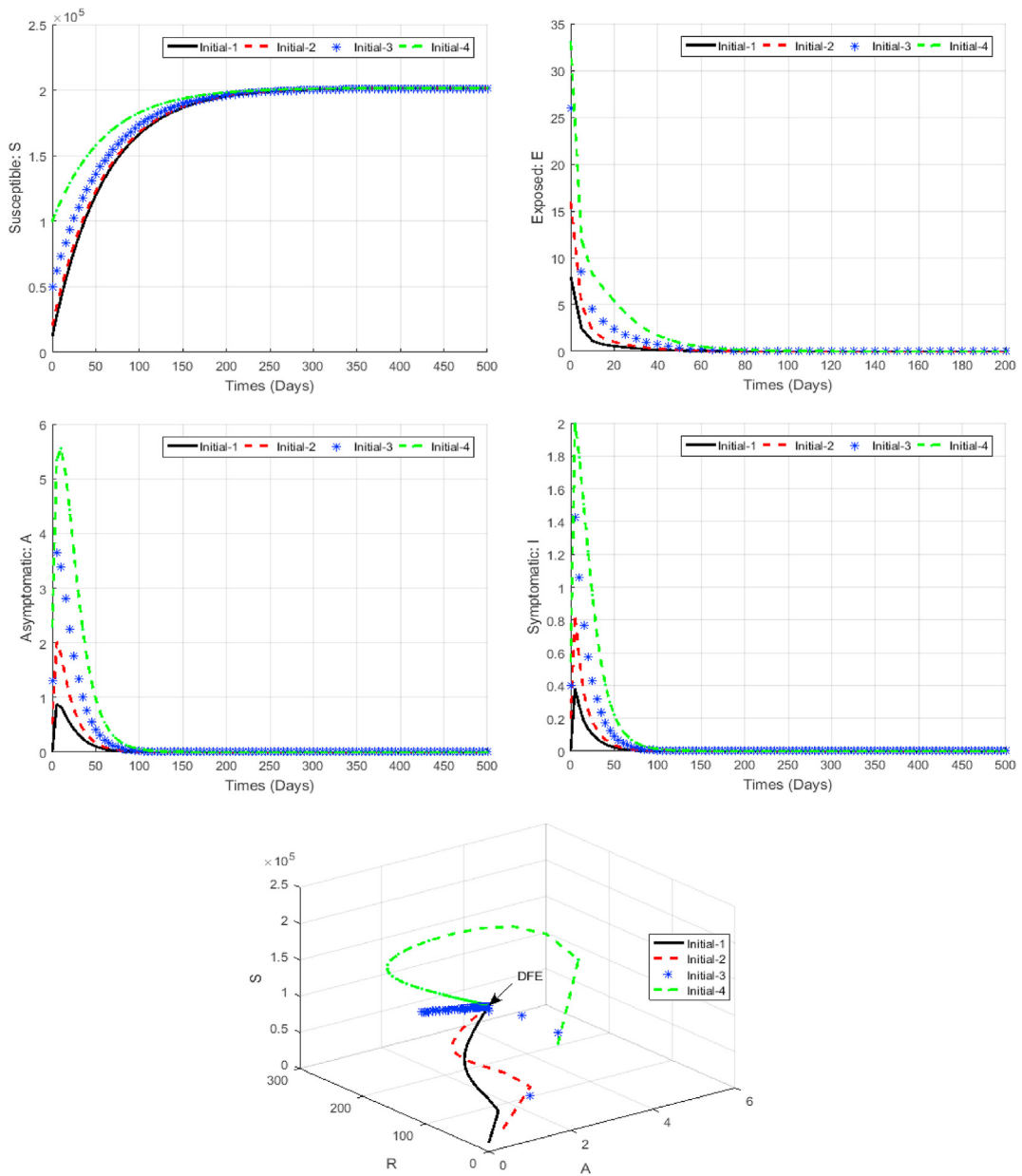


Fig. 5. Simulation of the COVID-19 model (2.2) as a function of time using various initial conditions and the parameter values from Table 1 except $\beta = 3.62 \times 10^{-7}$, $\eta_1 = 0.9$, $a_0 = 10^{-12}$, $\mu = 1/57$, $d_1 = 0.156986$, and $R_c = 0.3073 < 1$.

