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A new mathematical model for Zika virus transmission

Shahram Rezapour^{1,2,3}, Hakimeh Mohammadi^{4*} and Amin Jajarmi^{5,6}

*Correspondence:

hakimeh.mohammadi.02@gmail.com

⁴Department of Mathematics,
Miandoab Branch, Islamic Azad
University, Miandoab, Iran

Full list of author information is
available at the end of the article

Abstract

We present a new mathematical model for the transmission of Zika virus between humans as well as between humans and mosquitoes. In this way, we use the fractional-order Caputo derivative. The region of the feasibility of system and equilibrium points are calculated, and the stability of equilibrium point is investigated. We prove the existence of a unique solution for the model by using the fixed point theory. By using the fractional Euler method, we get an approximate solution to the model. Numerical results are presented to investigate the effect of fractional derivative on the behavior of functions and also to compare the integer-order derivative and fractional-order derivative results.

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1 Introduction

Zika virus was first detected in monkeys in 1947, and the first cases of Zika virus infection were reported in 1952 in Uganda and the Republic of Tanzania. Between 1960 and 1980, there were few human infections with Zika virus in Asia and Africa, until in 2007 in the Yap Islands, then in 2013 in the France Polynesia and the Pacific Ocean, and in 2015 in Brazil the widespread outbreak of the virus was reported. Subsequently, it has spread to other countries around the world, so far Zika virus infection has been recorded in 86 countries.

Fever, rash, conjunctivitis, muscle and joint pain, malaise, or headache are symptoms of this disease. According to the report of World Health Organization (WHO), Zika virus infection during pregnancy can cause infants to be born with microcephaly and other congenital malformations, known as congenital Zika syndrome. Also, at the result of this disease, other complications of pregnancy including preterm birth and miscarriage may happen. In adults and children, Zika virus infection is associated with neurologic complications including Guillain–Barre syndrome, neuropathy, and myelitis.

Zika is a virus that is spread mostly by *Aedes* mosquitoes. Besides that, this virus can be transmitted from mother to baby during pregnancy or around the time of birth. Also, it can spread through blood transfusions and sexual contact.

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The study of diseases dynamics is a dominating theme for many biologists and mathematicians (see, for example, [1–3]). It has been studied by many researchers that fractional extensions of mathematical models of integer order represent the natural fact in a very systematic way such as in the approach of Baleanu et al. [4–17]. Studies on the methods of solving fractional differential equations and the application of fractional systems have also been conducted [18–24]. In recent years, fractional-order derivatives have expanded and have been widely used in modeling real-world phenomena and investigating the process of disease transmission and control (see, for example, [25–35]). Also, some studies in the biological models with fractional-order derivative have been conducted in recent years [36–39]. During last years researchers have been using some mathematical models to simulate the transmission of Zika virus [40–43].

In mathematical models of Zika virus transmission it is assumed that the virus is usually transmitted from mosquitoes to humans, while according to WHO, in addition to the transmission through mosquitoes, Zika virus is transmitted through infected blood as well as through sexual contact with an infected person. In this article, we consider a mathematical model based on both ways of transmitting the virus. Also, according to the good results of fractional-order derivative in the modeling of real-world phenomena in recent years, we use Caputo fractional-order derivative instead of the integer-order derivative in this model.

The structure of the paper is as follows. In Sect. 2 some basic definitions and concepts of fractional calculus are recalled. The transmission model of Zika virus with fractional-order derivative is presented in Sect. 3, and the equilibrium points and the reproduction number are calculated. The existence and uniqueness of solution for the system are proved in Sect. 4. Numerical method and numerical results are presented in Sect. 5.

2 Preliminaries

In this section, we recall some basic concepts of fractional differential calculus.

Definition 2.1 ([44]) For an integrable function g , the Caputo derivative of fractional order $\nu \in (0, 1)$ is given by

$${}^C D^\nu g(t) = \frac{1}{\Gamma(m - \nu)} \int_0^t \frac{g^{(m)}(\nu)}{(t - \nu)^{\nu - m + 1}} d\nu, \quad m = [\nu] + 1.$$

Also, the corresponding fractional integral of order ν with $\text{Re}(\nu) > 0$ is given by

$${}^C I^\nu g(t) = \frac{1}{\Gamma(\nu)} \int_0^t (t - \nu)^{\nu - 1} g(\nu) d\nu.$$

Definition 2.2 ([45, 46]) For $g \in H^1(c, d)$ and $d > c$, the Caputo–Fabrizio derivative of fractional order $\nu \in (0, 1)$ for g is given by

$${}^{CF} D^\nu g(t) = \frac{M(\nu)}{(1 - \nu)} \int_c^t \exp\left(\frac{-\nu}{1 - \nu}(t - \nu)\right) g'(\nu) d\nu,$$

where $t \geq 0$, $M(\nu)$ is a normalization function that depends on ν and $M(0) = M(1) = 1$. If $g \notin H^1(c, d)$ and $0 < \nu < 1$, this derivative for $g \in L^1(-\infty, d)$ is given by

$${}^{CF} D^\nu g(t) = \frac{\nu M(\nu)}{(1 - \nu)} \int_{-\infty}^d (g(t) - g(\nu)) \exp\left(\frac{-\nu}{1 - \nu}(t - \nu)\right) d\nu.$$

Also, the corresponding *CF* fractional integral is presented by

$${}^{CF}I^\nu g(t) = \frac{2(1-\nu)}{(2-\nu)M(\nu)}g(t) + \frac{2\nu}{(2-\nu)M(\nu)} \int_0^t g(v) dv.$$

The Laplace transform is one of the important tools in solving differential equations that are defined below for two kinds of fractional derivative.

Definition 2.3 ([44]) The Laplace transform of Caputo fractional differential operator of order ν is given by

$$L[{}^C D^\nu g(t)](s) = s^\nu Lg(t) - \sum_{i=0}^{m-1} s^{\nu-i-1} g^{(i)}(0), \quad m-1 < \nu \leq m \in N,$$

which can also be obtained in the form

$$L[{}^C D^\nu g(t)] = \frac{s^m L[g(t)] - s^{m-1}g(0) - s^{m-1}g'(0) - \dots - g^{(m-1)}}{s^{m-\nu}}.$$

3 Model formulation

In this section, we provide a mathematical model for the transmission of Zika virus using the Caputo derivative of fractional order. We divide the human population N_h into two groups: susceptible people S_h and infected people I_h so that $N_h = S_h + I_h$. Similarly, we divide the total number of mosquitoes N_m into two groups: susceptible mosquitoes S_m and infected mosquitoes I_m so that $N_m = S_m + I_m$. To describe the mechanism of the spread of Zika virus, we consider the compartmental mathematical model as follows:

$$\begin{cases} \frac{dS_h}{dt} = \Lambda_h - \beta_1 S_h I_h - \beta_2 S_h I_m - k_1 S_h, \\ \frac{dI_h}{dt} = \beta_1 S_h I_h + \beta_2 S_h I_m - k_1 I_h, \\ \frac{dS_m}{dt} = \Lambda_m - \mu S_m I_h - k_2 S_m, \\ \frac{dI_m}{dt} = \mu S_m I_h - k_2 I_m, \end{cases} \tag{1}$$

with the initial conditions $S_h(0) = S_{0h}, I_h(0) = I_{0h}, S_m(0) = S_{0m}, I_m(0) = I_{0m}$.

