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# A new fractional SIRS-SI malaria disease model with application of vaccines, antimalarial drugs, and spraying

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## Abstract

The present paper deals with a new fractional SIRS-SI model describing the transmission of malaria disease. The SIRS-SI malaria model is modified by using the Caputo–Fabrizio fractional operator for the inclusion of memory. We also suggest the utilization of vaccines, antimalarial medicines, and spraying for the treatment and control of the malaria disease. The theory of fixed point is utilized to examine the existence of the solution of a fractional SIRS-SI model describing spreading of malaria. The uniqueness of the solution of SIRS-SI model for malaria is also analyzed. It is shown that the treatments have great impact on the dynamical system of human and mosquito populations. The numerical simulation of fractional SIRS-SI malaria model is performed with the aid of HATM and Maple packages to show the effect of different parameters of the treatment of malaria disease. The numerical results for fractional SIRS-SI malaria model reveal that the recommended approach is very accurate and effective.

**Keywords:** Fractional SIRS-SI malaria model; Fixed point theorem; Caputo–Fabrizio fractional operator; HATM

## 1 Introduction

Malaria is a life-threatening mosquito-borne blood illness in the developing portion of the globe and especially in Asia and Africa. It is caused by a plasmodium parasite. The reports by the WHO show that malaria is a major risk to human life and remains a high-risk infectious illness. The financial load on the infected areas by malaria disease is massive and requires serious public health attention. Mathematical modeling of infectious diseases is a very strong tool to understand the dynamical system of disease spreading and control strategies. Recently many scientists and mathematicians have suggested mathematical models for malaria transmission. Augusto et al. [1] reported a SIR having infection rate of nonlinear type with the aid of vaccination for the human population. Abdulahi et al. [2] analyzed the outcomes of treatment as a control variable on the malaria transmission process. Mandal et al. [3] formulated the mathematical representation of malaria transmission with the situation of person-to-person transmission via blood transfusions and malaria-infected ladies having pregnancy. Chiyaka et al. [4] suggested a modified mathematical modeling by assuming that the persons also belonging to the recov-

ered group have a probability to be susceptible. Rafikov et al. [5] reported an efficient control strategy of malaria vector with the aid of genetically altered mosquitoes. Yang [6] suggested another approach for describing the malaria transmission connected with global warming and local socioeconomic circumstances. In a recent work, Senthamarai et al. [7] utilized the HAM to examine the spreading of malaria illness in an SIRS-SI model. However, all these approaches and mathematical models have their own limitation due to local nature of integer-order derivatives. Therefore fractional derivative approaches are suggested in mathematical modeling of biological and physical systems [8–21].

Very recently, Caputo and Fabrizio [22] reported a novel operator namely the Caputo–Fabrizio (CF) fractional operator involving a nonsingular kernel. Furthermore, the additional properties of this operator were put up by Losada and Nieto [23]. The suitability and efficiency of the CF fractional operator have been demonstrated by many researchers. Singh et al. [24] reported a mathematical model for computer viruses incorporating the CF fractional operator. Singh et al. [25] suggested an innovative idea for mathematical modeling of giving up smoking dynamics with the aid of the CF fractional operator. Djida and Atangana [26] studied a water flow within a confined aquifer connected to an arbitrary order operator in the terms of the Caputo–Fabrizio and many others. Inspired by ongoing investigations on the CF fractional derivative and their effectiveness, we employ this derivative in SIRS-SI malaria model for inclusion of memory. The existence and uniqueness of the solution of SIRS-SI model describing spreading of malaria involving memory effects is shown by applying the fixed-point theory. The numerical computation for the fractional SIRS-SI malaria model is executed by using HATM [27–31] and Padé approximation [32]. The present paper is developed as follows. In Sect. 2, we present the required results pertaining to the CF operator of arbitrary order. In Sect. 3, we give a mathematical formulation of fractional SIRS-SI malaria model. Section 4 concerns with the existence and uniqueness analysis of solution of SIRS-SI model representing the malaria having fractional order. In Sect. 5, we use the efficiency of HATM for examining fractional SIRS-SI malaria model. In Sect. 6, we investigate the effect of various parameters on humans and mosquitoes. Finally, in Sect. 7, we conclude.

## 2 Preliminaries

**Definition 1** Let  $\psi \in H^1(\ell_1, \ell_2)$ ,  $\ell_2 > \ell_1$ ,  $\lambda \in (0, 1]$ . Then the CF fractional operator [22] is expressed as

$$\begin{aligned} D_t^\lambda(\psi(t)) &= \frac{M(\lambda)}{1-\lambda} \int_{\ell_1}^t \psi'(\theta) \exp\left[-\lambda \frac{t-\theta}{1-\lambda}\right] d\theta, \quad 0 < \lambda < 1, \\ &= \frac{d\psi}{dt}, \quad \lambda = 1. \end{aligned} \quad (1)$$

In this expression  $M(\lambda)$  satisfies the condition  $M(0) = M(1) = 1$  [22].

**Definition 2** The integral operator of fractional order corresponding to the CF fractional derivative is defined as [23]

$$I_t^\lambda(\psi(t)) = \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} \psi(t) + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t \psi(\zeta) d\zeta, \quad t \geq 0. \quad (2)$$

**Definition 3** The Laplace transform of  ${}_0^{CF}D_t^\lambda \psi(t)$  is represented as

$$L[{}_0^{CF}D_t^\lambda \psi(t)] = M(\lambda) \frac{sL[\psi(t)] - \psi(0)}{s + \lambda(1 - s)}. \tag{3}$$

### 3 Fractional SIRS-SI malaria model with exponential law

We suppose that the population of human is categorized into the following three groups:

- Group I: Susceptible human denoted by  $S_h$ ;
- Group II: Infected human denoted by  $I_h$ ;
- Group III: Recovered human denoted by  $R_h$ .

The population of mosquito is categorized into the following two groups:

- Group I: Susceptible mosquito denoted by  $S_m$ ;
- Group II: Infected mosquito denoted by  $I_m$ .

Let us suppose that the person who is born is transferred to the susceptible group with fixed rate  $\alpha_h$  per unit time. Peoples in the susceptible group are transferred to the infected group because of blood exchanging with rate  $\omega\gamma_1$  per unit time (here  $\omega$  indicates the average number of blood transfusions between the susceptible and infected groups in a fixed time period, whereas  $\gamma_1$  stands for the probability of disease transfer from an infected person to a susceptible person) or through an infected mosquito bite with rate  $\xi\gamma_2$  per unit time (here  $\xi$  stands for the average number of infected mosquito bites on a susceptible person in a fixed time period, whereas  $\gamma_2$  stands for the probability of disease transfer to susceptible persons through infected mosquitoes). Persons in the susceptible group transfer into the recovered group because of vaccination with rate  $\delta$  per unit time. The persons in the susceptible group expire with rate  $\eta_h$ . A newly born baby is infected by malaria because of inbred at rate  $\mu$  per unit time. The persons in infected group can transfer to the recovered group because of using antimalarial drugs at rate  $c\nu$  per unit time (here  $c$  stands for the rate of people healing, and  $\nu$  indicates the potency of anti-malarial medicines). Peoples in the infected group can expire with rate  $\eta_h$  and die because of malaria disease with rate  $\varepsilon$  per unit time. Peoples in recovered group die with rate  $\eta_h$  per unit time. Moreover, mosquitoes are born and transferred to the susceptible group at a fixed rate  $\alpha_m$  per unit time. The mosquitoes in the susceptible group can transfer into the infected group by biting of mosquito to the infected persons with rate  $e\gamma_3$  per unit time (here  $e$  stand for the average number of susceptible mosquito bites to the infected persons in a fixed time period, and  $\gamma_3$  indicates the possibility of disease passing to susceptible mosquitoes from the infected peoples) or can expire with rate  $\eta_m$  per unit time. The average per capita rate of loss of immunity is  $\beta$  per unit time. The mosquitoes belonging to the susceptible and infected groups can expire due to exercise of spraying with rate  $\sigma$  per unit time. The mosquitoes belonging to the infected group can die with rate  $\eta_m$  per unit time. The governing equations for the SIRS-SI malaria model are presented

as follows:

$$\begin{aligned}
 \frac{dS_h}{dt} &= \alpha_h + \beta R_h - (\omega\gamma_1 I_h + \xi\gamma_2 I_m)S_h - (\delta + \eta_h)S_h, \\
 \frac{dI_h}{dt} &= \mu I_h + (\omega\gamma_1 I_h + \xi\gamma_2 I_m)S_h - (\eta_h + \varepsilon + cv)I_h, \\
 \frac{dR_h}{dt} &= cvI_h - (\eta_h + \beta)R_h + \delta S_h, \\
 \frac{dS_m}{dt} &= \alpha_m - (e\gamma_3 I_h + \eta_m + \sigma)S_m, \\
 \frac{dI_m}{dt} &= e\gamma_3 I_h S_m - (\eta_m + \sigma)I_m.
 \end{aligned}
 \tag{4}$$