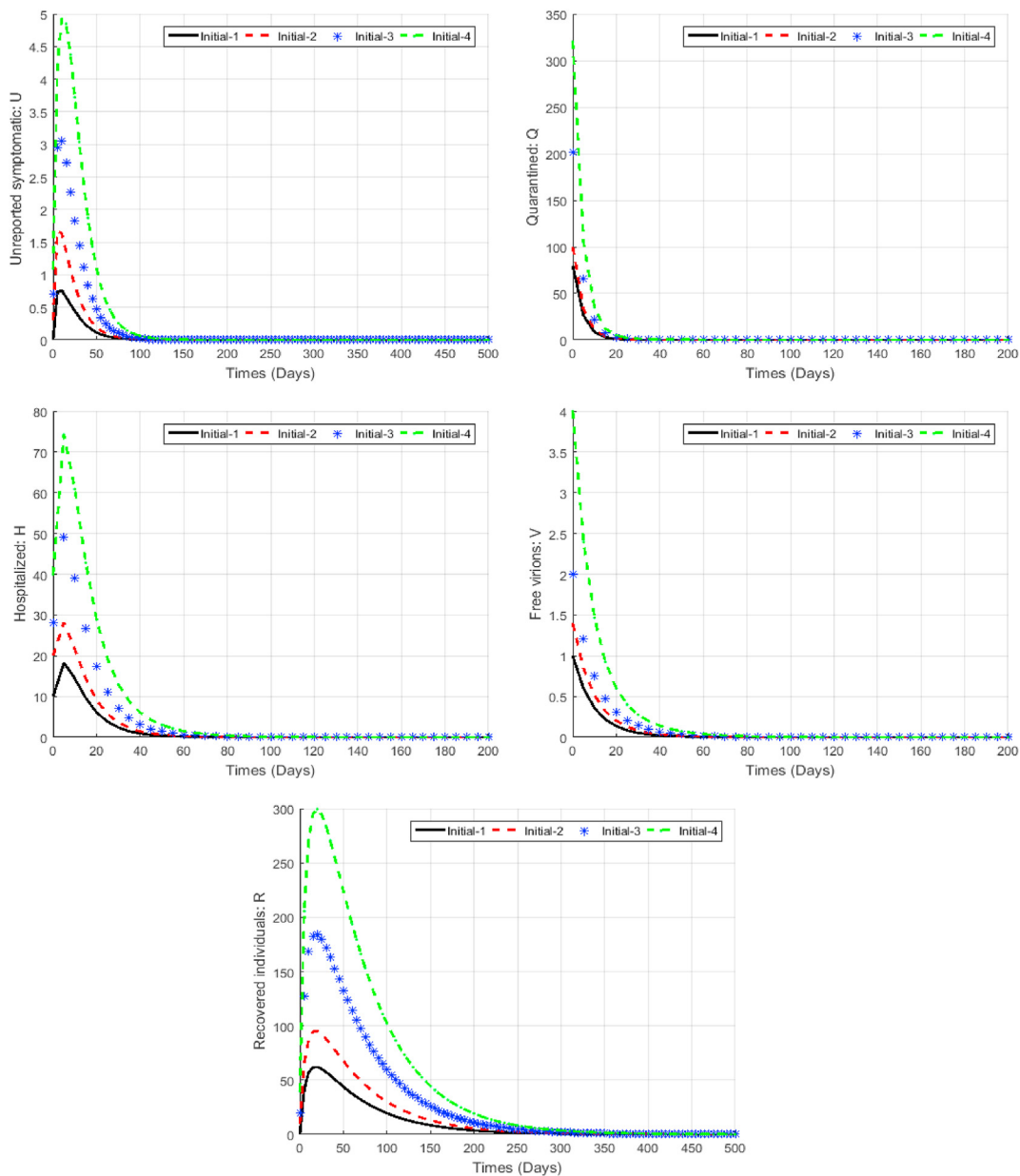


Fig. 6. Simulation of the COVID-19 model (2.2) as a function of time using various initial conditions and the parameter values from Table 1 except $\beta = 3.62 \times 10^{-7}$, $\eta_1 = 0.9$, $a_0 = 10^{-12}$, $\mu = 1/57$, $d_1 = 0.156986$, and $\mathcal{R}_c = 0.3073 < 1$.

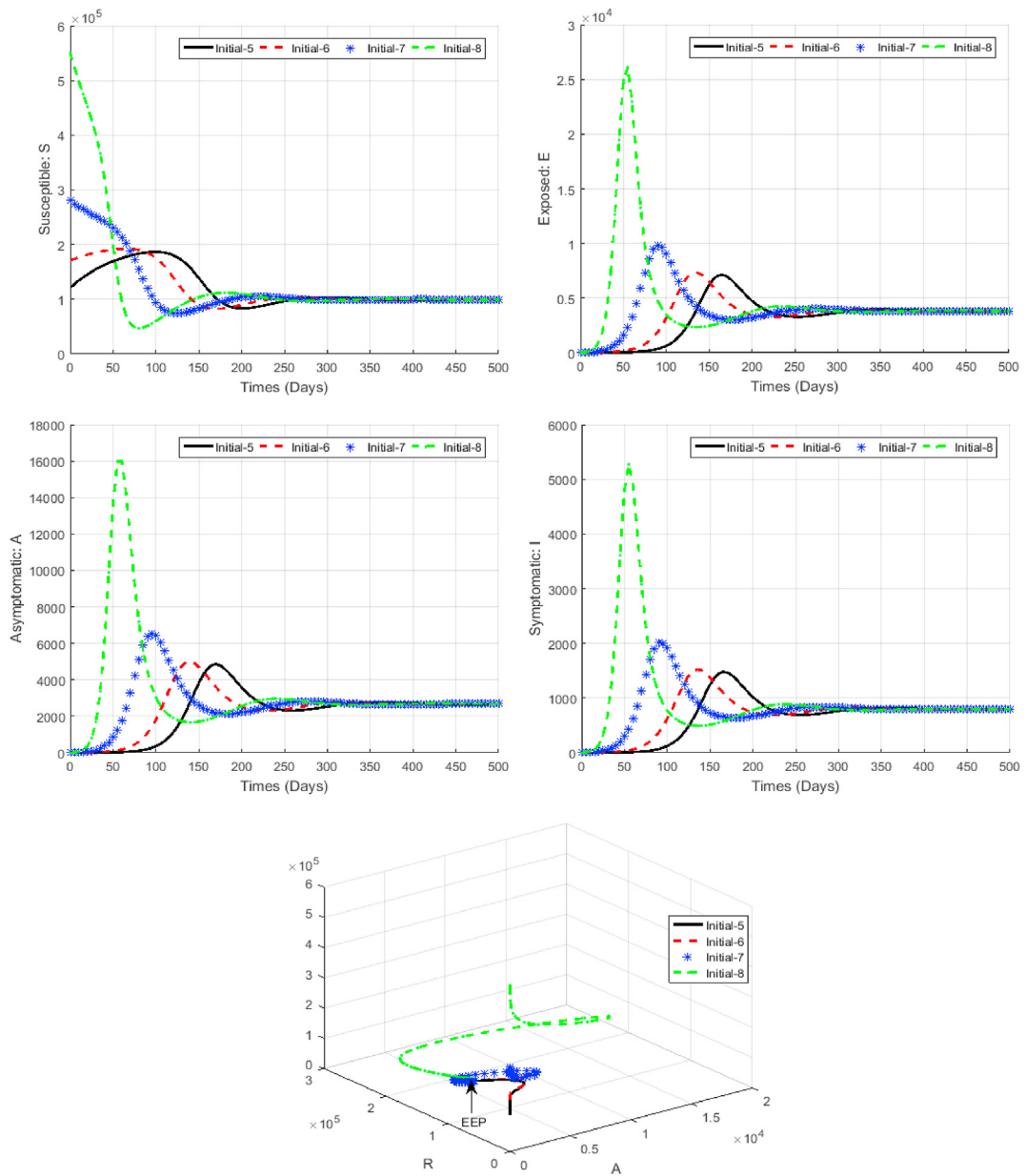


Fig. 7. Simulation of the COVID-19 model (2.2) as a function of time using various initial conditions and the parameter values from Table 1 except $\beta = 2.08 \times 10^{-6}$, $\eta_1 = 0.49$, $a_0 = 10^{-7}$, $\mu = 1/59$, $d_1 = 0.156986$, and $\mathcal{R}_c = 1.6543 > 1$.