The model parameters are: the recruitment rate of human population Λ_h , the recruitment rate of mosquito population Λ_m , the effective contact rate human to human β_1 , the effective contact rate mosquitoes to human β_2 , the effective contact rate human to mosquitoes μ , the natural death rate of human k_1 , the natural death rate of mosquitoes k_2 .

Model (1) does not include the internal memory effects of the system. To improve the model, we change the first-order time derivative to the Caputo fractional derivative of order ν . With this change, the right- and left-hand sides will not have the same dimension. To solve this problem, we use an auxiliary parameter θ , having the dimension of sec., to change the fractional operator so that the sides have the same dimension ([47, 48]). According to the explanation presented, the transmission model of Zika virus for $t \geq 0$ and

$\nu \in (0, 1)$ is given as follows:

$$\begin{cases} \theta^{\nu-1} {}^C D_t^\nu S_h(t) = \Lambda_h - \beta_1 S_h I_h - \beta_2 S_h I_m - k_1 S_h, \\ \theta^{\nu-1} {}^C D_t^\nu I_h(t) = \beta_1 S_h I_h + \beta_2 S_h I_m - k_1 I_h, \\ \theta^{\nu-1} {}^C D_t^\nu S_m(t) = \Lambda_m - \mu S_m I_h - k_2 S_m, \\ \theta^{\nu-1} {}^C D_t^\nu I_m(t) = \mu S_m I_h - k_2 I_m, \end{cases} \tag{2}$$

where the initial conditions are $S_h(0) = S_{0h}, I_h(0) = I_{0h}, S_m(0) = S_{0m}, I_m(0) = I_{0m}$.

3.1 Nonnegative solution

Consider $\Phi = \{(S_h, I_h, S_m, I_m) \in R_4^+ : N_h(t) \leq \frac{\Lambda_h}{k_1}, N_m(t) \leq \frac{\Lambda_m}{k_2}\}$, we show that the closed set Φ is the region of the feasibility of system (2).

Lemma 3.1 *The closed set Φ is positively invariant with respect to fractional system (2).*

Proof To obtain the fractional derivative of the total population, we add the first two relations in system (2). So

$$\theta^{\nu-1} {}^C D_t^\nu N_h(t) = \Lambda_h - k_1 N_h(t),$$

where $N_h(t) = S_h(t) + I_h(t)$. Using the Laplace transform, we obtain

$$N_h(t) = N_h(0)E_\nu(-k_1 \theta^{1-\nu} t^\nu) + \int_0^t \Lambda_h \theta^{1-\nu} \eta^{\nu-1} E_{\nu,\nu}(-k_1 \theta^{1-\nu} \eta^\nu) d\eta,$$

where $N_h(0)$ is the initial human population size, and the terms $E_\nu, E_{\nu,\nu}$ in the above equation are represented by the Mittag-Leffler function and its general form defined by

$$E_\nu(z) = \sum_{j=0}^{\infty} \frac{z^j}{\Gamma(1 + j\nu)}, \quad E_{\nu,\nu}(z) = \sum_{j=0}^{\infty} \frac{z^j}{\Gamma(\nu + j\nu)}, \quad \nu > 0.$$

With some calculations, we get

$$\begin{aligned} N_h(t) &= N_h(0)E_\nu(-k_1 \theta^{1-\nu} t^\nu) + \int_0^t \Lambda_h \theta^{1-\nu} \eta^{\nu-1} \sum_{i=0}^{\infty} \frac{(-1)^i k_1^i \theta^{i(1-\nu)} \eta^{i\nu}}{\Gamma(i\nu + \nu)} d\eta \\ &= \frac{\Lambda_h \theta^{1-\nu}}{k_1 \theta^{1-\nu}} + E_\nu(-k_1 \theta^{1-\nu} t^\nu) \left(N_h(0) - \frac{\Lambda_h \theta^{1-\nu}}{k_1 \theta^{1-\nu}} \right), \\ &= \frac{\Lambda_h}{k_1} + E_\nu(-k_1 \theta^{1-\nu} t^\nu) \left(N_h(0) - \frac{\Lambda_h}{k_1} \right). \end{aligned}$$

Thus, if $N_h(0) \leq \frac{\Lambda_h}{k_1}$, then for $t > 0, N_h(t) \leq \frac{\Lambda_h}{k_1}$. Similarly, we can prove for N_m that if $N_m(0) \leq \frac{\Lambda_m}{k_2}$, then for $t > 0, N_m(t) \leq \frac{\Lambda_m}{k_2}$. Consequently, the closed set Φ is positively invariant with respect to fractional model (2). \square

3.2 Equilibrium points and reproduction number

To determine the equilibrium points of fractional order system (2), we solve the following equations:

$${}^C D^\nu S_h(t) = {}^C D^\nu I_h(t) = {}^C D^\nu S_m(t) = {}^C D^\nu I_m(t) = 0.$$

By solving the above algebraic equations, we obtain two equilibrium points of system (2). The disease-free equilibrium point is obtained as $E^0 = (\frac{\Lambda_h}{k_1}, 0, \frac{\Lambda_m}{k_2})$. In addition, if $R_0 > 1$, then system (2) has a positive endemic equilibrium point $E^* = (S_h^*, I_h^*, S_m^*, I_m^*)$, where

$$S_h^* = \frac{k_2 k_1}{(\beta_2 \mu S_m^* + k_2 \beta_1)},$$

$$I_h^* = \frac{\Lambda_h \beta_2 \mu S_m^* + \Lambda_h k_2 \beta_1 - k_2 k_1^2}{k_1 (\beta_2 \mu S_m^* + k_2 \beta_1)},$$

$$I_m^* = \frac{\mu (\Lambda_h \beta_2 \mu S_m^* + \Lambda_h k_2 \beta_1 - k_2 k_1^2) S_m^*}{k_1 (\beta_2 \mu S_m^* + k_2 \beta_1) k_2}.$$

Also, R_0 is the basic reproduction number and is obtained using the next generation method [49]. To find R_0 , we first consider the system as follows:

$${}^C D^\nu \Psi(t) = F(\Psi(t)) - V(\Psi(t)),$$

where

$$F(\Psi(t)) = \theta^{1-\nu} \begin{bmatrix} \beta_1 S_h I_h + \beta_2 S_h I_m \\ \mu S_m I_h \end{bmatrix}$$

and

$$V(\Psi(t)) = \theta^{1-\nu} \begin{bmatrix} k_1 I_h \\ k_2 I_m \end{bmatrix}.$$

At E^0 , the Jacobian matrix for F and V is obtained as follows:

$$J_F(E_0) = \theta^{1-\nu} \begin{bmatrix} \frac{\beta_1 \Lambda_h}{k_1} & \frac{\beta_2 \Lambda_h}{k_2} \\ \frac{\mu \Lambda_m}{k_2} & 0 \end{bmatrix}, \quad J_V(E_0) = \theta^{1-\nu} \begin{bmatrix} k_1 & 0 \\ 0 & k_2 \end{bmatrix}.$$

FV^{-1} is the next generation matrix for the system (2), and the basic reproduction number is obtained from $R_0 = \rho(FV^{-1})$, where $\rho(FV^{-1})$ is the eigenvalue of matrix FV^{-1} . We get

$$R_0 = \frac{\beta_1 k_2 \Lambda_h + \sqrt{\beta_1^2 \Lambda_h^2 k_2^2 + 4 k_1^2 \beta_2 \mu \Lambda_h \Lambda_m}}{2 k_2 k_1^2}.$$

This basic reproduction number R_0 is an epidemiologic metric used to describe the contagiousness or transmissibility of infectious agents.