Since the integer-order derivative is local in nature, the presented SIRS-SI malaria model (4) does not describe different effects on humans and mosquitoes in efficient manner. Therefore, to include the memory effects in the description of malaria disease, we extend the model (4) by employing the newly proposed Caputo–Fabrizio fractional derivative as follows:

$$\begin{aligned}
 {}^{CF}D_0^\lambda S_h &= \alpha_h + \beta R_h - (\omega\gamma_1 I_h + \xi\gamma_2 I_m)S_h - (\delta + \eta_h)S_h, \\
 {}^{CF}D_0^\lambda I_h &= \mu I_h + (\omega\gamma_1 I_h + \xi\gamma_2 I_m)S_h - (\eta_h + \varepsilon + cv)I_h, \\
 {}^{CF}D_0^\lambda R_h &= cvI_h - (\eta_h + \beta)R_h + \delta S_h, \\
 {}^{CF}D_0^\lambda S_m &= \alpha_m - (e\gamma_3 I_h + \eta_m + \sigma)S_m, \\
 {}^{CF}D_0^\lambda I_m &= e\gamma_3 I_h S_m - (\eta_m + \sigma)I_m,
 \end{aligned}
 \tag{5}$$

with the initial conditions

$$S_h(0) = c_1, \quad I_h(0) = c_2, \quad R_h(0) = c_3, \quad S_m(0) = c_4, \quad I_m(0) = c_5.
 \tag{6}$$

Let us suppose that  $B$  is the Banach space of continuous real-valued functions defined on an interval  $I$  with the associated norm

$$\|(S_h, I_h, R_h, S_m, I_m)\| = \|S_h\| + \|I_h\| + \|R_h\| + \|S_m\| + \|I_m\|.
 \tag{7}$$

In Eq. (7), we have  $\|S_h\| = \sup\{|S_h(t)| : t \in I\}$ ,  $\|I_h\| = \sup\{|I_h(t)| : t \in I\}$ ,  $\|R_h\| = \sup\{|R_h(t)| : t \in I\}$ ,  $\|S_m\| = \sup\{|S_m(t)| : t \in I\}$ , and  $\|I_m\| = \sup\{|I_m(t)| : t \in I\}$ . Specifically,  $B = E(I) \times E(I) \times E(I) \times E(I) \times E(I)$ , where  $E(I)$  stands for the Banach space of continuous real-valued functions on  $I$  and the associated sup norm.

#### 4 Existence and uniqueness analysis

This section deals with the existence and uniqueness analysis of the solution of the fractional SIRS-SI malaria model with exponential law. It is very important to know about the existence and uniqueness of the solution of any mathematical model in natural sciences. Therefore we examine the existence and uniqueness of the solution of fractional SIRS-SI malaria model by using fixed point theory [33–35].

We apply the fractional integral operator (2) to Eq. (5), which gives

$$\begin{aligned}
 S_h(t) - S_h(0) &= {}_0^{\text{CF}}I_t^\lambda \{ \alpha_h + \beta R_h - (\omega\gamma_1 I_h + \xi\gamma_2 I_m) S_h - (\delta + \eta_h) S_h \}, \\
 I_h(t) - I_h(0) &= {}_0^{\text{CF}}I_t^\lambda \{ \mu I_h + (\omega\gamma_1 I_h + \xi\gamma_2 I_m) S_h - (\eta_h + \varepsilon + cv) I_h \}, \\
 R_h(t) - R_h(0) &= {}_0^{\text{CF}}I_t^\lambda \{ cv I_h - (\eta_h + \beta) R_h + \delta S_h \}, \\
 S_m(t) - S_m(0) &= {}_0^{\text{CF}}I_t^\lambda \{ \alpha_m - (e\gamma_3 I_h + \eta_m + \sigma) S_m \}, \\
 I_m(t) - I_m(0) &= {}_0^{\text{CF}}I_t^\lambda \{ e\gamma_3 I_h S_m - (\eta_m + \sigma) I_m \}.
 \end{aligned}
 \tag{8}$$

Using the notation suggested in [23], we have

$$\begin{aligned}
 &S_h(t) - S_h(0) \\
 &= \frac{2(1 - \lambda)}{(2 - \lambda)M(\lambda)} \{ \alpha_h + \beta R_h(t) - (\omega\gamma_1 I_h(t) + \xi\gamma_2 I_m(t)) S_h(t) - (\delta + \eta_h) S_h(t) \} \\
 &\quad + \frac{2\lambda}{(2 - \lambda)M(\lambda)} \int_0^t \{ \alpha_h + \beta R_h(\varsigma) - (\omega\gamma_1 I_h(\varsigma) + \xi\gamma_2 I_m(\varsigma)) S_h(\varsigma) \\
 &\quad - (\delta + \eta_h) S_h(\varsigma) \} d\varsigma, \\
 &I_h(t) - I_h(0) \\
 &= \frac{2(1 - \lambda)}{(2 - \lambda)M(\lambda)} \{ \mu I_h(t) + (\omega\gamma_1 I_h(t) + \xi\gamma_2 I_m(t)) S_h(t) - (\eta_h + \varepsilon + cv) I_h(t) \} \\
 &\quad + \frac{2\lambda}{(2 - \lambda)M(\lambda)} \int_0^t \{ \mu I_h(\varsigma) + (\omega\gamma_1 I_h(\varsigma) + \xi\gamma_2 I_m(\varsigma)) S_h(\varsigma) \\
 &\quad - (\eta_h + \varepsilon + cv) I_h(\varsigma) \} d\varsigma, \\
 &R_h(t) - R_h(0) \\
 &= \frac{2(1 - \lambda)}{(2 - \lambda)M(\lambda)} \{ cv I_h(t) - (\eta_h + \beta) R_h(t) + \delta S_h(t) \} \\
 &\quad + \frac{2\lambda}{(2 - \lambda)M(\lambda)} \int_0^t \{ cv I_h(\varsigma) - (\eta_h + \beta) R_h(\varsigma) + \delta S_h(\varsigma) \} d\varsigma, \\
 &S_m(t) - S_m(0) \\
 &= \frac{2(1 - \lambda)}{(2 - \lambda)M(\lambda)} \{ \alpha_m - (e\gamma_3 I_h(t) + \eta_m + \sigma) S_m(t) \} \\
 &\quad + \frac{2\lambda}{(2 - \lambda)M(\lambda)} \int_0^t \{ \alpha_m - (e\gamma_3 I_h(\varsigma) + \eta_m + \sigma) S_m(\varsigma) \} d\varsigma, \\
 &I_m(t) - I_m(0) \\
 &= \frac{2(1 - \lambda)}{(2 - \lambda)M(\lambda)} \{ e\gamma_3 I_h(t) S_m(t) - (\eta_m + \sigma) I_m(t) \} \\
 &\quad + \frac{2\lambda}{(2 - \lambda)M(\lambda)} \int_0^t \{ e\gamma_3 I_h(\varsigma) S_m(\varsigma) - (\eta_m + \sigma) I_m(\varsigma) \} d\varsigma.
 \end{aligned}
 \tag{9}$$

For clarity, we express

$$\begin{aligned}
 \Omega_1(t, S_h) &= \alpha_h + \beta R_h(t) - (\omega\gamma_1 I_h(t) + \xi\gamma_2 I_m(t))S_h(t) - (\delta + \eta_h)S_h(t), \\
 \Omega_2(t, I_h) &= \mu I_h(t) + (\omega\gamma_1 I_h(t) + \xi\gamma_2 I_m(t))S_h(t) - (\eta_h + \varepsilon + c\nu)I_h(t), \\
 \Omega_3(t, R_h) &= c\nu I_h(t) - (\eta_h + \beta)R_h(t) + \delta S_h(t), \\
 \Omega_4(t, S_m) &= \alpha_m - (e\gamma_3 I_h(t) + \eta_m + \sigma)S_m(t), \\
 \Omega_5(t, I_m) &= e\gamma_3 I_h(t)S_m(t) - (\eta_m + \sigma)I_m(t).
 \end{aligned}
 \tag{10}$$

**Theorem 1** *The kernels  $\Omega_1, \Omega_2, \Omega_3, \Omega_4,$  and  $\Omega_5$  satisfy the Lipschitz condition and contraction if*

$$0 \leq \omega\gamma_1 a_2 + \xi\gamma_2 a_5 + (\delta + \eta_h) < 1.$$

*Proof* We initiate with  $\Omega_1$ . For two functions  $S_h$  and  $S_{h1}$ , we have

$$\begin{aligned}
 &\|\Omega_1(t, S_h) - \Omega_1(t, S_{h1})\| \\
 &= \|\{S_h(t) - S_{h1}(t)\}(\omega\gamma_1 I_h(t) + \xi\gamma_2 I_m(t)) - \{S_h(t) - S_{h1}(t)\}(\delta + \eta_h)\|.
 \end{aligned}
 \tag{11}$$