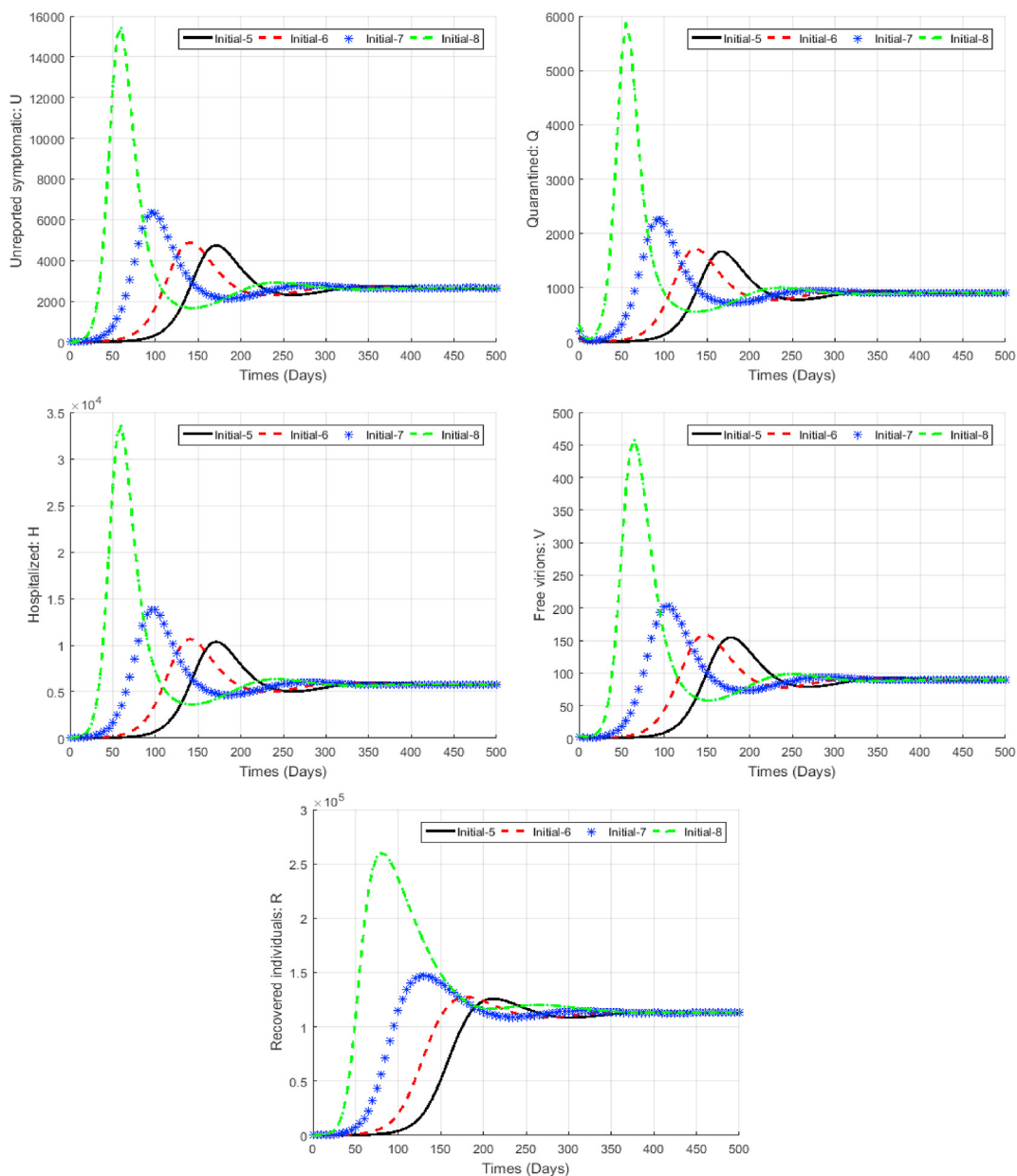


Fig. 8. Simulation of the COVID-19 model (2.2) as a function of time using various initial conditions and the parameter values from Table 1 except $\beta = 2.08 \times 10^{-6}$, $\eta_1 = 0.49$, $a_0 = 10^{-7}$, $\mu = 1/59$, $d_1 = 0.156986$, and $\mathcal{R}_c = 1.6543 > 1$.

We clearly observe from Fig. 12 that the cumulative number of new predicted active cases is higher when quarantine and hospitalization are not performed than when these control measures are implemented. This means that when condition $\eta_1 < \min\{\eta_{1\epsilon}, \eta_{1d_0}\}$ is satisfied, the use of quarantine and hospitalization could have positive impact on the community. But on Fig. 13, we see that the cumulative number of new predicted active cases is higher when quarantine and hospitalization are used than when these control measures are not implemented. This means that when condition $\eta_1 > \max\{\eta_{1\epsilon}, \eta_{1d_0}\}$ is satisfied, the use of quarantine and hospitalization could have negative impact to the community. The contour plots of Fig. 3 show the subordination of control reproduction number \mathcal{R}_c on the quarantine rate ϵ and the hospitalization rate d_0 for Cameroon.

Finally, the optimality system constituted of the established state equation (5.1), adjoint equation (5.3), control characterization (5.4)–(5.6) and corresponding initial and final conditions are carried out by using the forward-backward method. The algorithm starts by solving the state variables equations with a guess for the controls over the simulated time using an iterative method with forward fourth order Runge Kutta scheme. The state variables system with an initial guess is solved

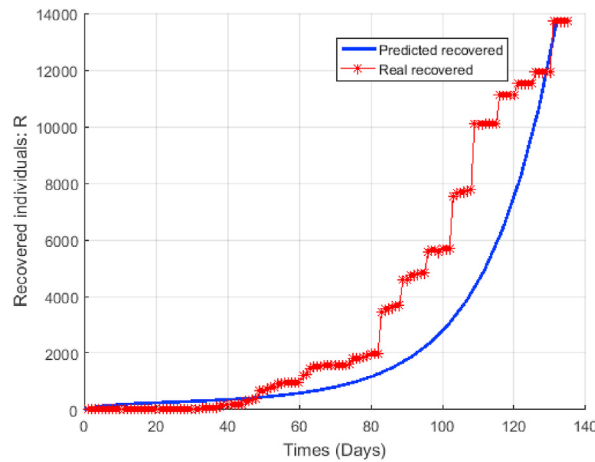


Fig. 9. Fitted results from the COVID-19 model (2.2) using the parameter values from Table 1 except $\beta = 2.08 \times 10^{-6}$, $\eta_1 = 0.49$, $a_0 = 10^{-7}$, $\mu = 1/59$, $d_1 = 0.156986$, and $\mathcal{R}_c = 1.6543 > 1$. Here, the red starred line indicates the real recovered cases and the blue line indicates the predicted recovered individuals.