3.3 Stability of equilibrium point

To investigate the stability of the equilibrium point, we first consider the Jacobian matrix of system (2) as follows:

$$J = \theta^{1-\nu} \begin{bmatrix} -\beta_1 I_h - \beta_2 I_m - k_1 & -\beta_1 S_h & 0 & -\beta_2 S_h \\ \beta_1 I_h + \beta_2 I_m & \beta_1 S_h - k_1 & 0 & \beta_2 S_h \\ 0 & -\mu S_m & -\mu I_h - k_2 & 0 \\ 0 & \mu S_m & \mu I_h & -k_2 \end{bmatrix}.$$

At E_0 , the Jacobian matrix of system (2) is

$$J = \theta^{1-\nu} \begin{bmatrix} -k_1 & -\beta_1 \frac{\Lambda_h}{k_1} & 0 & -\beta_2 \frac{\Lambda_h}{k_1} \\ 0 & \beta_1 \frac{\Lambda_h}{k_1} - k_1 & 0 & \beta_2 \frac{\Lambda_h}{k_1} \\ 0 & -\mu \frac{\Lambda_m}{k_2} & -k_2 & 0 \\ 0 & \mu \frac{\Lambda_m}{k_2} & 0 & -k_2 \end{bmatrix}.$$

Theorem 3.2 *If $R_0 < 1$, the equilibrium point E^0 of system (2) is locally asymptotically stable.*

Proof At the disease-free equilibrium point E^0 , the characteristic equation of the Jacobian matrix is $\det(\lambda I - J(E^0)) = 0$. Then we obtain

$$\theta^{1-\nu}(\lambda + k_1)(\lambda + k_2)(\lambda^2 + B\lambda + C) = 0,$$

where $B = k_1 + k_2 - \frac{\beta_1 \Lambda_h}{k_1}$ and $C = 2k_1 k_2 - \frac{k_2 \beta_1 \Lambda_h}{k_1} - \frac{\mu \beta_2 \Lambda_m \Lambda_h}{k_1 k_2}$. By simplifying the above equations, the eigenvalues of characteristic equation are obtained as $\lambda_1 = -k_1$, $\lambda_2 = -k_2$ and the roots of the equation

$$\lambda^2 + B\lambda + C = 0.$$

If $R_0 < 1$, since all of the parameters are positive, then

$$\frac{\beta_1 k_2 \Lambda_h + \sqrt{\beta_1^2 \Lambda_h^2 k_2^2}}{2k_2 k_1^2} < 1 \Rightarrow \frac{\beta_1 \Lambda_h}{k_1} < k_1 \Rightarrow B = k_1 + k_2 - \frac{\beta_1 \Lambda_h}{k_1} > k_2 > 0.$$

Also, from $R_0 < 1$ we have

$$\begin{aligned} \frac{\sqrt{4k_1^2 \beta_2 \mu \Lambda_h \Lambda_m}}{2k_2 k_1^2} &< 1 \\ \Rightarrow \frac{\beta_2 \mu \Lambda_m \Lambda_h}{k_1 k_2} &< k_1 k_2, \\ \Rightarrow 2k_1 k_2 - \frac{k_2 \beta_1 \Lambda_h}{k_1} - \frac{\mu \beta_2 \Lambda_m \Lambda_h}{k_1 k_2} &> 2k_1 k_2 - k_1 k_2 - k_1 k_2 \Rightarrow C > 0. \end{aligned}$$

Since $B > 0$, $C > 0$, applying the Routh–Hurwitz criteria, we obtain that E_0 is locally asymptotically stable. □

4 Existence and uniqueness of solution

To show that the system has a unique solution, we write system (2) as follows:

$$\begin{cases} \theta^{\nu-1} {}^C D_t^\nu S_h(t) = W_1(t, S_h(t)), \\ \theta^{\nu-1} {}^C D_t^\nu I_h(t) = W_2(t, I_h(t)), \\ \theta^{\nu-1} {}^C D_t^\nu S_m(t) = W_3(t, S_m(t)), \\ \theta^{\nu-1} {}^C D_t^\nu I_m(t) = W_4(t, I_m(t)). \end{cases}$$

By applying integral on both sides of the above equations, we have

$$\begin{cases} S_h(t) - S_h(0) = \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t W_1(\tau, S_h)(t - \tau)^{\nu-1} d\tau, \\ I_h(t) - I_h(0) = \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t W_2(\tau, I_h)(t - \tau)^{\nu-1} d\tau, \\ S_m(t) - S_m(0) = \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t W_3(\tau, S_m)(t - \tau)^{\nu-1} d\tau, \\ I_m(t) - I_m(0) = \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t W_4(\tau, I_m)(t - \tau)^{\nu-1} d\tau. \end{cases} \tag{3}$$

We show that the kernels $W_i, i = 1, 2, 3, 4$, satisfy the Lipschitz condition and contraction.

Theorem 4.1 *The kernel W_1 satisfies the Lipschitz condition and contraction if the following inequality holds:*

$$0 \leq \beta_1 u_1 + \beta_2 u_2 + k_1 < 1.$$

Proof For S_h and S_{1h} , we have

$$\begin{aligned} \|W_1(t, S_h) - W_1(t, S_{1h})\| &= \|\beta_1 I_h(S_h - S_{1h}) + \beta_2 I_m(S_h - S_{1h}) + k_1(S_h - S_{1h})\|, \\ &\leq \beta_1 \|I_h\| \|S_h - S_{1h}\| + \beta_2 \|I_m\| \|S_h - S_{1h}\| + k_1 \|S_h - S_{1h}\|, \\ &\leq (\beta_1 \|I_h\| + \beta_2 \|I_m\| + k_1) \|S_h - S_{1h}\|, \\ &\leq (\beta_1 u_1 + \beta_2 u_2 + k_1) \|S_h - S_{1h}\|. \end{aligned}$$

Suppose that $M_1 = \beta_1 u_1 + \beta_2 u_2 + k_1$, where $\|I_h(t)\| \leq u_1, \|I_m\| \leq u_2$ are bounded functions, then

$$\|W_1(t, S_h) - W_1(t, S_{1h})\| \leq M_1 \|S_h(t) - S_{1h}(t)\|. \tag{4}$$

Thus, for W_1 , the Lipschitz condition is obtained, and if $0 \leq \beta_1 u_1 + \beta_2 u_2 + k_1 < 1$ then W_1 is a contraction. \square

Similarly, we can prove that $W_i, i = 2, 3, 4$, satisfies the Lipschitz condition as follows:

$$\begin{cases} \|W_2(t, I_h) - W_2(t, I_{1h})\| \leq M_2 \|I_h(t) - I_{1h}(t)\|, \\ \|W_3(t, S_m) - W_3(t, S_{1m})\| \leq M_3 \|S_m(t) - S_{1m}(t)\|, \\ \|W_4(t, I_m) - W_4(t, I_{1m})\| \leq M_4 \|I_m(t) - I_{1m}(t)\|, \end{cases}$$

where $\|S_h(t)\| \leq u_3, \|S_m(t)\| \leq u_4$, and $M_2 = \beta_1 u_3 + k_1, M_3 = \mu u_1 + k_2, M_4 = k_2$ are bounded functions, if $0 \leq M_i < 1, i = 2, 3, 4$, then $W_i, i = 2, 3, 4$, are contraction.