On applying the properties of norm on Eq. (11), it yields

$$\begin{aligned}
 \|\Omega_1(t, S_h) - \Omega_1(t, S_{h1})\| &\leq \|\{S_h(t) - S_{h1}(t)\}(\omega\gamma_1 I_h(t) + \xi\gamma_2 I_m(t))\| \\
 &\quad + \|\{S_h(t) - S_{h1}(t)\}(\delta + \eta_h)\| \\
 &\leq \{\omega\gamma_1 \|I_h(t)\| + \xi\gamma_2 \|I_m(t)\| + (\delta + \eta_h)\} \|S_h(t) - S_{h1}(t)\| \\
 &\leq \{\omega\gamma_1 a_2 + \xi\gamma_2 a_5 + (\delta + \eta_h)\} \|S_h(t) - S_{h1}(t)\| \\
 &\leq b_1 \|S_h(t) - S_{h1}(t)\|.
 \end{aligned}
 \tag{12}$$

Taking  $b_1 = \omega\gamma_1 a_2 + \xi\gamma_2 a_5 + (\delta + \eta_h)$ , where  $\|S_h(t)\| \leq a_1, \|I_h(t)\| \leq a_2, \|R_h(t)\| \leq a_3, \|S_m(t)\| \leq a_4,$  and  $\|I_m(t)\| \leq a_5$  are bounded functions, we get

$$\|\Omega_1(t, S_h) - \Omega_1(t, S_{h1})\| \leq b_1 \|S_h(t) - S_{h1}(t)\|.$$

Thus the Lipschitz condition is satisfied for  $\Omega_1$ . Furthermore, if

$$0 \leq \omega\gamma_1 a_2 + \xi\gamma_2 a_5 + (\delta + \eta_h) < 1, \quad \text{then it is also a contraction.}$$

Similarly, we can prove that the kernels  $\Omega_2(t, I_h), \Omega_3(t, R_h), \Omega_4(t, S_m),$  and  $\Omega_5(t, I_m)$  satisfy the Lipschitz conditions

$$\begin{aligned}
 \|\Omega_2(t, I_h) - \Omega_2(t, I_{h1})\| &\leq b_2 \|I_h(t) - I_{h1}(t)\|, \\
 \|\Omega_3(t, R_h) - \Omega_3(t, R_{h1})\| &\leq b_3 \|R_h(t) - R_{h1}(t)\|, \\
 \|\Omega_4(t, S_m) - \Omega_4(t, S_{m1})\| &\leq b_4 \|S_m(t) - S_{m1}(t)\|, \\
 \|\Omega_5(t, I_m) - \Omega_5(t, I_{m1})\| &\leq b_5 \|I_m(t) - I_{m1}(t)\|.
 \end{aligned}
 \tag{14}$$

On using the notations of the earlier stated kernels, Eq. (9) reduces to the system

$$\begin{aligned}
 S_h(t) &= S_h(0) + \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)}\Omega_1(t, S_h) + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t \Omega_1(\zeta, S_h) d\zeta, \\
 I_h(t) &= I_h(0) + \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)}\Omega_2(t, I_h) + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t \Omega_2(\zeta, I_h) d\zeta, \\
 R_h(t) &= R_h(0) + \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)}\Omega_3(t, R_h) + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t \Omega_3(\zeta, R_h) d\zeta, \\
 S_m(t) &= S_m(0) + \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)}\Omega_4(t, S_m) + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t \Omega_4(\zeta, S_m) d\zeta, \\
 I_m(t) &= I_m(0) + \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)}\Omega_5(t, I_m) + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t \Omega_5(\zeta, I_m) d\zeta.
 \end{aligned}
 \tag{15}$$

Next, we construct the following recursive formulas:

$$\begin{aligned}
 S_{hn}(t) &= \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)}\Omega_1(t, S_{h(n-1)}) + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t (\Omega_1(\zeta, S_{h(n-1)})) d\zeta, \\
 I_{hn}(t) &= \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)}\Omega_2(t, I_{h(n-1)}) + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t (\Omega_2(\zeta, I_{h(n-1)})) d\zeta, \\
 R_{hn}(t) &= \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)}\Omega_3(t, R_{h(n-1)}) + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t (\Omega_3(\zeta, R_{h(n-1)})) d\zeta, \\
 S_{mn}(t) &= \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)}\Omega_4(t, S_{m(n-1)}) + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t (\Omega_4(\zeta, S_{m(n-1)})) d\zeta, \\
 I_{mn}(t) &= \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)}\Omega_5(t, I_{m(n-1)}) + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t (\Omega_5(\zeta, I_{m(n-1)})) d\zeta,
 \end{aligned}
 \tag{16}$$

along with the initial conditions

$$\begin{aligned}
 S_{h0} &= S_h(0), & I_{h0} &= I_h(0), & R_{h0} &= R_h(0), \\
 S_{m0} &= S_m(0), & I_{m0} &= I_m(0).
 \end{aligned}
 \tag{17}$$

We express the difference between the succession terms as

$$\begin{aligned}
 \varpi_{1n}(t) &= S_{hn}(t) - S_{h(n-1)}(t) \\
 &= \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} (\Omega_1(t, S_{h(n-1)}) - \Omega_1(t, S_{h(n-2)})) \\
 &\quad + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t (\Omega_1(\zeta, S_{h(n-1)}) - \Omega_1(\zeta, S_{h(n-2)})) d\zeta, \\
 \varpi_{2n}(t) &= I_{hn}(t) - I_{h(n-1)}(t) \\
 &= \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} (\Omega_2(t, I_{h(n-1)}) - \Omega_2(t, I_{h(n-2)})) \\
 &\quad + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t (\Omega_2(\zeta, I_{h(n-1)}) - \Omega_2(\zeta, I_{h(n-2)})) d\zeta,
 \end{aligned}$$

$$\begin{aligned}
 \varpi_{3n}(t) &= R_{hn}(t) - R_{h(n-1)}(t) \\
 &= \frac{2(1-\lambda)}{(2-\lambda)\mathcal{M}(\lambda)} (\mathcal{Q}_3(t, R_{h(n-1)}) - \mathcal{Q}_3(t, R_{h(n-2)})) \\
 &\quad + \frac{2\lambda}{(2-\lambda)\mathcal{M}(\lambda)} \int_0^t (\mathcal{Q}_3(\zeta, R_{h(n-1)}) - \mathcal{Q}_3(\zeta, R_{h(n-2)})) d\zeta, \\
 \varpi_{4n}(t) &= S_{mn}(t) - S_{m(n-1)}(t) \\
 &= \frac{2(1-\lambda)}{(2-\lambda)\mathcal{M}(\lambda)} (\mathcal{Q}_4(t, S_{m(n-1)}) - \mathcal{Q}_4(t, S_{m(n-2)})) \\
 &\quad + \frac{2\lambda}{(2-\lambda)\mathcal{M}(\lambda)} \int_0^t (\mathcal{Q}_4(\zeta, S_{h(n-1)}) - \mathcal{Q}_4(\zeta, S_{m(n-2)})) d\zeta, \\
 \varpi_{5n}(t) &= I_{mn}(t) - I_{m(n-1)}(t) \\
 &= \frac{2(1-\lambda)}{(2-\lambda)\mathcal{M}(\lambda)} (\mathcal{Q}_5(t, I_{m(n-1)}) - \mathcal{Q}_5(t, I_{m(n-2)})) \\
 &\quad + \frac{2\lambda}{(2-\lambda)\mathcal{M}(\lambda)} \int_0^t (\mathcal{Q}_5(\zeta, I_{h(n-1)}) - \mathcal{Q}_5(\zeta, I_{m(n-2)})) d\zeta.
 \end{aligned}
 \tag{18}$$

It is worth observing that

$$\begin{aligned}
 S_{hn}(t) &= \sum_{i=0}^n \varpi_{1i}(t), & I_{hn}(t) &= \sum_{i=0}^n \varpi_{2i}(t), & R_{hn}(t) &= \sum_{i=0}^n \varpi_{3i}(t), \\
 S_{mn}(t) &= \sum_{i=0}^n \varpi_{4i}(t), & I_{mn}(t) &= \sum_{i=0}^n \varpi_{5i}(t).
 \end{aligned}
 \tag{19}$$