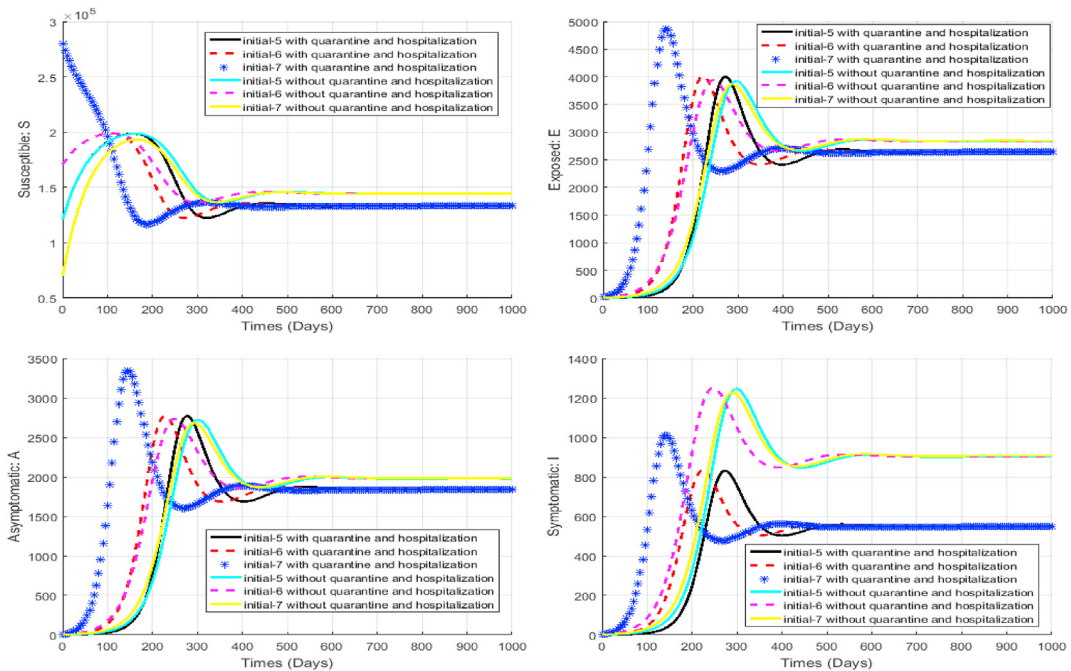


Fig. 10. Time plots for COVID-19 model (5.1) with quarantine and hospitalization (solid line) or without quarantine and hospitalization (dashed line) using various initial conditions. The parameter values are as given in Table 1, except $\beta = 1.55 \times 10^{-6}$, $\eta_1 = 0.49$, $a_0 = 10^{-7}$, $\mu = 1/59$, $d_1 = 0.156986$, and $\mathcal{R}_c = 1.2331 > 1$.

forward in time and then the adjoint system (5.3) is solved backward in time by a backward fourth order Runge Kutta scheme. This iterative process breaks off when the current state, adjoint, and control values converge sufficiently. Here, we choose the initial condition $(S(0), E(0), A(0), I(0), U(0), Q(0), H(0), V(0), R(0)) = (450000, 8, 0, 0, 0, 80, 10, 1, 0)$ to illustrate the control strategies. We choose the upper bound b_1 of u_1 equal to 0.8, owing to the reasonable case in Cameroon that it took at least average 3 days to quarantine people who have been exposed to COVID-19. We choose the upper bound b_2 of u_2 similarly to u_1 and the upper bound b_3 of u_3 equal to 0.7. Considering the weight coefficients associated with E, A, I, U, Q, H and V , we choose $B_1 = 100, B_2 = 500, B_3 = 2000, B_4 = 700, B_5 = 100, B_6 = 1500, B_7 = 800, R_1 = 3.5 \times 10^7, R_2 = 10^7$ and $R_3 = 2.5 \times 10^8$ to illustrate the optimal strategies. We suppose that the weight coefficient R_3 associated with control u_3 is greater than R_1 and R_2 which have close values associated with the controls u_1 and u_2 , respectively. These assumptions are based on the fact that: the cost associated with u_1 includes the cost of monitoring and quarantining schedule, and the cost associated with u_2 includes the

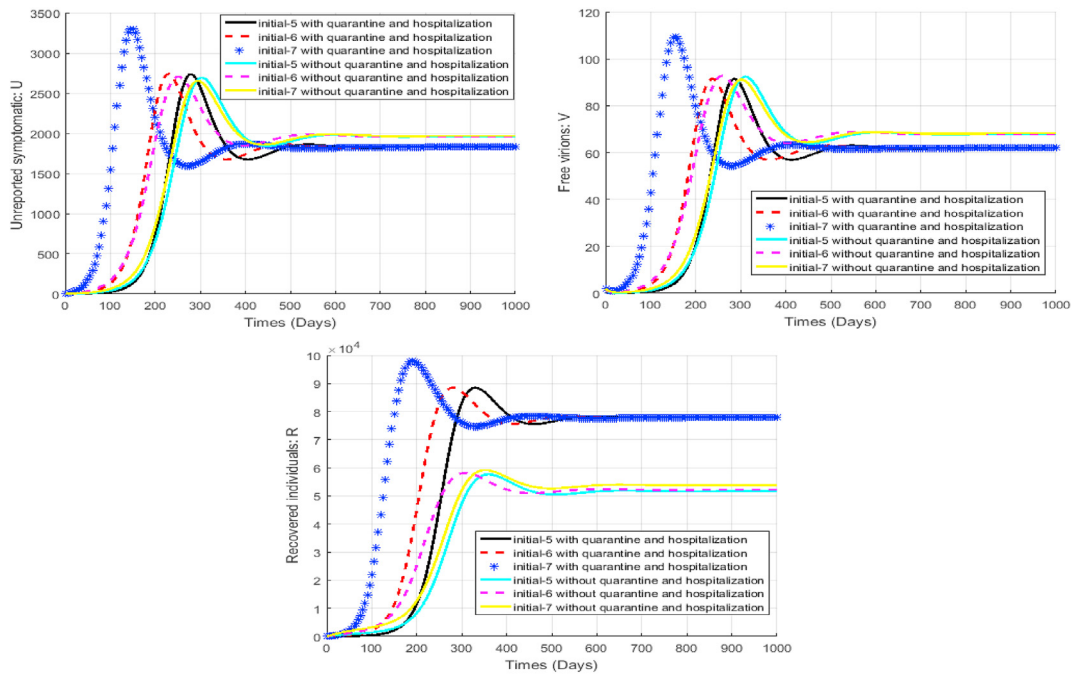


Fig. 11. Time plots for COVID-19 model (2.2) with quarantine and hospitalization (solid line) or without quarantine and hospitalization (dashed line) using various initial conditions. The parameter values are as given in Table 1, except $\beta = 1.55 \times 10^{-6}$, $\eta_1 = 0.49$, $a_0 = 10^{-7}$, $\mu = 1/59$, $d_1 = 0.156986$, and $\mathcal{R}_c = 1.2331 > 1$.

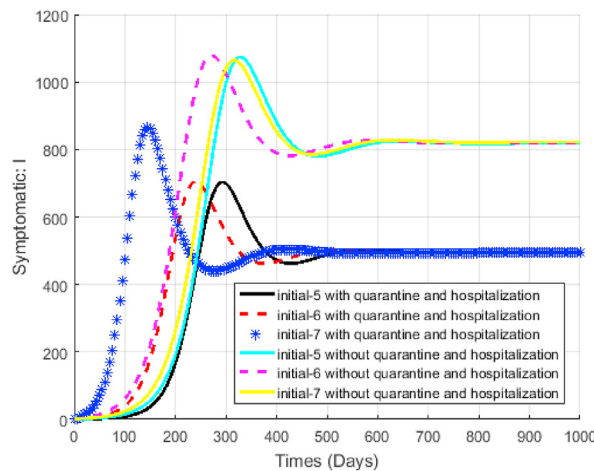


Fig. 12. Simulation of the COVID-19 model (2.2) giving the cumulative number of new cases of infection as a function of time and using various initial conditions. The parameter values are as given in Table 1, except $\beta = 1.55 \times 10^{-6}$, $a_0 = 10^{-7}$, $\mu = 1/57$, $d_1 = 0.999$, and $\eta_1 = 0.4$, so that $\mathcal{R}_c = 1.2241 > 1$, $\eta_{1e} = 0.9905$, $\eta_{1d_0} = 0.4145$ and $\eta_1 < \min\{\eta_{1e}, \eta_{1d_0}\}$.

cost of hospital special medical treatment resource, while the cost associated with u_3 includes the cost of hydro-alcoholic gel, disinfectant products and face masks. We observe on Figs. 14 and 15 that when the controls are used, the unreported symptomatic infectious individuals cases decrease faster than when the strategies are not applied. Moreover, in the presence of control measures, we have less infectious individuals than in the absence of the control. Also, the compliance with barrier measures such as the regular washing of hands, the use of hydro-alcoholic gel, wearing face masks, social distancing rules and disinfected surfaces can significantly reduce the number of infected and infectious individuals as well as the concentration of virus in the environment. Thus, the disease could infect a large part of the population if these measures are not followed.