According to system (3), consider the following recursive forms:

$$\begin{aligned} \Phi_{1n}(t) &= S_{nh}(t) - S_{(n-1)h}(t) = \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t (W_1(\tau, S_{(n-1)h}) - W_1(\tau, S_{(n-2)h}))(t - \tau)^{\nu-1} d\tau, \\ \Phi_{2n}(t) &= I_{nh}(t) - I_{(n-1)h}(t) = \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t (W_2(\tau, I_{(n-1)h}) - W_2(\tau, I_{(n-2)h}))(t - \tau)^{\nu-1} d\tau, \\ \Phi_{3n}(t) &= S_{nh}(t) - S_{(n-1)h}(t) = \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t (W_3(\tau, S_{(n-1)h}) - W_3(\tau, S_{(n-2)h}))(t - \tau)^{\nu-1} d\tau, \\ \Phi_{4n}(t) &= I_{nh}(t) - I_{(n-1)h}(t) = \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t (W_4(\tau, I_{(n-1)h}) - W_4(\tau, I_{(n-2)h}))(t - \tau)^{\nu-1} d\tau, \end{aligned}$$

with the initial conditions $S_{0h}(t) = S_h(0)$, $I_{0h}(t) = I_h(0)$, $S_{0m}(t) = S_m(0)$, and $I_{0m}(t) = I_m(0)$. We take the norm of the first equation in the above system, then

$$\begin{aligned} \|\Phi_{1n}(t)\| &= \|S_{nh}(t) - S_{(n-1)h}(t)\| \\ &= \left\| \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t (W_1(\tau, S_{(n-1)h}) - W_1(\tau, S_{(n-2)h}))(t - \tau)^{\nu-1} d\tau \right\| \\ &\leq \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t \|W_1(\tau, S_{(n-1)h}) - W_1(\tau, S_{(n-2)h}))(t - \tau)^{\nu-1}\| d\tau. \end{aligned}$$

By Lipschitz condition (4), we have

$$\|\Phi_{1n}(t)\| \leq \frac{\theta^{1-\nu}}{\Gamma(\nu)} M_1 \int_0^t \|\Phi_{1(n-1)}(\tau)\| d\tau. \tag{5}$$

In a similar way, we obtained

$$\begin{aligned} \|\Phi_{2n}(t)\| &\leq \frac{\theta^{1-\nu}}{\Gamma(\nu)} M_2 \int_0^t \|\Phi_{2(n-1)}(\tau)\| d\tau, \\ \|\Phi_{3n}(t)\| &\leq \frac{\theta^{1-\nu}}{\Gamma(\nu)} M_3 \int_0^t \|\Phi_{3(n-1)}(\tau)\| d\tau, \\ \|\Phi_{4n}(t)\| &\leq \frac{\theta^{1-\nu}}{\Gamma(\nu)} M_4 \int_0^t \|\Phi_{4(n-1)}(\tau)\| d\tau. \end{aligned} \tag{6}$$

Then we can obtain

$$\begin{aligned} S_{nh}(t) &= \sum_{i=1}^n \Phi_{1i}(t), & I_{nh}(t) &= \sum_{i=1}^n \Phi_{2i}(t), \\ S_{nm}(t) &= \sum_{i=1}^n \Phi_{3i}(t), & I_{nm}(t) &= \sum_{i=1}^n \Phi_{4i}(t). \end{aligned}$$

We prove the existence of a solution in the next theorem.

Theorem 4.2 *The fractional model of Zika virus (2) has a solution if there exists t_1 such that*

$$\frac{\theta^{1-\nu}}{\Gamma(\nu)} t_1 M_i < 1.$$

Proof From the recursive technique and Eq. (5) and Eq. (6), we conclude that

$$\begin{aligned} \|\Phi_{1n}(t)\| &\leq \|S_{nh}(0)\| \left[\frac{\theta^{1-\nu}}{\Gamma(\nu)} M_1 t \right]^n, \\ \|\Phi_{2n}(t)\| &\leq \|I_{nh}(0)\| \left[\frac{\theta^{1-\nu}}{\Gamma(\nu)} M_2 t \right]^n, \\ \|\Phi_{3n}(t)\| &\leq \|S_{nm}(0)\| \left[\frac{\theta^{1-\nu}}{\Gamma(\nu)} M_3 t \right]^n, \\ \|\Phi_{4n}(t)\| &\leq \|I_{nm}(0)\| \left[\frac{\theta^{1-\nu}}{\Gamma(\nu)} M_4 t \right]^n. \end{aligned}$$

Then the system has a solution, and also it is continuous. Now we show that the above functions construct a solution for model (2). We assume that

$$\begin{aligned} S_h(t) - S_h(0) &= S_{nh}(t) - B_{1n}(t), \\ I_h(t) - I_h(0) &= I_{nh}(t) - B_{2n}(t), \\ S_m(t) - S_m(0) &= S_{nm}(t) - B_{3n}(t), \\ I_m(t) - I_m(0) &= I_{nm}(t) - B_{4n}(t). \end{aligned}$$

Thus

$$\begin{aligned} \|B_{1n}(t)\| &= \left\| \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t (W_1(\tau, S_h) - W_1(\tau, S_{(n-1)h})) d\tau \right\| \\ &\leq \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t \|W_1(\tau, S_h) - W_1(\tau, S_{(n-1)h})\| d\tau \\ &\leq \frac{\theta^{1-\nu}}{\Gamma(\nu)} M_1 \|S_h - S_{(n-1)h}\| t. \end{aligned}$$

By repeating the method, we obtain

$$\|B_{1n}(t)\| \leq \left[\frac{\theta^{1-\nu}}{\Gamma(\nu)} t \right]^{n+1} M_1^{n+1} h.$$

At t_1 , we get

$$\|B_{1n}(t)\| \leq \left[\frac{\theta^{1-\nu}}{\Gamma(\nu)} t_1 \right]^{n+1} M_1^{n+1} h.$$

Taking limit on recent equation as n approaches ∞ , we obtain $\|B_{1n}(t)\| \rightarrow 0$. In the same way, we can show that $\|B_{in}(t)\| \rightarrow 0, i = 2, 3, 4$. This completes the proof. \square