Now we easily obtain the following result:

$$\begin{aligned}
 \|\varpi_{1n}(t)\| &= \|S_{hn}(t) - S_{h(n-1)}(t)\| \\
 &= \left\| \frac{2(1-\lambda)}{(2-\lambda)\mathcal{M}(\lambda)} (\mathcal{Q}_1(t, S_{h(n-1)}) - \mathcal{Q}_1(t, S_{h(n-2)})) \right. \\
 &\quad \left. + \frac{2\lambda}{(2-\lambda)\mathcal{M}(\lambda)} \int_0^t (\mathcal{Q}_1(\zeta, S_{h(n-1)}) - \mathcal{Q}_1(\zeta, S_{h(n-2)})) d\zeta \right\|.
 \end{aligned}
 \tag{20}$$

Applying the triangle inequality to Eq. (20), we get

$$\begin{aligned}
 \|S_{hn}(t) - S_{h(n-1)}(t)\| &\leq \frac{2(1-\lambda)}{(2-\lambda)\mathcal{M}(\lambda)} (\mathcal{Q}_1(t, S_{h(n-1)}) - \mathcal{Q}_1(t, S_{h(n-2)})) \\
 &\quad + \frac{2\lambda}{(2-\lambda)\mathcal{M}(\lambda)} \left\| \int_0^t (\mathcal{Q}_1(\zeta, S_{h(n-1)}) - \mathcal{Q}_1(\zeta, S_{h(n-2)})) d\zeta \right\|.
 \end{aligned}
 \tag{21}$$

It is already proved that the kernels satisfy the Lipschitz condition, so Eq. (21) gives

$$\begin{aligned}
 \|S_{hn}(t) - S_{h(n-1)}(t)\| &\leq \frac{2(1-\lambda)}{(2-\lambda)\mathcal{M}(\lambda)} b_1 \|S_{h(n-1)} - S_{h(n-2)}\| \\
 &\quad + \frac{2\lambda}{(2-\lambda)\mathcal{M}(\lambda)} b_1 \int_0^t \|S_{h(n-1)} - S_{h(n-2)}\| d\zeta.
 \end{aligned}
 \tag{22}$$

Consequently, we arrive at the subsequent result

$$\|\varpi_{1n}(t)\| \leq \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_1 \|\varpi_{1(n-1)}(t)\| + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_1 \int_0^t \|\varpi_{1(n-1)}(\varsigma)\| d\varsigma. \tag{23}$$

Using the same process, we derive the following results:

$$\begin{aligned} \|\varpi_{2n}(t)\| &\leq \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_2 \|\varpi_{2(n-1)}(t)\| + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_2 \int_0^t \|\varpi_{2(n-1)}(\varsigma)\| d\varsigma, \\ \|\varpi_{3n}(t)\| &\leq \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_3 \|\varpi_{3(n-1)}(t)\| + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_3 \int_0^t \|\varpi_{3(n-1)}(\varsigma)\| d\varsigma, \\ \|\varpi_{4n}(t)\| &\leq \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_4 \|\varpi_{4(n-1)}(t)\| + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_4 \int_0^t \|\varpi_{4(n-1)}(\varsigma)\| d\varsigma, \\ \|\varpi_{5n}(t)\| &\leq \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_5 \|\varpi_{5(n-1)}(t)\| + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_5 \int_0^t \|\varpi_{5(n-1)}(\varsigma)\| d\varsigma. \end{aligned} \tag{24}$$

Taking (23) and (24) into account, we obtain the existence of the solution of the considered model. □

**Theorem 2** *The SIRS-SI malaria model involving the CF fractional operator expressed in Eq. (5) has a solution if there exists  $t_0$  such that*

$$\frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_1 + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_1 t_0 < 1.$$

*Proof* As we know, the functions  $S_h(t)$ ,  $I_h(t)$ ,  $R_h(t)$ ,  $S_m(t)$ , and  $I_m(t)$  are bounded. Using the results presented in Eqs. (23)–(24) and utilizing the recursive algorithm, we get

$$\begin{aligned} \|\varpi_{1n}(t)\| &\leq \|S_{hn}(0)\| \left[ \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_1 + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_1 t \right]^n, \\ \|\varpi_{2n}(t)\| &\leq \|I_{hn}(0)\| \left[ \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_2 + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_2 t \right]^n, \\ \|\varpi_{3n}(t)\| &\leq \|R_{hn}(0)\| \left[ \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_3 + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_3 t \right]^n, \\ \|\varpi_{4n}(t)\| &\leq \|S_{mn}(0)\| \left[ \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_4 + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_4 t \right]^n, \\ \|\varpi_{5n}(t)\| &\leq \|I_{mn}(0)\| \left[ \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_5 + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_5 t \right]^n. \end{aligned} \tag{25}$$

Hence the solution of the considered model exists and is continuous.

Now, to show that Eq. (15) is a solution of the model (5), we take

$$\begin{aligned} S_h(t) - S_h(0) &= S_{nh}(t) - A_n(t), \\ I_h(t) - I_h(0) &= I_{hn}(t) - B_n(t), \\ R_h(t) - R_h(0) &= R_{hn}(t) - C_n(t), \\ S_m(t) - S_m(0) &= S_{mn}(t) - D_n(t), \\ I_m(t) - I_m(0) &= I_{mn}(t) - E_n(t). \end{aligned} \tag{26}$$

Thus we have

$$\begin{aligned} \|A_n(t)\| &= \left\| \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} (\Omega_1(t, S_h) - \Omega_1(t, S_{h(n-1)})) \right. \\ &\quad \left. + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t (\Omega_1(\zeta, S_h) - \Omega_1(\zeta, S_{h(n-1)})) d\zeta \right\| \\ &\leq \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} \|(\Omega_1(t, S_h) - \Omega_1(t, S_{h(n-1)}))\| \\ &\quad + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t \|(\Omega_1(\zeta, S_h) - \Omega_1(\zeta, S_{h(n-1)}))\| d\zeta \\ &\leq \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_1 \|S_h - S_{h(n-1)}\| + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_1 \|S_h - S_{h(n-1)}\| t. \end{aligned} \tag{27}$$

Using this process recursively, we get

$$\|A_n(t)\| \leq \left( \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} + \frac{2\lambda}{(2-\lambda)M(\lambda)} t \right)^{n+1} b_1^{n+1} a_1. \tag{28}$$

Then at  $t_0$ , we have

$$\|A_n(t)\| \leq \left( \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} + \frac{2\lambda}{(2-\lambda)M(\lambda)} t_0 \right)^{n+1} b_1^{n+1} a_1. \tag{29}$$

Taking the limit on Eq. (29) as  $n$  tends to infinity gives

$$\|A_n(t)\| \rightarrow 0.$$

Similarly, we get

$$\|B_n(t)\| \rightarrow 0, \quad \|C_n(t)\| \rightarrow 0, \quad \|D_n(t)\| \rightarrow 0, \quad \text{and} \quad \|E_n(t)\| \rightarrow 0.$$

This completes the proof of the existence theorem.

Next, we prove the uniqueness of a solution of the fractional SIRS-SI malaria model (5).

Let us assume that there exists another system of solutions of SIRS-SI malaria model (5),  $S_h^*(t)$ ,  $I_h^*(t)$ ,  $R_h^*(t)$ ,  $S_m^*(t)$ , and  $I_m^*(t)$ . Then

$$\begin{aligned} S_h(t) - S_h^*(t) &= \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} (\Omega_1(t, S_h) - \Omega_1(t, S_h^*)) \\ &\quad + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t (\Omega_1(\zeta, S) - \Omega_1(\zeta, S_h^*)) d\zeta. \end{aligned} \tag{30}$$

Taking the norms gives

$$\begin{aligned} \|S_h(t) - S_h^*(t)\| &\leq \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} \|\Omega_1(t, S_h) - \Omega_1(t, S_h^*)\| \\ &\quad + \frac{2\lambda}{(2-\lambda)M(\lambda)} \int_0^t \|(\Omega_1(\zeta, S) - \Omega_1(\zeta, S_h^*))\| d\zeta. \end{aligned} \tag{31}$$

Employing the results presented in (13) and (14), we get

$$\begin{aligned} \|S(t) - S_h^*(t)\| &\leq \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_1 \|S_h(t) - S_h^*(t)\| \\ &\quad + \frac{2\lambda}{(2-\lambda)M(\lambda)} b_1 t \|S_h(t) - S_h^*(t)\|, \end{aligned} \tag{32}$$

which gives

$$\|S_h(t) - S_h^*(t)\| \left( 1 - \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_1 - \frac{2\lambda}{(2-\lambda)M(\lambda)} b_1 t \right) \leq 0. \tag{33}$$

□

**Theorem 3** *The fractional SIRS-SI malaria model (5) has a unique solution if*

$$\left( 1 - \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} \gamma_1 - \frac{2\lambda}{(2-\lambda)M(\lambda)} \gamma_1 t \right) > 0. \tag{34}$$

