#### FRACTIONAL TRANSMISSION MODEL FOR ZIKA VIRUS

Faith O. Oderhohwo<sup>\*1</sup>, Newton I. Okposo<sup>2</sup>, Augustine O. Atonuje<sup>3</sup>

<sup>1,2,3</sup> Department of Mathematics, Delta State University, PMB 1, Abraka, Delta State, Nigeria

Correspondence: Faith O. Oderhohwo, faiththemore@yahoo.com

#### Abstract

In this work, we studied the dynamics of a Zika virus model within the framework of the Caputo fractional derivative. Using a fixedpoint approach, we establish conditions for which the considered fractional model admits a unique system of solutions. The two-step Adams-Bashforth numerical scheme incorporating the fractional order parameter index  $\sigma$  is then used to furnish numerical simulations demonstrating the behaviour of the model state variables with respect to distinct values of the fractional order parameter index.

As the value of  $\sigma$  increases from 0.7 to 1, there is decrease in the number of susceptible individuals and then a gradual increase after some time t, until it steadies at equilibrium.

It was also observed that as the value of the fractional order parameter increases from 0.7 to 1 the number of exposed and infected individual decreases while the number of recovered individual increases after some time. Furthermore as the value of susceptible and exposed vector decreases, the number of infected vector increases and then decreases after some time.

**Keywords:** Fixed-point; Numerical scheme; Fractional derivatives; Fractional order parameter; Zika virus.

### Introduction

Zika virus was named after the Zika forest in Uganda where it was first discovered over 70 years ago by some researcher who were working on YFVD and was first isolated in a rhesus monkey. The virus belongs to the genus Flavi virus family with about 53 different species and has no specified antiviral drug or vaccine (Dick et al., 1952). Nigeria discovered and reported her first human cases in year 1954. Zika virus disease is caused by bites of infected female Aedes mosquitoes "Aedes aegypti "which are phylogenetically related to the ones that cause mosquito borne flavi viruses. This virus can be found in Blood, Urine, Saliva, semen, breast milk.

It is well documented that the transmission of Zika virus is possible via three main route, namely, between human to human via sexual interaction, transplacental transmission between an infected mother and the new born child, unscreened blood transfusion etc, human to vector and vector to human. This allows us to take into account transmissions arising from both human-to-human interactions as well as interactions between vector and human.

Prevention of Zika virus involves deliberate measures which include prevention of arbo viral infection by using mosquito treated nets, wearing long gear covering body, insecticides, repellents, or bird nets. Efforts towards having a vaccine to prevent Zika virus are currently ongoing (Barrett, 2018).

Several authors have studied different mathematical models describing the spread of Zika virus within the framework of systems of ordinary differential equations within the framework of classical (or integer-order) and fractional derivatives, however, models with integer-order derivatives do not adequately account for hereditary and memory effects associated with many biological processes (Rezapour et. al., 2020), the fractional order operators

In the present section, we present some fundamental definitions and properties on fractional differential and integral operators related to the Caputo type. incorporate hereditary properties and provides good description of the memory effects associated with many physical systems.

In this article, we employ a SEIR-type epidemic modeling framework comprising of two sub-populations, namely, human and mosquito vector sub-populations to investigate the dynamics of a fractional Zika virus model with seven epidemiological compartments consisting of susceptible, exposed, infectious and recovered humans as well as susceptible, exposed and infectious mosquitoes within the framework of the Caputo fractional derivative. For each of the mentioned subpopulations, a compartmental model is constructed to simulate Zika virus transmission while an interaction between both sub-populations occurs through mosquito bites. We incorporate a linear incidence term for both human to human and vector to human transmissions.

#### 2 Preliminaries

**Definition 1.** (Podlubny, 1999; Caputo, 1967) The Riemann-Liouville fractional integral of order  $\sigma$  of a function  $g \in C_{\mu,\mu} \ge$ - 1 is defined as

$$I_{t}^{\sigma}[g(t)] = \begin{cases} g(t), & \sigma = 0, t > 0, \\ \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t - \tau)^{\sigma - 1} g(\tau) d\tau, & \sigma > 0, t > 0, \end{cases}$$
(2.1)

where  $0 < \sigma < 1$ .