In the following, we show that system (2) has a unique solution. We suppose that the system has another solution such as $S_{1h}(t), I_{1h}(t), S_{1m}(t),$ and $I_{1m}(t),$ then we have

$$S_h(t) - S_{1h}(t) = \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t (W_1(\tau, S_h) - W_1(\tau, S_{1h})) d\tau.$$

By taking the norm from this equation, we obtain

$$\|S_h(t) - S_{1h}(t)\| = \frac{\theta^{1-\nu}}{\Gamma(\nu)} \int_0^t \|W_1(\tau, S_h) - W_1(\tau, S_{1h})\| d\tau.$$

It follows from Lipschitz condition (4) that

$$\|S_h(t) - S_{1h}(t)\| \leq \frac{\theta^{1-\nu}}{\Gamma(\nu)} M_1 t \|S_h(t) - S_{1h}(t)\|.$$

Then

$$\|S_h(t) - S_{1h}(t)\| \left(1 - \frac{\theta^{1-\nu}}{\Gamma(\nu)} M_1 t\right) \leq 0. \tag{7}$$

Theorem 4.3 *The solution of the transmission model of Zika virus is unique if the following condition holds:*

$$1 - \frac{\theta^{1-\nu}}{\Gamma(\nu)} M_1 t > 0.$$

Proof Suppose that condition (7) holds

$$\|S_h(t) - S_{1h}(t)\| \left(1 - \frac{\theta^{1-\nu}}{\Gamma(\nu)} M_1 t\right) \leq 0.$$

Then $\|S_h(t) - S_{1h}(t)\| = 0$. So, we obtain $S_h(t) = S_{1h}(t)$. Similarly, we can show the same equality for I_h, S_m, I_m . □

5 Numerical results

Using the fractional Euler method for Caputo derivative, we present the approximate solutions for the transmission model of Zika virus [50]. We present simulations for investigating the dynamics of the system.

5.1 Numerical method

We consider system (2) in the compact form as follows:

$$\theta^{\nu-1} {}^C D_t^\nu u(t) = p(t, u(t)), \quad u(0) = u_0, 0 \leq t \leq T < \infty, \tag{8}$$

where $u = (S_h, I_h, S_m, I_m) \in R_+^4$, $u_0 = (S_{0h}, I_{0h}, S_{0m}, I_{0m})$ is the initial vector, and $p(t) \in R$ is a continuous vector function satisfying the Lipschitz condition

$$\|p(u_1(t)) - p(u_2(t))\| \leq r \|u_1(t) - u_2(t)\|, \quad r > 0.$$

Applying the fractional integral operator corresponding Caputo derivative to equation (8), we obtain

$$u(t) = \theta^{1-\nu} [u_0 + I^\nu p(u(t))], \quad 0 \leq t \leq T < \infty.$$

Set $q = \frac{T-0}{N}$ and $t_n = nq$, where $t \in [0, T]$ and N is a natural number and $n = 0, 1, 2, \dots, N$. Let u_n be the approximation of $u(t)$ at $t = t_n$. Using the fractional Euler method [50], we get

$$u_{n+1} = \theta^{1-\nu} \left[u_0 + \frac{q^\nu}{\Gamma(\nu + 1)} \sum_{j=0}^n z_{n+1,j} p(t_j, u_j) \right], \quad j = 0, 1, 2, \dots, N - 1,$$

where

$$z_{n+1,j} = (n + 1 - j)^\nu - (n - j)^\nu, \quad j = 0, 1, 2, \dots, n.$$

The stability analysis of the obtained scheme has been proved in Theorem (3.1) in [50].

Thus, the solution of system (2) is written as follows:

$$\begin{aligned} S_{(n+1)h} &= \theta^{1-\nu} \left[S_{0h} + \frac{q^\nu}{\Gamma(\nu + 1)} \sum_{j=0}^n z_{n+1,j} g_1(t_j, u_j) \right], \\ I_{(n+1)h} &= \theta^{1-\nu} \left[I_{0h} + \frac{q^\nu}{\Gamma(\nu + 1)} \sum_{j=0}^n z_{n+1,j} g_2(t_j, u_j) \right], \\ S_{(n+1)m} &= \theta^{1-\nu} \left[S_{0m} + \frac{q^\nu}{\Gamma(\nu + 1)} \sum_{j=0}^n z_{n+1,j} g_3(t_j, u_j) \right], \\ I_{(n+1)m} &= \theta^{1-\nu} \left[I_{0m} + \frac{q^\nu}{\Gamma(\nu + 1)} \sum_{j=0}^n z_{n+1,j} g_4(t_j, u_j) \right], \end{aligned}$$

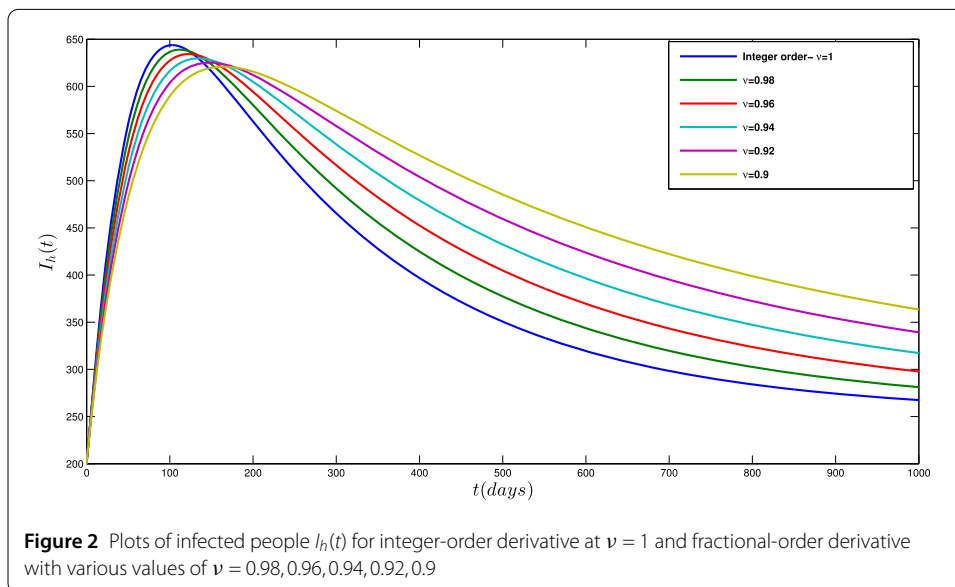
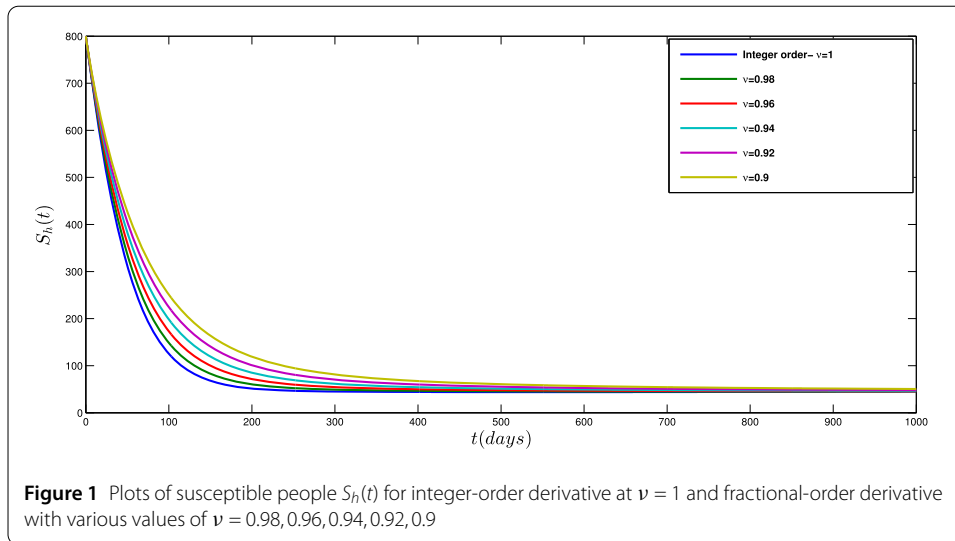
where $z_{n+1,j} = (n + 1 - j)^\nu - (n - j)^\nu$, $g_1(t, u(t)) = \Lambda_h - \beta_1 S_h(t) I_h(t) - \beta_2 I_m(t)(t) S_h(t) - k_1 S_h(t)$, $g_2(t, u(t)) = \beta_1 S_h(t) I_h(t) + \beta_2 I_m(t)(t) S_h(t) - k_1 I_h(t)$, $g_3(t, u(t)) = \Lambda_m - \mu S_m(t) I_h(t) - k_2 S_m(t)$, $g_4(t, u(t)) = \mu S_m(t) I_h(t) - k_2 I_m(t)$.