*Proof* From Eq. (33) we have

$$\|S_h(t) - S_h^*(t)\| \left( 1 - \frac{2(1-\lambda)}{(2-\lambda)M(\lambda)} b_1 - \frac{2\lambda}{(2-\lambda)M(\lambda)} b_1 t \right) \leq 0. \tag{35}$$

Using Eq. (34) and properties of a norm in Eq. (35) gives

$$\|S_h(t) - S_h^*(t)\| = 0.$$

Thus we can see that

$$S_h(t) = S_h^*(t). \tag{36}$$

Using a similar procedure, we easily prove that

$$I_h = I_h^*, \quad R_h = R_h^*, \quad S_m = S_m^*, \quad I_m = I_m^*. \tag{37}$$

Thus the fractional SIRS-SI malaria model (5) has a unique solution. □

**5 HATM for fractional SIRS-SI malaria model**

In this section, we simulate the numerical results for the fractional SIRS-SI malaria model by using HATM. Firstly, we apply the Laplace transform to fractional SIRS-SI malaria model (5), which yields

$$\begin{aligned} \frac{sL[S_h] - S_h(0)}{s + \lambda(1-s)} &= L[\alpha_h + \beta R_h - (\omega\gamma_1 I_h + \xi\gamma_2 I_m)S_h - (\delta + \eta_h)S_h], \\ \frac{sL[I_h] - I_h(0)}{s + \lambda(1-s)} &= L[\mu I_h + (\omega\gamma_1 I_h + \xi\gamma_2 I_m)S_h - (\eta_h + \varepsilon + cv)I_h], \\ \frac{sL[R_h] - R_h(0)}{s + \lambda(1-s)} &= L[cv I_h - (\eta_h + \beta)R_h + \delta S_h], \end{aligned} \tag{38}$$

$$\frac{sL[S_m] - S_m(0)}{s + \lambda(1 - s)} = L[\alpha_m - (e\gamma_3 I_h + \eta_m + \sigma)S_m],$$

$$\frac{sL[I_m] - I_m(0)}{s + \lambda(1 - s)} = L[e\gamma_3 I_h S_m - (\eta_m + \sigma)I_m].$$

By simplification this gives

$$L[S_h] - \frac{c_1}{s} - \frac{[s + \lambda(1 - s)]\alpha_h}{s^2} - \frac{s + \lambda(1 - s)}{s} L[\beta R_h - (\omega\gamma_1 I_h + \xi\gamma_2 I_m)S_h - (\delta + \eta_h)S_h] = 0,$$

$$L[I_h] - \frac{c_2}{s} - \frac{s + \lambda(1 - s)}{s} L[\mu I_h + (\omega\gamma_1 I_h + \xi\gamma_2 I_m)S_h - (\eta_h + \varepsilon + cv)I_h] = 0,$$

$$L[R_h] - \frac{c_3}{s} - \frac{s + \lambda(1 - s)}{s} L[cv I_h - (\eta_h + \beta)R_h + \delta S_h] = 0,$$

$$L[S_m] - \frac{c_4}{s} - \frac{[s + \lambda(1 - s)]\alpha_m}{s^2} - \frac{s + \lambda(1 - s)}{s} L[-(e\gamma_3 I_h + \eta_m + \sigma)S_m] = 0,$$

$$L[I_m] - \frac{c_5}{s} - \frac{s + \lambda(1 - s)}{s} L[e\gamma_3 I_h S_m - (\eta_m + \sigma)I_m] = 0.$$
(39)

We present the nonlinear operators as

$$N_1[\varphi_1(t; z)] = L[\varphi_1(t; z)] - \frac{c_1}{s} - \frac{[s + \lambda(1 - s)]\alpha_h}{s^2} - \frac{s + \lambda(1 - s)}{s} L[\beta\varphi_3(t; z) - (\omega\gamma_1\varphi_2(t; z) + \xi\gamma_2\varphi_5(t; z))\varphi_1(t; z) - (\delta + \eta_h)\varphi_1(t; z)] = 0,$$

$$N_2[\varphi_2(t; z)] = L[\varphi_2(t; z)] - \frac{c_2}{s} - \frac{s + \lambda(1 - s)}{s} L[\mu\varphi_2(t; z) + (\omega\gamma_1\varphi_2(t; z) + \xi\gamma_2\varphi_4(t; z))\varphi_1(t; z) - (\eta_h + \varepsilon + cv)\varphi_2(t; z)] = 0,$$

$$N_3[\varphi_3(t; z)] = L[\varphi_3(t; z)] - \frac{c_3}{s} - \frac{s + \lambda(1 - s)}{s} L[cv\varphi_2(t; z) - (\eta_h + \beta)\varphi_3(t; z) + \delta\varphi_1(t; z)] = 0,$$

$$N_4[\varphi_4(t; z)] = L[\varphi_4(t; z)] - \frac{c_4}{s} - \frac{[s + \lambda(1 - s)]\alpha_m}{s^2} - \frac{s + \lambda(1 - s)}{s} L[-(e\gamma_3\varphi_2(t; z) + \eta_m + \sigma)\varphi_4(t; z)],$$

$$N_5[\varphi_5(t; z)] = L[\varphi_5(t; z)] - \frac{c_5}{s} - \frac{s + \lambda(1 - s)}{s} L[e\gamma_3\varphi_2(t; z)\varphi_4(t; z) - (\eta_m + \sigma)\varphi_5(t; z)] = 0,$$
(40)

and thus we have

$$\mathfrak{R}_{1,k}(\vec{S}_{h(k-1)}) = L[S_{h(k-1)}] - \left( \frac{c_1}{s} + \frac{[s + \lambda(1 - s)]\alpha_h}{s^2} \right) (1 - \chi_k) - \frac{s + \lambda(1 - s)}{s} L \left[ \beta R_{h(k-1)} - \omega\gamma_1 \left( \sum_{r=0}^{k-1} I_{hr} S_{h(k-1-r)} \right) - \xi\gamma_2 \left( \sum_{r=0}^{k-1} I_{mr} S_{h(k-1-r)} \right) - (\delta + \eta_h) S_{h(k-1)} \right],$$

$$\begin{aligned}
 \mathfrak{N}_{2,k}(\vec{I}_{h(k-1)}) &= L[I_{h(k-1)}] - \frac{c_2}{s}(1 - \chi_k) \\
 &\quad - \frac{s + \lambda(1 - s)}{s} L \left[ \mu I_{h(k-1)} + \omega \gamma_1 \left( \sum_{r=0}^{k-1} I_{hr} S_{h(k-1-r)} \right) \right. \\
 &\quad \left. + \xi \gamma_2 \left( \sum_{r=0}^{k-1} S_{mr} S_{h(k-1-r)} \right) - (\eta_h + \varepsilon + c\nu) I_{h(k-1)} \right], \\
 \mathfrak{N}_{3,k}(\vec{R}_{h(k-1)}) &= L[R_{h(k-1)}] - \frac{c_3}{s}(1 - \chi_k) \\
 &\quad - \frac{s + \lambda(1 - s)}{s} L [c\nu I_{h(k-1)} - (\eta_h + \beta) R_{h(k-1)} + \delta S_{h(k-1)}], \\
 \mathfrak{N}_{4,k}(\vec{S}_{m(k-1)}) &= L[S_{m(k-1)}] - \left( \frac{c_4}{s} + \frac{[s + \lambda(1 - s)]\alpha_m}{s^2} \right) (1 - \chi_k) \\
 &\quad - \frac{s + \lambda(1 - s)}{s} L \left[ -e\gamma_3 \left( \sum_{r=0}^{k-1} I_{hr} S_{m(k-1-r)} \right) - (\eta_m + \sigma) S_{m(k-1)} \right], \\
 \mathfrak{N}_{5,k}(\vec{I}_{m(k-1)}) &= L[I_{m(k-1)}] - \frac{c_5}{s}(1 - \chi_k) \\
 &\quad - \frac{s + \lambda(1 - s)}{s} L \left[ e\gamma_3 \left( \sum_{r=0}^{k-1} I_{hr} S_{m(k-1-r)} \right) - (\eta_m + \sigma) I_{m(k-1)} \right].
 \end{aligned} \tag{41}$$

Further, the deformation equations of  $k$ th order are expressed as

$$\begin{aligned}
 L[S_{hk}(t) - \chi_k S_{h(k-1)}(t)] &= \hbar \mathfrak{N}_{1,k}(\vec{S}_{h(k-1)}), \\
 L[I_{hk}(t) - \chi_k I_{h(k-1)}(t)] &= \hbar \mathfrak{N}_{2,k}(\vec{I}_{h(k-1)}), \\
 L[R_{hk}(t) - \chi_k R_{h(k-1)}(t)] &= \hbar \mathfrak{N}_{3,k}(\vec{R}_{h(k-1)}), \\
 L[S_{mk}(t) - \chi_k S_{m(k-1)}(t)] &= \hbar \mathfrak{N}_{4,k}(\vec{S}_{m(k-1)}), \\
 L[I_{mk}(t) - \chi_k I_{m(k-1)}(t)] &= \hbar \mathfrak{N}_{5,k}(\vec{I}_{m(k-1)}).
 \end{aligned} \tag{42}$$