Fig. 16 depicts the extremal control behaviour of u_1 , u_2 and u_3 . In order to minimize the total infected individuals, $E + A + I + U + Q + H$ and the concentration of virus in the environment, V , the optimal control u_1 stays at its upper bound for a

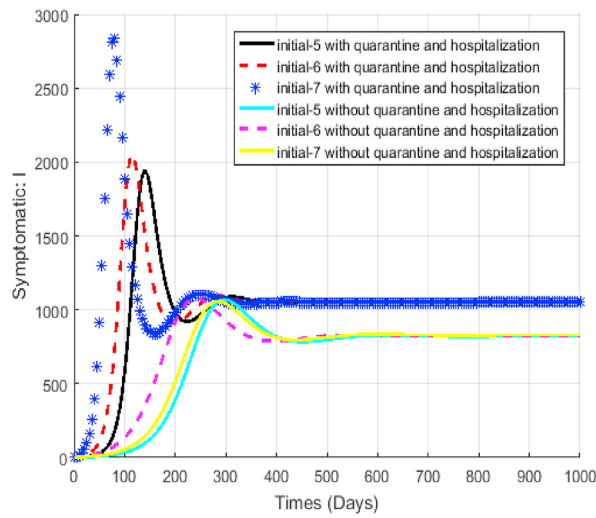


Fig. 13. Simulation of the COVID-19 model (2.2) giving the cumulative number of new cases of infection as a function of time and using various initial conditions. The parameter values are as given in Table 1, except $\beta = 1.55 \times 10^{-6}$, $a_0 = 10^{-7}$, $\mu = 1/57$, $\epsilon = 0.01$, $d_1 = 0.999$, and $\eta_1 = 0.9906$, so that $\mathcal{R}_c = 1.5112 > 1$, $\eta_{1\epsilon} = 0.9905$, $\eta_{1d_0} = 0.4145$ and $\eta_1 > \max\{\eta_{1\epsilon}, \eta_{1d_0}\}$.

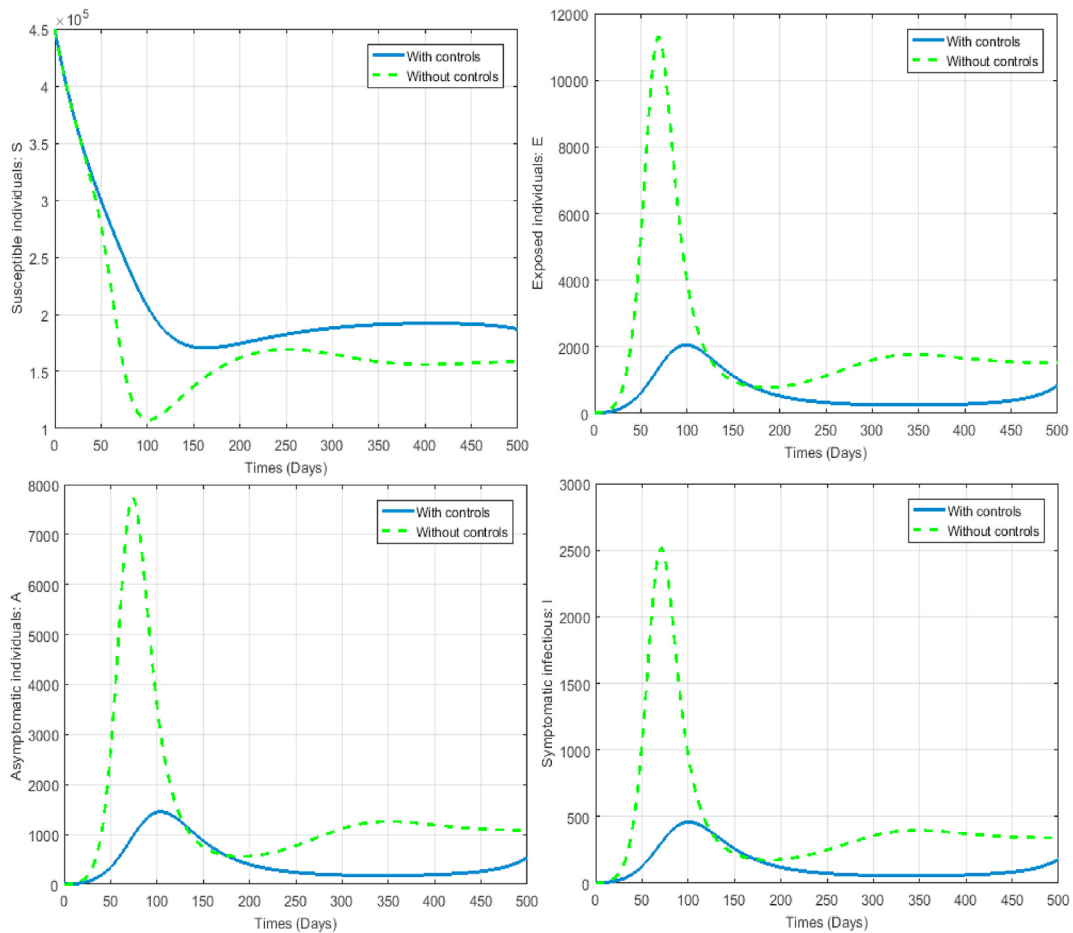


Fig. 14. Time plots for COVID-19 model (5.1) with control (solid line) or without control (dashed line). The parameter values are as given in Table 1, except $\gamma_1 = 0.7$, $\gamma_2 = 0.5$, $\beta = 1.55 \times 10^{-6}$, $\eta_1 = 0.49$, $a_0 = 10^{-7}$, $\mu = 1/57$.