5.2 Simulation

In this section, using numerical results, we investigate the behavior of the answers of the transmission model of Zika virus obtained from system (2). The numerical values of the model parameters are considered as $\Lambda_h = 1.2$, $\Lambda_m = 0.3$, $k_1 = 0.004$, $k_2 = 0.0014$, $\beta_1 = 0.125 \times 10^{-4}$, $\beta_2 = 0.4 \times 10^{-4}$, $\mu = 0.475 \times 10^{-5}$, and we take its modification parameter as $\theta = 0.99$. Also, the initial values are considered as $S_h(0) = 800$, $I_h(0) = 200$, $S_m = 600$, $I_m = 300$.

Figure 1 shows susceptible people S_h and Fig. 2 shows infected people I_h for the integer-order derivative $\nu = 1$ and fractional-order derivative $\nu = 0.98, 0.96, 0.94, 0.92, .09$. As Fig. 1 shows, the behavior of S_h in both types of integer-order and fractional-order derivative is the same and decreasing, that is, over time, all healthy people are exposed to the disease, but the obtained numerical values are different, and as the derivative order decreases, the resulting numerical value increases.

In Fig. 2, you can see that the behavior of I_h is the same in both derivatives, and the resulting numerical values are different. As the derivative order decreases, the resulting numerical value for I_h increases, and this difference in the obtained value is significant

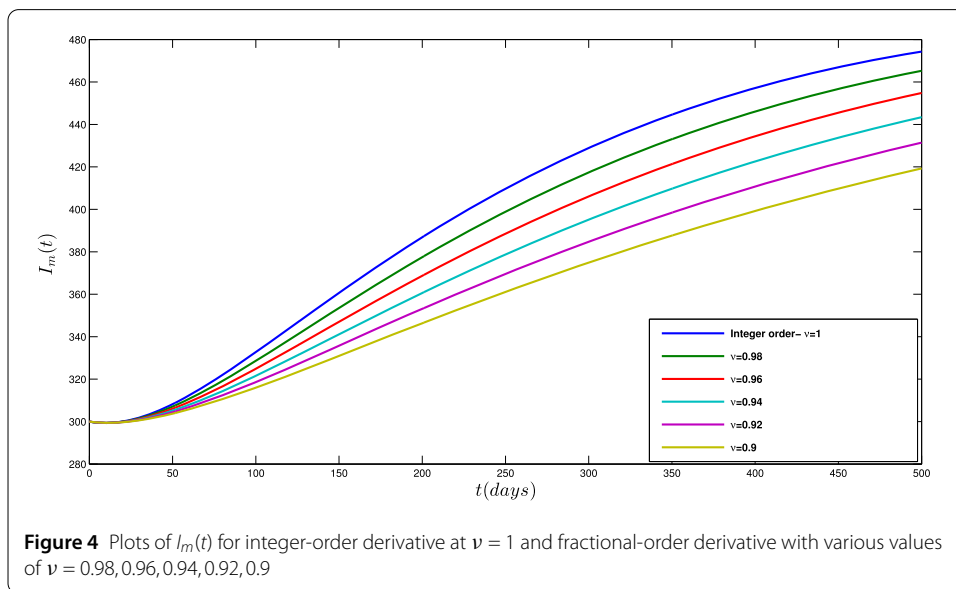
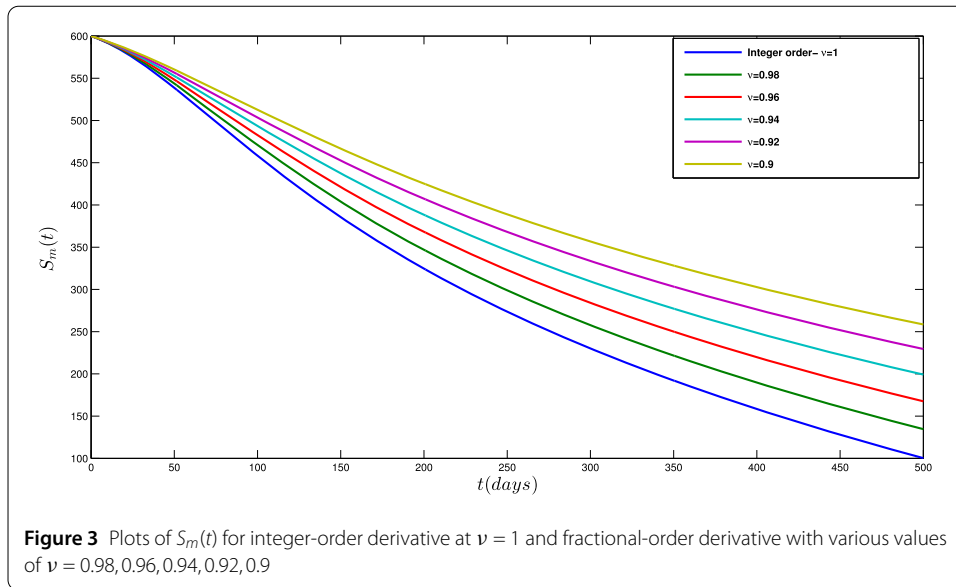


over time. Figure 2 also shows that I_h passes the peak in the first 100 days and the number of infected people gradually decreases and tends to the equilibrium point.

Figures 3 and 4 show susceptible mosquitoes S_m and infected mosquitoes I_m , respectively. In these diagrams, you can see that the behavior of the functions is the same in both derivatives and the resulting numerical values are different. These figures also show that over time the population of healthy mosquitoes decreases and they are more exposed to the disease, while the number of infected mosquitoes increases.