Applying the inverse Laplace transform to Eq. (42) yields

$$\begin{aligned}
 S_{hk}(t) &= \chi_k S_{h(k-1)}(t) + \hbar L^{-1} [\mathfrak{N}_{1,k}(\vec{S}_{h(k-1)})], \\
 I_{hk}(t) &= \chi_k I_{h(k-1)}(t) + \hbar L^{-1} [\mathfrak{N}_{2,k}(\vec{I}_{h(k-1)})], \\
 R_{hk}(t) &= \chi_k R_{h(k-1)}(t) + \hbar L^{-1} [\mathfrak{N}_{3,k}(\vec{R}_{h(k-1)})], \\
 S_{mk}(t) &= \chi_k S_{m(k-1)}(t) + \hbar L^{-1} [\mathfrak{N}_{4,k}(\vec{S}_{m(k-1)})], \\
 I_{mk}(t) &= \chi_k I_{m(k-1)}(t) + \hbar L^{-1} [\mathfrak{N}_{5,k}(\vec{I}_{m(k-1)})].
 \end{aligned} \tag{43}$$

Taking the initial guess  $S_{h0}(t) = c_1 + \{1 + \lambda(t - 1)\}\alpha_h$ ,  $I_{h0}(t) = c_2$ ,  $R_{h0}(t) = c_3$ ,  $S_{m0}(t) = c_4 + \{1 + \lambda(t - 1)\}\alpha_m$ ,  $I_{m0}(t) = c_5$  and solving Eq. (43) for  $k = 1, 2, 3, \dots$ , we get the values of  $S_{hk}(t)$ ,  $I_{hk}(t)$ ,  $R_{hk}(t)$ ,  $S_{mk}(t)$ , and  $I_{mk}(t)$  for  $k \geq 1$ .

So, the solution of fractional SIRS-SI malaria model (5) is given as

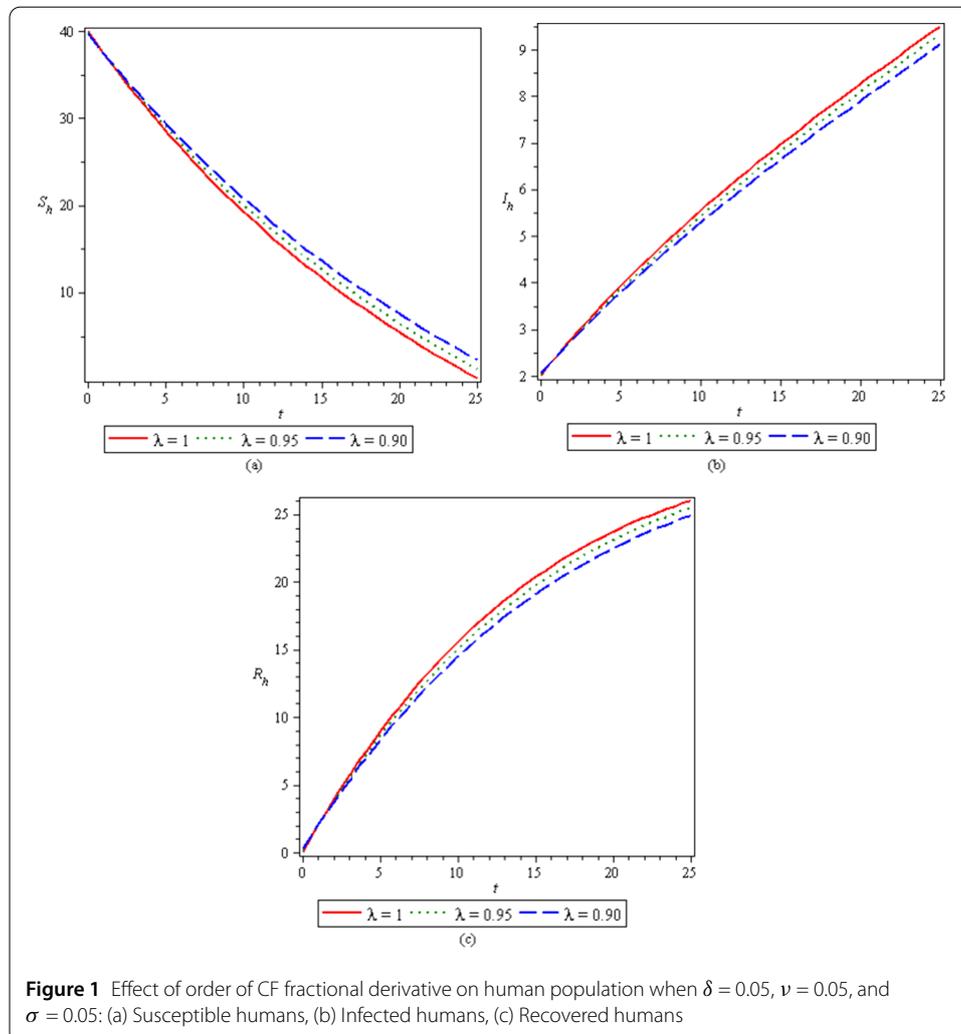
$$\begin{aligned}
 S_h(t) &= S_{h0} + S_{h1} + S_{h2} + \dots, \\
 I_h(t) &= I_{h0} + I_{h1} + I_{h2} + \dots, \\
 R_h(t) &= R_{h0} + R_{h1} + R_{h2} + \dots,
 \end{aligned} \tag{44}$$

$$S_m(t) = S_{m0} + S_{m1} + S_{m2} + \dots,$$

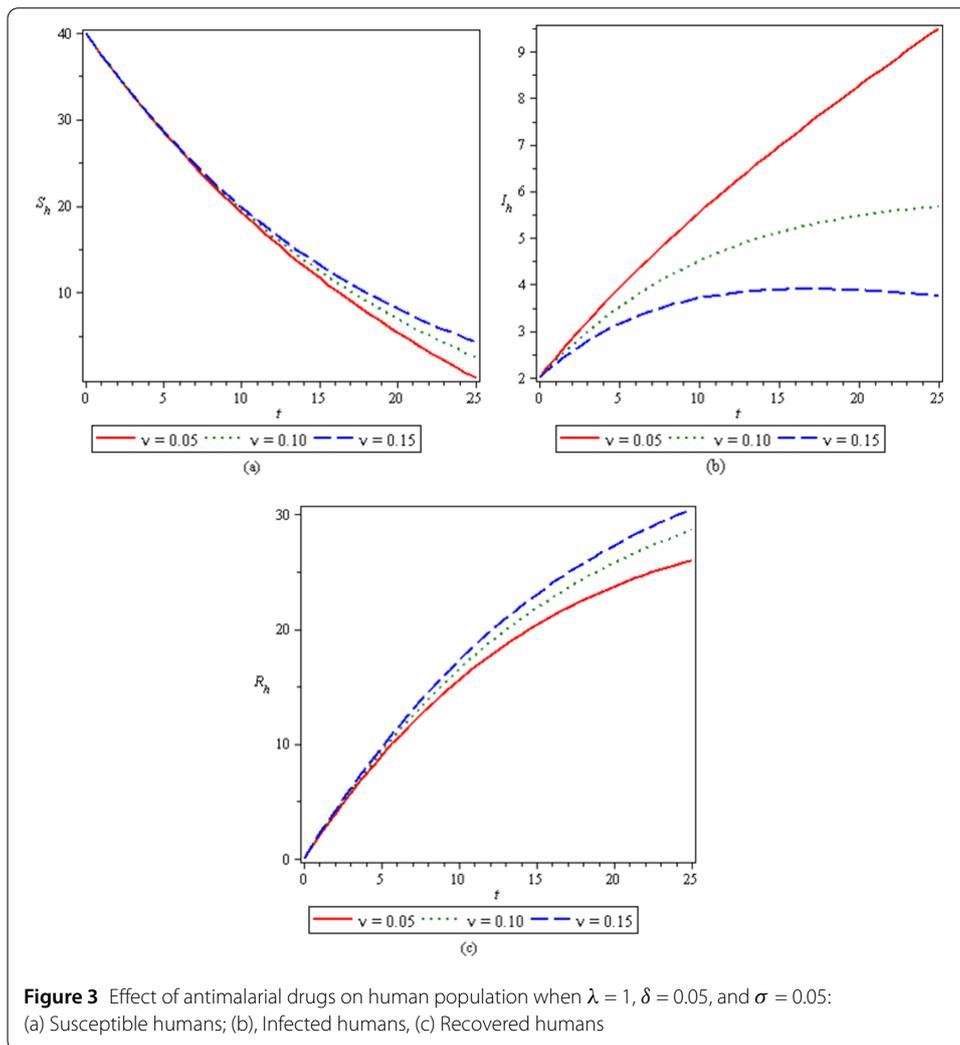
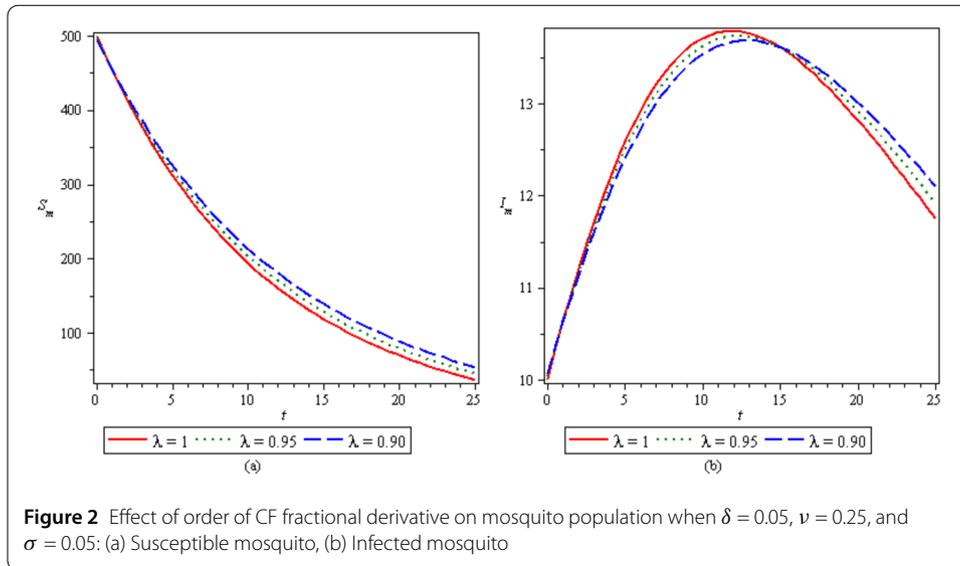
$$I_m(t) = I_{m0} + I_{m1} + I_{m2} + \dots.$$