**Definition 2.** (Podlubny, 1999; Caputo, 1967) The Caputo fractional derivative of

order  $\sigma$  of a function  $g \in C^m_{-1}, m \in \mathbb{N} \cup \{0\}$ is defined as

$${}_{0}^{C}D_{t}^{\sigma}[g(t)] = \begin{cases} g^{(m)}(t) \coloneqq \frac{d^{m}g(t)}{dt^{m}}, & \sigma = m, \\ \frac{1}{\Gamma(m-\sigma)} \int_{0}^{t} g^{(m)}(t)(t-\tau)^{m-\sigma-1} d\tau, m-1 < \sigma < m. \end{cases}$$
(2.2)

**Lemma 1.** (Atangana and Owolabi, 2018) Let  $\sigma \in (0,1)$  and  $\mathcal{H} \in C([0,T], \mathbb{R}_+)$  be a nonlinear function. Then the fractional IVP in Caputo derivatives

$$\begin{cases} {}^{\mathbb{C}}_{0}D^{\sigma}_{t}\theta(t) = \mathcal{H}(t,\theta(t)), t \in [0,T], \\ \theta(0) = \theta_{0}, \end{cases}$$
(2.3)

has a unique solution given as

$$\theta(t) = \theta_0 + \frac{1}{\Gamma(\sigma)} \int_0^t (t-s)^{\sigma-1} \mathcal{H}(s,\theta(s)) ds.$$
(2.4)

#### 3

#### **MODEL FORMULATION**

In the present consideration, we consider the fractional version of the model proposed by Ali et. al. (2022). the total human population denoted by  $N_h(t)$  is subdivided into four independent epidemiological compartments, namely, susceptible humans  $S_h(t),$ exposed humans  $E_h(t)$ , infected humans  $I_h(t)$ , and recovered humans  $R_h(t)$ , such that  $N_h(t) = S_h(t) + E_h(t) + I_h(t) +$ 

 $R_h(t)$  while the entire vector population denoted by  $N_v(t)$  is subdivided into three independent epidemiological compartments, namely, susceptible vectors  $S_v(t)$ , exposed vectors  $E_v(t)$  and infected mosquito  $I_v(t)$  such that  $N_v(t) = S_v(t) + E_v(t) + I_v(t)$ .

Susceptible humans and mosquitoes are recruited into the susceptible compartments  $S_h$  and  $S_v$  at rates  $\Pi_h$  and  $\Pi_v$ , respectively. We represent by  $\lambda S_h = (\lambda_1 + \lambda_2)S_h$  the incidence rate of infection in the human population where  $\lambda_1 = \beta_h I_v$  is the rate at which susceptible individuals acquire infection due to effective contact with an infected vector and  $\lambda_2 = \rho \beta_h I_h$  is the rate at which susceptible individuals acquire infection due to sexual interaction with infected individuals. Here,  $\beta_h$  is the effective contact rate between susceptible humans and infected mosquitoes while  $\rho$  is a modification parameter that accounts for the relative infectiousness of individuals in the  $I_h$  relative to those in the  $I_v$ compartment. Similarly, we represent by  $\beta_{v}I_{h}S_{v}$  the incidence rate of the susceptible vector population where  $\beta_{\nu}$  denotes the transmission rate from infected humans to susceptible mosquito. The disease induced mortality rate is denoted by  $\delta$ . Natural mortality rates due for the human and vector subpopulations are denoted by  $\mu_h$ and  $\mu_{\nu}$  respectively. Lastly,  $\gamma$  and  $\tau$  are the natural and treatment rates. Following the above description for the interrelationship between compartments, we arrive at the following coupled system of nonlinear ordinary differential equations describing the Zika virus dynamics:

$$\begin{cases} \frac{S_h(t)}{dt} = \Pi_h - \beta_h S_h (I_v + \rho I_h) - \mu_h S_h \\ \frac{E_h(t)}{dt} = \beta_h S_h (I_v + \rho I_h) - (\mu_h + \chi) E_h \\ \frac{I_h(t)}{dt} = \chi E_h - (\mu_h + \gamma + \tau) I_h \\ \frac{R_h(t)}{dt} = \gamma I_h - \mu_h R_h \end{cases}$$
(3.1)  
$$\frac{S_v(t)}{dt} = \Pi_v - \beta_v S_v I_v - \mu_v S_v \\ \frac{E_v(t)}{dt} = \beta_v S_v I_v - (\mu_v + \delta) E_v \\ \frac{I_v(t)}{dt} = \delta E_v - \mu_v I_v, \end{cases}$$

With associated initial conditions

Sh(0) = Sh0, Eh(0) = Eh0, Ih(0) = Ih0, Rh(0) = Rh0, Sv(0) = Sv0, Ev(0) = Ev0, Iv(0) = Iv0.

Next, we extend the classical model (3.2) to a fractional model by incorporating the time-fractional derivative in place of the classical ordinary derivative for each equation in the system. We will consider the fractional model in the Caputo sense, namely,

$$\begin{cases} {}^{\mathbb{C}}_{0}D_{t}^{\sigma}S_{h}(t) = \Pi_{h} - \beta_{h}S_{h}(I_{v} + \rho I_{h}) - \mu_{h}S_{h}, \\ {}^{\mathbb{C}}_{0}D_{t}^{\sigma}E_{h}(t) = \beta_{h}S_{h}(I_{v} + \rho I_{h}) - (\mu_{h} + \chi)E_{h}, \\ {}^{\mathbb{C}}_{0}D_{t}^{\sigma}I_{h}(t) = \chi E_{h} - (\mu_{h} + \gamma + \tau)I_{h}, \\ {}^{\mathbb{C}}_{0}D_{t}^{\sigma}R_{h}(t) = \gamma I_{h} - \mu_{h}R_{h}, \\ {}^{\mathbb{C}}_{0}D_{t}^{\sigma}S_{v}(t) = \Pi_{v} - \beta_{v}S_{v}I_{h} - \mu_{h}S_{h}, \\ {}^{\mathbb{C}}_{0}D_{t}^{\sigma}E_{v}(t) = \beta_{v}S_{v}I_{h} - (\mu_{h} + \delta)E_{v}, \\ {}^{\mathbb{C}}_{0}D_{t}^{\sigma}I_{v}(t) = \delta E_{v} - \mu_{v}I_{v}, \end{cases}$$
(3.2)

with associated non-negative initial conditions

$$S_{h}(0) = S_{h}, E_{h}(0) = E_{h0}, I_{h}(0) = I_{h0}, R_{h}(0) = R_{h0},$$
  

$$S_{v}(0) = S_{v0}, E_{v}(0) = E_{v0}, I_{v}(0) = I_{v0}$$
(3.3)

 ${}_{0}^{c}D_{t}^{\sigma}$  represents the fractional differential operator in the Caputo sense with  $0 < \sigma \le 1$  being the fractional parameter index.

### 4 Existence and Uniqueness of Solutions

Under certain conditions, the existence and uniqueness of solutions for the considered model with respect to the Caputo fractional derivative can be investigated via a fixedpoint technique. For the sake of convenience in our subsequent investigations, we make the following notations for the right-hand terms appearing in (3.2).