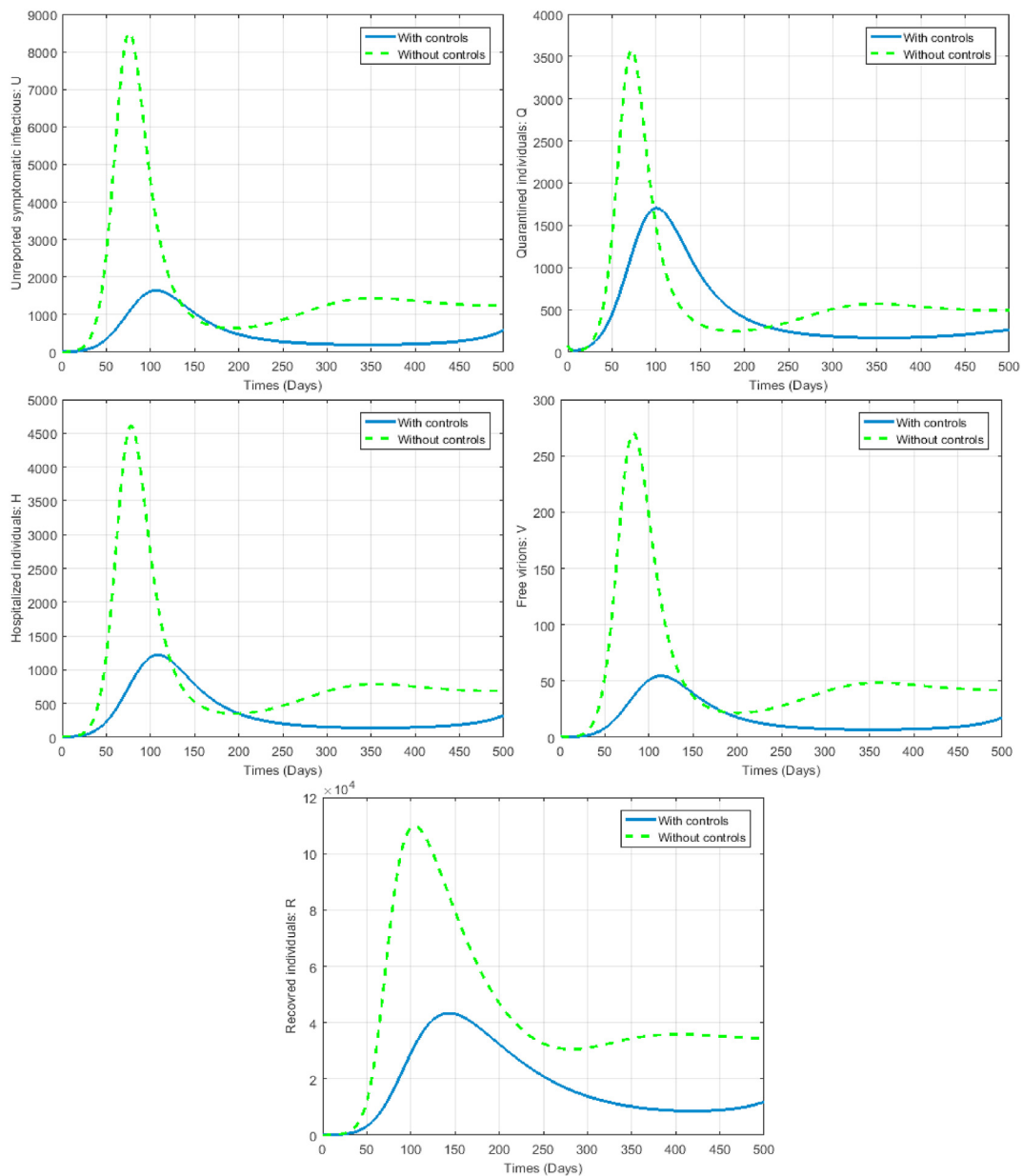


Fig. 15. Time plots for COVID-19 model (5.1) with control (solid line) or without control (dashed line). The parameter values are as given in Table 1, except $\gamma_1 = 0.7$, $\gamma_2 = 0.5$, $\beta = 1.55 \times 10^{-6}$, $\eta_1 = 0.49$, $a_0 = 10^{-7}$, $\mu = 1/57$.

short time, approximately 20 days and then steadily decreases to the lower bound in the remaining simulated time. Meanwhile, the optimal control u_2 starts at a lower level value zero, steadily increases to its upper bound and stays for short time, about 10 days, then steadily decreases to the lower bound in the simulated time until 500 days and, at the end, increases again to the level value (0.003). In the meantime, the optimal control u_3 also starts at a lower level value zero, steadily increases to an upper level value (8.7×10^{-5}) and stays for a short time, nearly up to 25 days, then is tapered off to a lower level (2.5×10^{-5}), and increases to its upper bound where it stays during two months and finally decreases steadily to the lower bound over the remaining simulated time.

Note that at the beginning of simulated time, the optimal control u_1 is staying at its upper bound in order to quarantine many exposed individuals (E) to prevent the increasing of the number of the infected classes. But at the beginning of simulated time, the optimal control u_2 seems to start by tracing, testing and then reaches its upper bound where it stays in order to hospitalize many symptomatic infectious individuals (I) to prevent the increasing of the number of people with

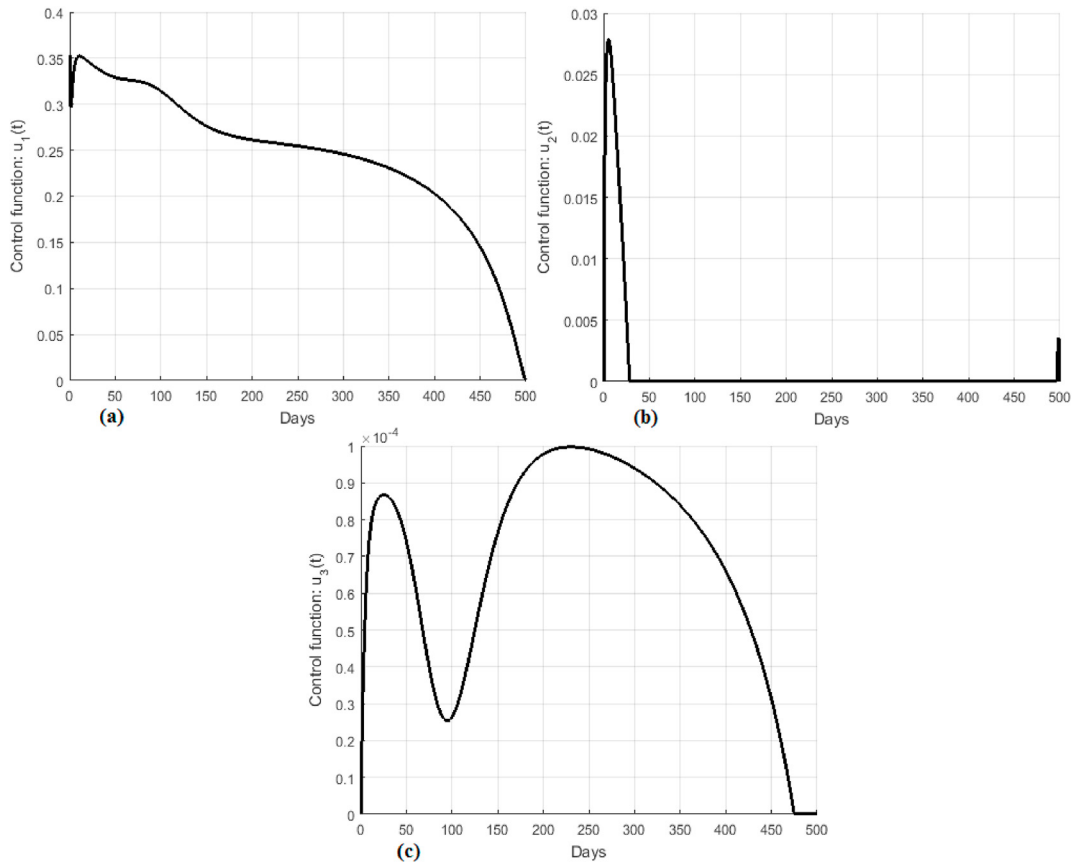


Fig. 16. The optimal control profiles (a) $u_1(t)$, (b) $u_2(t)$ and (c) $u_3(t)$ with $\gamma_1 = 0.7$, $\gamma_2 = 0.5$, $B_1 = 100$, $B_2 = 500$, $B_3 = 2000$, $B_4 = 700$, $B_5 = 100$, $B_6 = 1500$, $B_7 = 800$, $R_1 = 3.5 \times 10^7$, $R_2 = 10^7$ and $R_3 = 2.5 \times 10^8$.