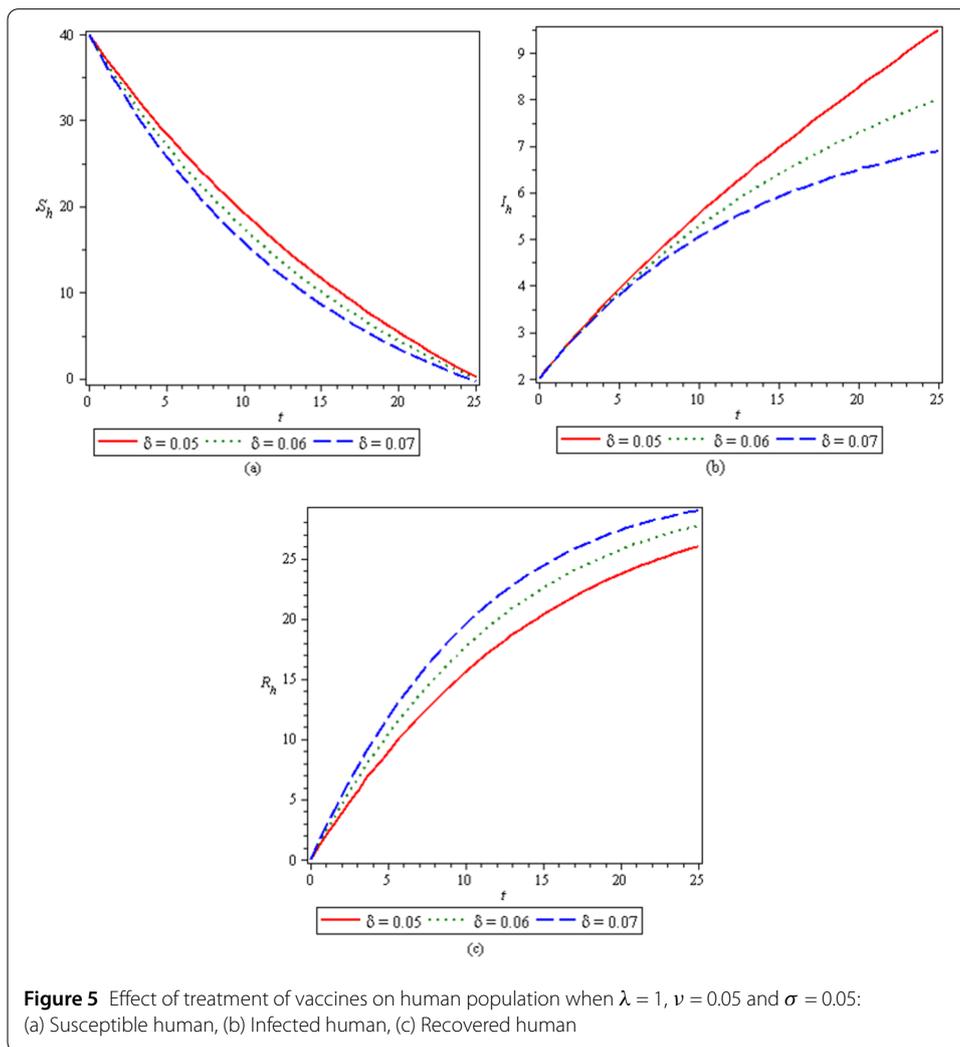
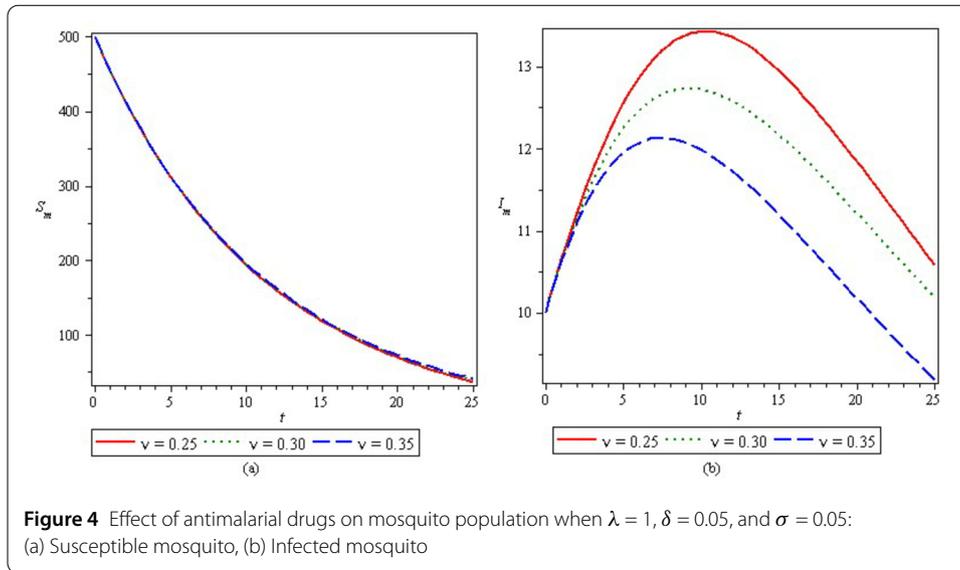
### 6 Numerical results and discussions