6 Conclusion

In this paper, a mathematical model for the transmission of Zika virus between humans and mosquitoes is presented using the Caputo fractional-order derivative. The region of the feasibility of system (2), the equilibrium points, and the reproduction number have been determined, and the stability of the equilibrium point E^0 has been checked. Using a fixed point theory, the existence of a unique solution for model (2) has been proven. In



the numerical section, the answers of system (2) are calculated using the Euler method, and the results are compared for the integer-order model and the fractional-order model in numerical results. The results show that the behavior of the obtained functions in both types of derivatives is the same, but the resulting numerical values are different, especially the difference in values increases over time.

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Authors' contributions

The authors declare that the study was realized in collaboration with equal responsibility. All authors read and approved the final manuscript.

Author details

¹Institute of Research and Development, Duy Tan University, Da Nang 550000, Vietnam. ²Faculty of Natural Sciences, Duy Tan University, Da Nang 550000, Vietnam. ³Department of Medical Research, China Medical University Hospital, China Medical University, Taichung, Taiwan. ⁴Department of Mathematics, Miandoab Branch, Islamic Azad University, Miandoab, Iran. ⁵Department of Electrical Engineering, University of Bojnord, P.O. Box, 94531-1339, Bojnord, Iran. ⁶Department of Mathematics, Near East University TRNC, Mersin 10, Turkey.

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References

1. Chen, T.M., Rui, J., Wang, Q.P., Zhao, Z.Y., Cui, J.A., Yin, L.: A mathematical model for simulating the phase-based transmissibility of a novel coronavirus. *Infect. Dis. Poverty* **9**, 24 (2020). <https://doi.org/10.1186/s40249-020-00640-3>
2. Dighe, A., Jombart, T., Van Kerkhove, M., Ferguson, N.: A mathematical model of the transmission of middle east respiratory syndrome coronavirus in dromedary camels (*Camelus dromedarius*). *Int. J. Infect. Dis.* **79**(S1), 03.002 (2019). <https://doi.org/10.1016/j.ijid.2018.11.023>
3. Tan, X., Yuan, L., Zhou, J., Zheng, Y., Yang, F.: Modeling the initial transmission dynamics of influenza A H1N1 in Guangdong province, China. *Int. J. Infect. Dis.* **17**, 479–484 (2013)
4. Alizadeh, S., Baleanu, D., Rezapour, S.: Analyzing transient response of the parallel RCL circuit by using the Caputo–Fabrizio fractional derivative. *Adv. Differ. Equ.* **2020**, 55 (2020). <https://doi.org/10.1186/s13662-020-2527-0>
5. Aydogan, S.M., Baleanu, D., Mousalou, A., Rezapour, S.: On high order fractional integro-differential equations including the Caputo–Fabrizio derivative. *Bound. Value Probl.* **2018**, 90 (2018)
6. Rezapour, S., Samei, M.E.: On the existence of solutions for a multi-singular pointwise defined fractional q-integro-differential equation. *Bound. Value Probl.* **2020**, 38 (2020). <https://doi.org/10.1186/s13661-020-01342-3>
7. Samei, M.E., Rezapour, S.: On a system of fractional q-differential inclusions via sum of two multi-term functions on a time scale. *Bound. Value Probl.* **2020**, 135 (2020). <https://doi.org/10.1186/s13661-020-01433-1>
8. Baleanu, D., Mousalou, A., Rezapour, S.: On the existence of solutions for some infinite coefficient-symmetric Caputo–Fabrizio fractional integro-differential equations. *Bound. Value Probl.* **2017**(1), 145 (2017). <https://doi.org/10.1186/s13661-017-0867-9>
9. Baleanu, D., Etemad, S., Rezapour, S.: On a fractional hybrid integro-differential equation with mixed hybrid integral boundary value conditions by using three operators. *Alex. Eng. J.* (2020). <https://doi.org/10.1016/j.aej.2020.04.053>
10. Baleanu, D., Nazemi, Z., Rezapour, S.: Attractivity for a k-dimensional system of fractional functional differential equations and global attractivity for a k-dimensional system of nonlinear fractional differential equations. *J. Inequal. Appl.* **2014**, 31 (2014). <https://doi.org/10.1186/1029-242X-2014-31>
11. Ghorbanian, R., Hedayati, V., Postolache, M., Rezapour, S.: Attractivity for a k-dimensional system of fractional functional differential equations and global attractivity for a k-dimensional system of nonlinear fractional differential equations. *J. Inequal. Appl.* **2014**, 319 (2014). <https://doi.org/10.1186/1029-242X-2014-319>
12. Agarwal, R.P., Baleanu, D., Hedayati, V., Rezapour, S.: Two fractional derivative inclusion problems via integral boundary conditions. *Appl. Math. Comput.* **257**, 205–212 (2015). <https://doi.org/10.1016/j.amc.2014.10.082>
13. Alsaedi, A., Baleanu, D., Etemad, S., Rezapour, S.: On coupled systems of time-fractional differential problems by using a new fractional derivative. *J. Funct. Spaces* **2016**, Article ID 4626940 (2016). <https://doi.org/10.1155/2016/4626940>
14. Baleanu, D., Hedayati, V., Rezapour, S.: On two fractional differential inclusions. *SpringerPlus* **5**, 882 (2016). <https://doi.org/10.1186/s40064-016-2564-z>
15. Aydogan, S.M., Nazemi, Z., Rezapour, S.: Positive solutions for a sum-type singular fractional integro-differential equation with m-point boundary conditions. *Sci. Bull. "Politeh." Univ. Buchar., Ser. A, Appl. Math. Phys.* **79**(1), 89–98 (2017)
16. Etemad, S., Rezapour, S., Samei, M.E.: On a fractional Caputo–Hadamard inclusion problem with sum boundary value conditions by using approximate endpoint property. *Math. Methods Appl. Sci.* (2020). <https://doi.org/10.1002/mma.6644>
17. Aydogan, S.M., Baleanu, D., Aguilar, J.F.G., Rezapour, S., Samei, M.E.: Approximate endpoint solutions for a class of fractional q-differential inclusions. *Fractals* **26**(8), 1–18 (2020). <https://doi.org/10.1142/S0218348X20400290>
18. Shiri, B., Wu, G.C., Baleanu, D.: Collocation methods for terminal value problems of tempered fractional differential equations. *Appl. Numer. Math.* **156**, 385–395 (2020). <https://doi.org/10.1016/j.apnum.2020.05.007>