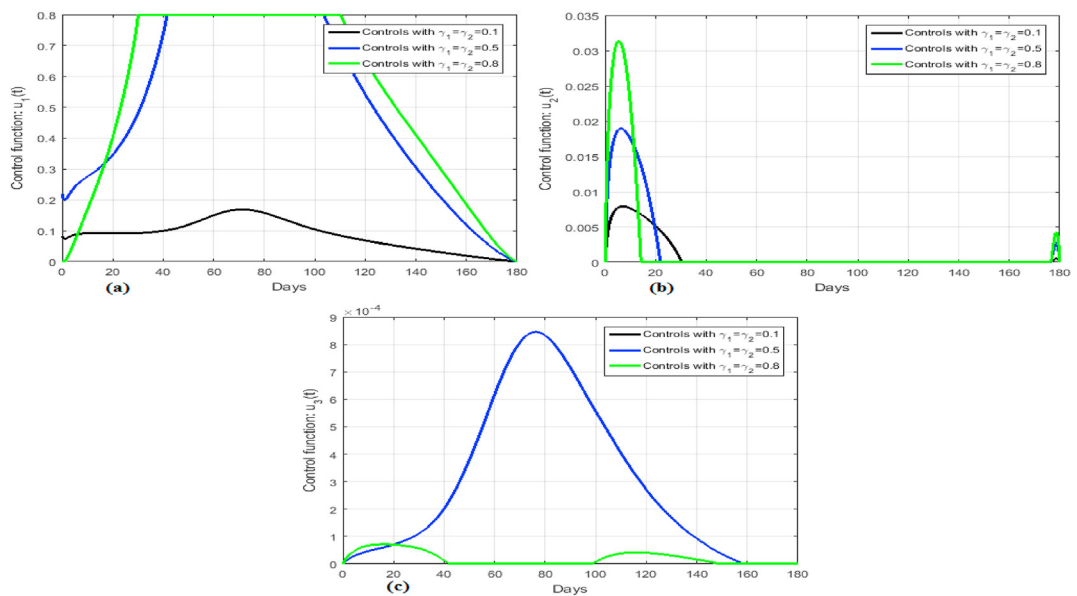


Fig. 17. The optimal control profiles (a) $u_1(t)$, (b) $u_2(t)$ and (c) $u_3(t)$ with $B_1 = 100$, $B_2 = 500$, $B_3 = 2000$, $B_4 = 700$, $B_5 = 100$, $B_6 = 1500$, $B_7 = 800$, $R_1 = 3.5 \times 10^7$, $R_2 = 10^7$ and $R_3 = 2.5 \times 10^8$.

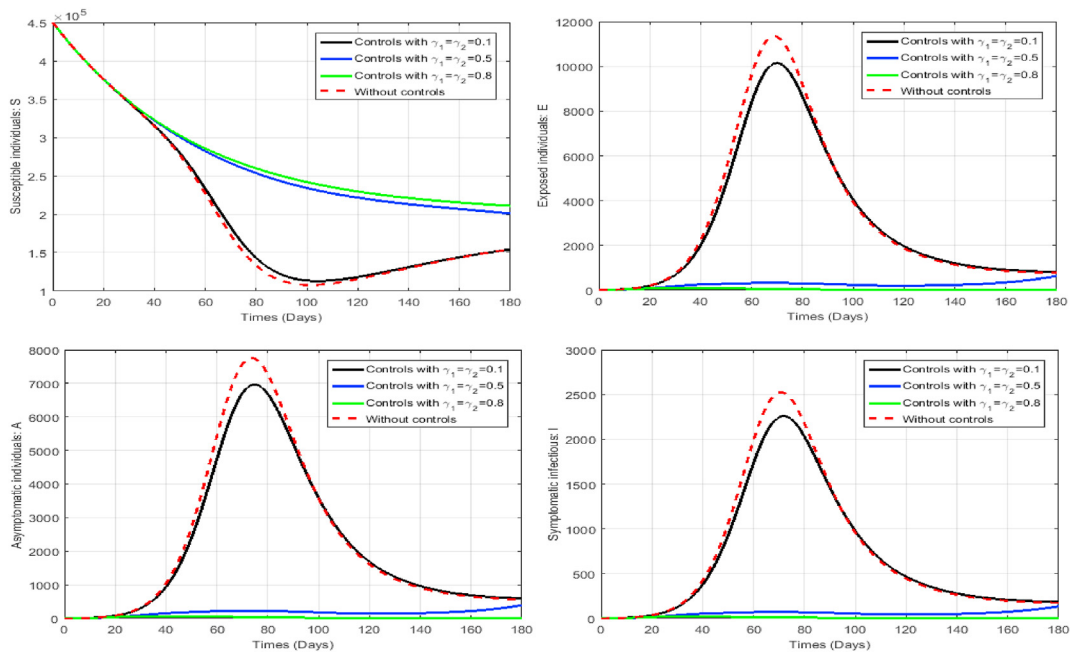


Fig. 18. Time plots for COVID-19 model (5.1) with control (solid lines) or without control (dashed lines). The parameter values are as given in Table 1, except $\beta = 1.55 \times 10^{-6}$, $\eta_1 = 0.49$, $a_0 = 10^{-7}$, $\mu = 1/57$.

clinical symptoms. Now, we see on Fig. 16 (c) that the optimal control u_3 implements the global effort of educational campaigns that run for over 250 days in order to prevent the increasing of the concentration of virus in the environment.

Fig. 17 shows the time dependent optimal control u_1 , u_2 and u_3 , for different values of special medical treatment rate γ_1 and mandatory quarantine rate γ_2 . From this figure, we can see that the higher the values of γ_1 and γ_2 , the more effective the controls u_1 and u_2 are, while the control u_3 is effective only at 50% of these values.

Figs. 18 and 19 illustrate how optimal control strategies change as the special medical treatment rate γ_1 and mandatory quarantine rate γ_2 vary. These Figures confirm that from 50% of the value of γ_1 and γ_2 , one could expect a considerable reduction of the infection in the community.

Fig. 20 represents the evolution number of positive cases in Cameroon from March 6 to July 20, 2020.