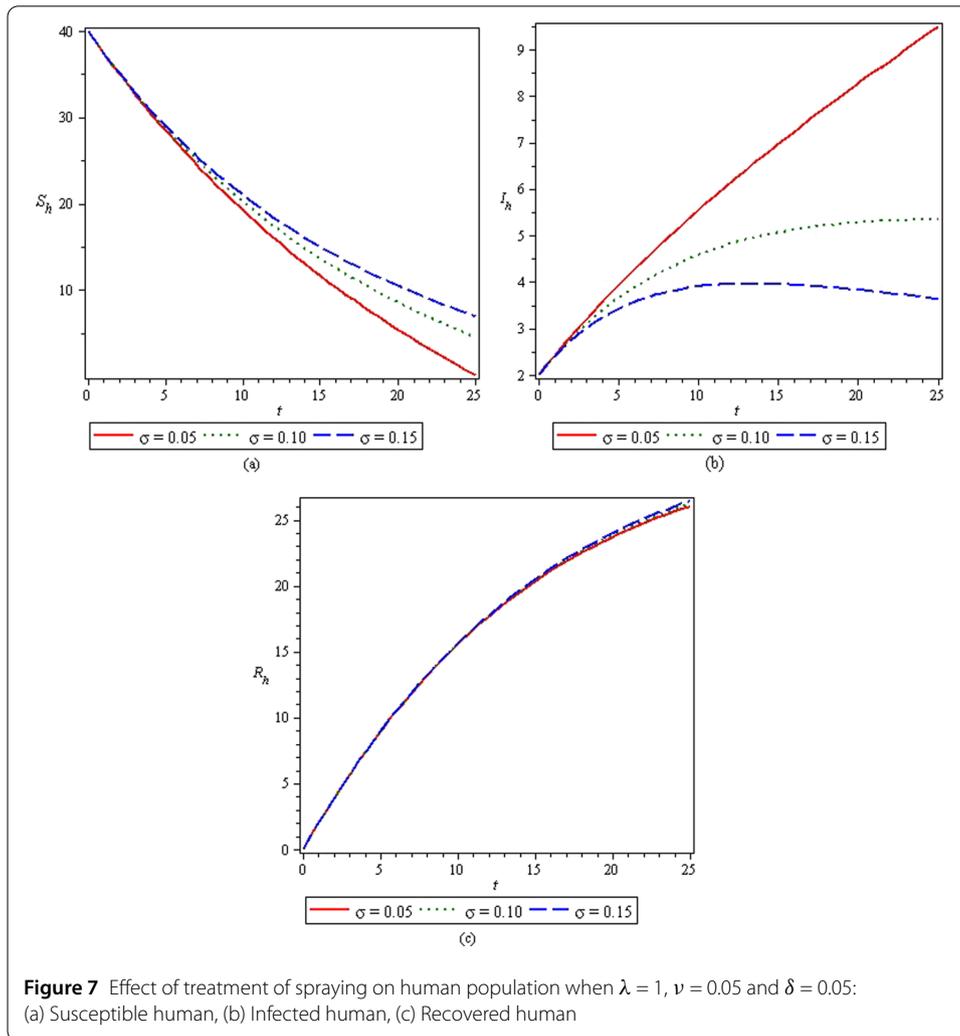
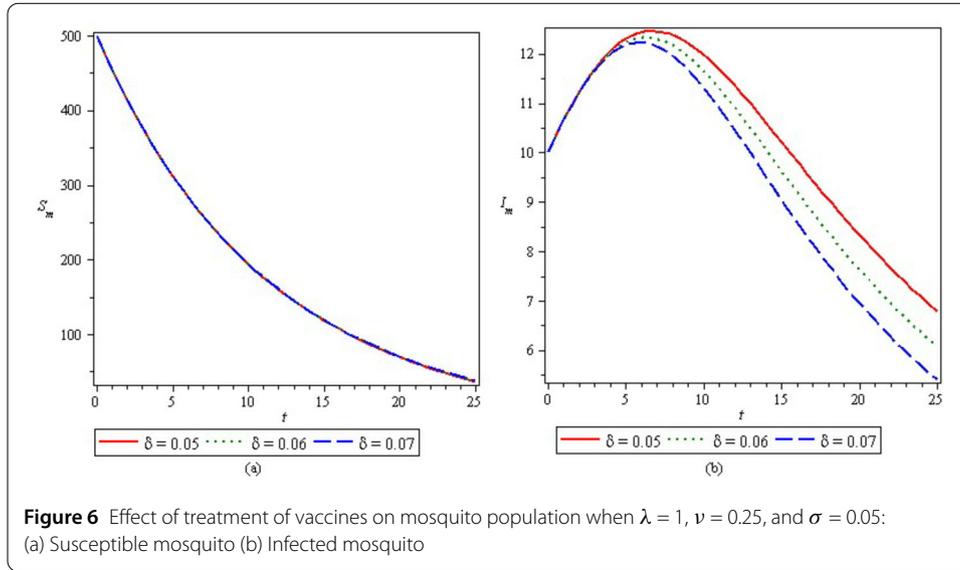
This part is devoted to numerical simulation for the fractional SIRS-SI malaria model. The numerical results for the model (5) are computed by using the HATM and Padé approximation. To compute numerical results, we have taken the values of various parameters from distinct reliable sources [1, 2, 5]. The numerical simulation is performed for  $\eta_h = 0.0004$ ,  $\eta_h = 0.04$ ,  $\varepsilon = 0.05$ ,  $\beta = 1/730$ ,  $\sigma = 1/730$ ,  $\alpha_h = 0.027$ ,  $\alpha_m = 0.13$ ,  $\omega = 0.038$ ,  $\xi = 0.13$ ,  $\gamma_1 = 0.02$ ,  $\gamma_2 = 0.010$ ,  $\gamma_3 = 0.072$ ,  $c = 0.611$ ,  $e = 0.022$ ,  $\mu = 0.005$ ,  $\sigma \in [0, 1]$ ,  $\delta \in [0, 1]$ , and  $\nu \in [0.01, 1]$ . The initial conditions are taken as  $S_h(0) = c_1 = 40$ ,  $I_h(0) = c_2 = 2$ ,  $R_h(0) = c_3 = 0$ ,  $S_m(0) = c_4 = 500$ , and  $I_m(0) = c_5 = 10$ . Figure 1 depicts the influence of order of the CF fractional operator on different groups of human population, that is, susceptible humans, infected humans, and recovered humans. Figure 2 demonstrates the influence of order of the CF fractional derivative on different groups of mosquito population, that is, susceptible mosquitoes and infected mosquitoes. Figure 3 shows the influence of anti-malarial drugs on different classes of human population. Figure 4 displays the impact of

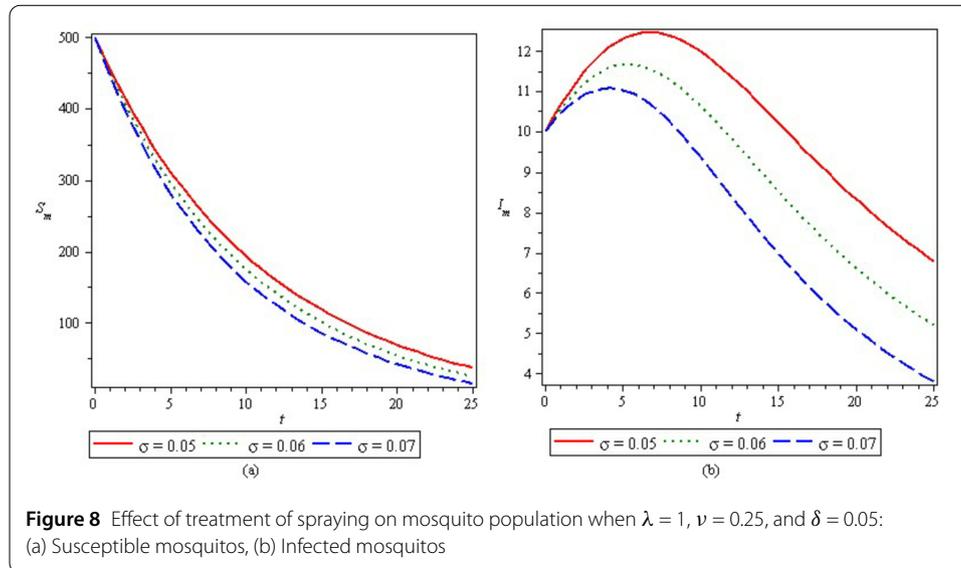


**Figure 1** Effect of order of CF fractional derivative on human population when  $\delta = 0.05$ ,  $\nu = 0.05$ , and  $\sigma = 0.05$ : (a) Susceptible humans, (b) Infected humans, (c) Recovered humans









antimalarial drugs on different classes of mosquito population. Figure 5 presents the influence of treatment of vaccines on various classes of human population. Figure 6 exhibits the effect of treatment of vaccines on various groups of mosquito population. In Fig. 7, the influence of treatment of spraying on various classes of human population is displayed. In Fig. 8, the impact of treatment of spraying on distinct classes of mosquito population is shown.

## 7 Conclusions

In this paper, we studied a fractional SIRS-SI malaria model transmission along with a number of cures such as the utilization of vaccines, antimalarial medicines, and spraying. The power of this model was inclusion of memory effects. The theory of fixed point was employed to examine the existence and uniqueness of solution of the considered fractional SIRS-SI model describing the spreading of malaria. The HATM and Padé approximation were applied to perform numerical simulation. The effects of order of the CF fractional derivative, vaccines, antimalarial drugs, and spraying on different groups of human populations and mosquito populations were analyzed. From the results we conclude that the CF fractional derivative is very useful for describing the treatment and control of the malaria disease and similar type of problems.

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### Competing interests

The authors declare that they have no competing interests.

### Authors' contributions

DK, JS, MAQ, and DB designed the study, developed the methodology, collected the data, performed the analysis, and wrote the manuscript. All authors read and approved the final manuscript.

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