19. Baleanu, D., Shiri, B.: Collocation methods for fractional differential equations involving non-singular kernel. *Chaos Solitons Fractals* **116**, 136–145 (2018)
20. Shiri, B., Baleanu, D.: System of fractional differential algebraic equations with applications. *Chaos Solitons Fractals* **120**, 203–212 (2019). <https://doi.org/10.1016/j.chaos.2019.01.028>
21. Ma, C.Y., Shiri, B., Wu, G.C., Baleanu, D.: New signal smoothing equations with short memory and variable order. *Optik* **218**, 164507 (2020). <https://doi.org/10.1016/j.ijleo.2020.164507>
22. Kumar, D., Singh, J., Baleanu, D.: On the analysis of vibration equation involving a fractional derivative with Mittag-Leffler law. *Math. Methods Appl. Sci.* **43**(1), 443–457 (2019)
23. Kumar, D., Singh, J., Tanwar, K., Baleanu, D.: A new fractional exothermic reactions model having constant heat source in porous media with power, exponential and Mittag-Leffler laws. *Int. J. Heat Mass Transf.* **138**, 1222–1227 (2019). <https://doi.org/10.1016/j.ijheatmasstransfer.2019.04.094>
24. Singh, J.: Analysis of fractional blood alcohol model with composite fractional derivative. *Chaos Solitons Fractals* **140**, 110127 (2020). <https://doi.org/10.1016/j.chaos.2020.110127>
25. Baleanu, D., Aydogan, S.M., Mohammadi, H., Rezapour, S.: On modelling of epidemic childhood diseases with the Caputo–Fabrizio derivative by using the Laplace Adomian decomposition method. *Alex. Eng. J.* (2020). <https://doi.org/10.1016/j.aej.2020.05.007>
26. Baleanu, D., Rezapour, S., Mohammadi, H.: Some existence results on nonlinear fractional differential equations. *Philos. Trans. R. Soc. Lond. A* **2013**, 371 (2013). <https://doi.org/10.1098/rsta.2012.0144>
27. Tuan, N.H., Mohammadi, H., Rezapour, S.: A mathematical model for Covid-19 transmission by using the Caputo fractional derivative. *Chaos Solitons Fractals* **140**, 110107 (2020). <https://doi.org/10.1016/j.chaos.2020.110107>
28. Baleanu, D., Jajarmi, A., Mohammadi, H., Rezapour, S.: Analysis of the human liver model with Caputo–Fabrizio fractional derivative. *Chaos Solitons Fractals* **134**, 109705 (2020)
29. Baleanu, D., Mohammadi, H., Rezapour, S.: Analysis of the model of HIV-1 infection of CD4+ T-cell with a new approach of fractional derivative. *Adv. Differ. Equ.* **2020**, 71 (2020)
30. Baleanu, D., Rezapour, S., Saberpour, Z.: On fractional integro-differential inclusions via the extended fractional Caputo–Fabrizio derivation. *Bound. Value Probl.* **2019**, 79 (2019)
31. Baleanu, D., Etemad, S., Rezapour, S.: A hybrid Caputo fractional modeling for thermostat with hybrid boundary value conditions. *Bound. Value Probl.* **2020**, 64 (2020). <https://doi.org/10.1186/s13661-020-01361-0>
32. Dokuyucu, M.A., Celik, E., Bulut, H., Baskonus, H.M.: Cancer treatment model with the Caputo–Fabrizio fractional derivative. *Eur. Phys. J. Plus* **133**, 92 (2018)
33. Khan, M.A., Hammouch, Z., Baleanu, D.: Modeling the dynamics of hepatitis E via the Caputo–Fabrizio derivative. *Math. Model. Nat. Phenom.* **14**(3), 311 (2019)
34. Ucar, E., Ozdemir, N., Altun, E.: Fractional order model of immune cells influenced by cancer cells. *Math. Model. Nat. Phenom.* **14**(3), 308 (2019)
35. Ullah, S., Khan, M.A., Farooq, M., Hammouch, Z., Baleanu, D.: A fractional model for the dynamics of tuberculosis infection using Caputo–Fabrizio derivative. *Discrete Contin. Dyn. Syst.* **13**(3), 975–993 (2020)
36. Singh, J., Kumar, D., Baleanu, D.: A new analysis of fractional fish farm model associated with Mittag-Leffler type kernel. *Int. J. Biomath.* **13**(2), 2050010 (2020)
37. Goswami, A., Singh, J., Kumar, D., Sunshila: An efficient analytical approach for fractional equal width equations describing hydro-magnetic waves in cold plasma. *Phys. A, Stat. Mech. Appl.* **524**, 563–575 (2019)
38. Srivastava, H.M., Dubey, V.P., Kumar, R., Singh, J., Kumar, D., Baleanu, D.: An efficient computational approach for a fractional-order biological population model with carrying capacity. *Chaos Solitons Fractals* **138**, 109880 (2020). <https://doi.org/10.1016/j.chaos.2020.109880>
39. Singh, J., Kilicmen, A., Kumar, D., Swroop, R., Ali, F.M.: Numerical study for fractional model of nonlinear predator–prey biological population dynamical system. *Therm. Sci.* **23**(6), 2017–2025 (2019)
40. Khalid, M., Samikhan, F.: Stability analysis of deterministic mathematical model for Zika virus. *Br. J. Math. Comput. Sci.* **19**(4), 1–10 (2016). <https://doi.org/10.9734/BJMCS/2016/29834>
41. Kibona, I.E., Yang, C.H.: SIR model of spread of Zika virus infections: Zikv linked to microcephaly simulations. *Health* **9**(8), 1190–1210 (2017). <https://doi.org/10.4236/health.2017.98086>
42. Maysaroh, A., Waluya, S.B., Wuryanto: Analisis dan simulasi model matematika penyakit Zika dengan satu serotip virus Zika. *UNNES J. Math.* **8**(1), 56–71 (2019)
43. Alkahtani, B.S.T., Atangana, A., Koca, I.: Novel analysis of the fractional Zika model using the Adams type predictor-corrector rule for non-singular and non-local fractional operators. *J. Nonlinear Sci. Appl.* **10**, 3191–3200 (2017). <https://doi.org/10.22436/jnsa.010.06.32>
44. Samko, S.G., Kilbas, A.A., Marichev, O.I.: *Fractional Integrals and Derivatives: Theory and Applications*. Gordon & Breach, Switzerland (1993)
45. Caputo, M., Fabrizio, M.: A new definition of fractional derivative without singular kernel. *Prog. Fract. Differ. Appl.* **1**(2), 73–85 (2015)
46. Losada, J., Nieto, J.J.: Properties of the new fractional derivative without singular kernel. *Prog. Fract. Differ. Appl.* **1**(2), 87–92 (2015)
47. Ullah, M.Z., Alzahrani, A.K., Baleanu, D.: An efficient numerical technique for a new fractional tuberculosis model with nonsingular derivative operator. *J. Taibah Univ. Sci.* **13**(1), 1147–1157 (2019)
48. Gomez-Aguilar, J.F., Rosales-Garcia, J.J., Bernal-Alvarado, J.J., Cordova-Fraga, T., Guzman-Cabrera, R.: Fractional mechanical oscillators. *Rev. Mex. Fis.* **58**, 348–352 (2012)
49. Van den Driessche, P., Watmough, J.: Reproduction numbers and subthreshold endemic equilibria for compartmental models of disease transmission. *Math. Biosci.* **180**, 29–48 (2002)
50. Li, C., Zeng, F.: The finite difference methods for fractional ordinary differential equations. *Numer. Funct. Anal. Optim.* **34**(2), 149–179 (2013)