7. Conclusion

In this paper, to understand the transmission dynamics of COVID-19 in Cameroon, we formulated a compartmental ordinary differential equations model. A particular stress has been placed on quarantine and hospitalized classes. More precisely, we studied the impact of quarantine and hospitalization on curtailing the spread of the disease. The model is completely analyzed and the strategies for effective control of the progress of the disease are suggested. Using the method developed by van den Driessche and Watmough (van den Driessche and Watmough, 2002), we obtained the control reproduction number \mathcal{R}_c of the model. We constructed a suitable Lyapunov function to prove that system (2.2) has a globally asymptotically stable disease-free equilibrium whenever the control reproduction number is less than unity. When the control reproduction number exceeds unity, the disease-free equilibrium loss its stability and gives rise to a unique endemic equilibrium. By a skillful construction of a suitable Lyapunov function we proved that the endemic equilibrium is globally asymptotically stable. The efficiency of the quarantine of exposed cases and the isolation of hospitalized cases is dependent on the size of the modification parameter for the reduction of infectiousness of hospitalized individuals η_1 . It is shown that the use of quarantine and hospitalization could have positive impact on the population if $\eta_1 < \min\{\eta_{1e}, \eta_{1d_0}\}$, no impact if $\eta_1 = \min\{\eta_{1e}, \eta_{1d_0}\}$, and harmful impact if $\eta_1 > \max\{\eta_{1e}, \eta_{1d_0}\}$. Adding to this investigation the optimal control problem, we suggest quarantine and hospitalization as good strategies for controlling the disease. Note that COVID-19 is still ongoing in Cameroon and in many other countries in the world. This investigation attempt to provide Cameroonian authorities with some in-depth understanding of the disease dynamics so as to help them take better decisions for fighting against this highly deadly pandemic.

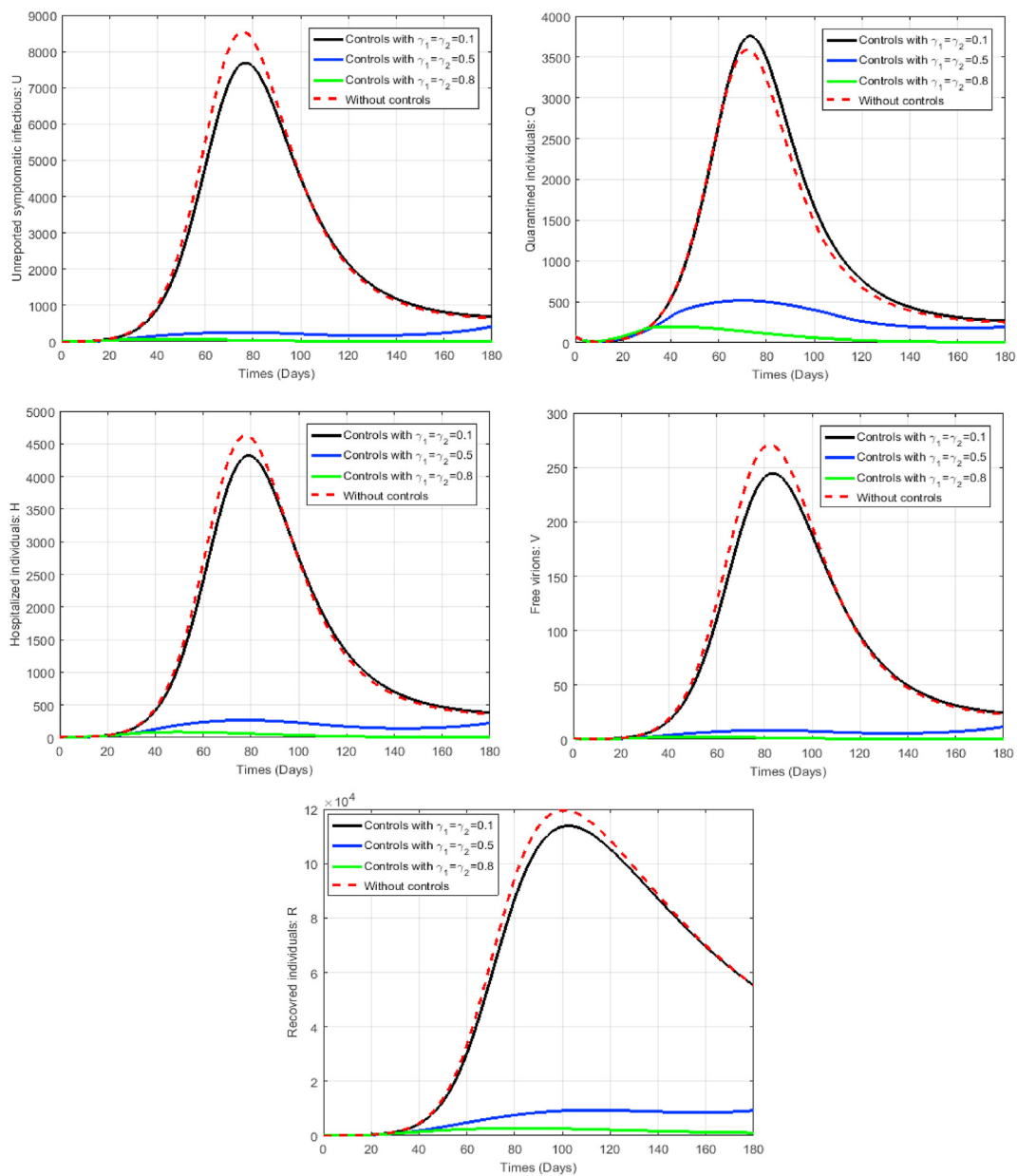


Fig. 19. Time plots for COVID-19 model (5.1) with control (solid lines) or without control (dashed lines). The parameter values are as given in Table 1, except $\beta = 1.55 \times 10^{-6}$, $\eta_1 = 0.49$, $a_0 = 10^{-7}$, $\mu = 1/57$.

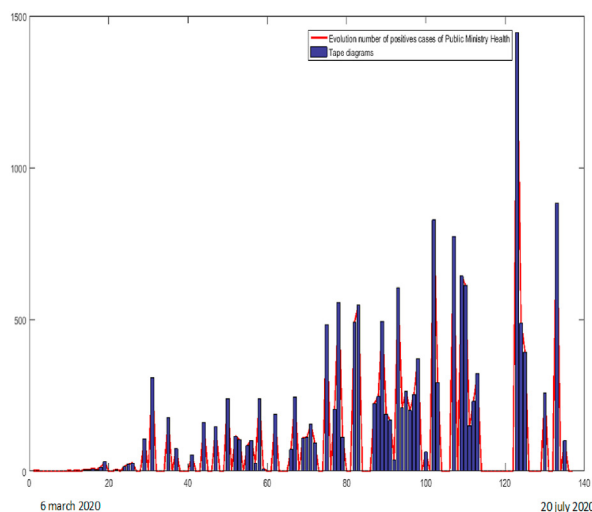


Fig. 20. Evolution number of positive cases. Source: Cameroon Ministry of Public Health.

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Data availability statement

The codes written to run most of the simulations presented in this work can be available upon simple request to the authors.

Declaration of interests

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.

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