$$\begin{cases} \mathbb{F}_{1}(t, S_{h}(t)) = \Pi_{h} - \beta_{h}S_{h}(I_{v} + \rho I_{h}) - \mu_{h}S_{h}, \\ \mathbb{F}_{2}(t, E_{h}(t)) = \beta_{h}E_{h}(I_{v} + \rho I_{h}) - (\mu_{h} + \chi)E_{h}, \\ \mathbb{F}_{3}(t, I_{h}(t)) = \chi E_{h} - (\mu_{h} + \gamma + \tau)I_{h}, \\ \mathbb{F}_{4}(t, R_{h}(t)) = \gamma I_{h} - \mu_{h}R_{h}, \\ \mathbb{F}_{5}(t, S_{v}(t)) = \Pi_{v} - \beta_{v}S_{v}I_{h} - \mu_{h}S_{h}, \\ \mathbb{F}_{6}(t, E_{v}(t)) = \beta_{v}E_{v}I_{h} - (\mu_{h} + \delta)E_{v}, \\ \mathbb{F}_{7}(t, I_{v}(t)) = \delta E_{v} - \mu_{v}I_{v}, \end{cases}$$
(4.1)

In view of Lemma 1, applying the the following equivalent system of Riemann-Liouville integral operator on fractional integral equations: both sides of each equation in (3.2) yields

$$\begin{cases} S_{h}(t) = S_{h}(0) + \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t-\tau)^{\sigma-1} \mathbb{F}_{1}(\tau, S_{h}\tau) d\tau, \\ E_{h}(t) = E_{h}(0) + \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t-\tau)^{\sigma-1} \mathbb{F}_{2}(\tau, E_{h}(\tau)) d\tau, \\ I_{h}(t) = I_{h}(0) + \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t-\tau)^{\sigma-1} \mathbb{F}_{3}(r, I_{h}(\tau)) d\tau, \\ R_{h}(t) = R_{h}(0) + \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t-\tau)^{\sigma-1} \mathbb{F}_{4}(\tau, R_{h}(r)) d\tau, \\ S_{v}(t) = S_{v}(0) + \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t-\tau)^{\sigma-1} \mathbb{F}_{5}(\tau, S_{v}(\tau)) d\tau, \\ E_{v}(t) = E_{v}(0) + \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t-\tau)^{\sigma-1} \mathbb{F}_{6}(\tau, E_{v}(\tau)) d\tau, \\ I_{v}(t) = I_{v}(0) + \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t-\tau)^{\sigma-1} \mathbb{F}_{7}(\tau, I_{v}(\tau)) d\tau, \end{cases}$$

Next, we show that the nonlinear functions  $\mathbb{F}_i$  (*i*= 1, 2, · · · 7) defined in (4.2) satisfy the Lipschitz condition. To this end, we first consider

$$\mathbb{F}_1(t, S_h(t)) = \Pi_h - \beta_h S_h(I_v + \rho I_h) - \mu_h S_h.$$

Let  $I_h$  and  $I_v$  be two bounded functions on  $t \in [0, T]$  such that  $||I_h(t)|| \leq \tau_1$  and  $||I_v(t)|| \leq \tau_2$ where  $||\cdot||$  denotes the maximum norm. Then for any two functions  $S_t^*(t)$  and  $S_t^{**}(t)$ , we have  $||\mathbb{F}_1(t, S_h^*(t)) - \mathbb{F}_1(t, S_t^{**}(t))|| = ||-\beta_h(I_v(t) + \rho I_h(t))(S_t^*(t) - S_h^{**}(t)) - \mu_h(S_h^*(t) - S_h^{**}(t))||$  $\leq [\beta_h(||I_v(t)|| + \rho ||I_h(t)||) + \mu_h]||S_h^*(t) - S_h^{**}(t)||$  $\leq [\beta_h(\tau_2 + \rho \tau_1) + \mu_h]||S_h^*(t) - S_h^{**}(t)||.$ 

This implies

$$\left\| \mathbb{F}_{1}(t, S_{h}^{*}(t)) - \mathbb{F}_{1}(t, S_{t}^{**}(t)) \right\| \leq \mathcal{K}_{1} \| S_{h}^{*}(t) - S_{h}^{**}(t) \|$$
(4.3)

where  $\mathcal{K}_1 = \beta_h(\tau_2 + \rho \tau_1) + \mu_h$ . It follows that,  $\mathbb{F}_1(t, S_h(t))$  satisfies the Lipschitz condition. Moreover, is a contraction if  $0 \leq \beta_h(\tau_2 + \rho \tau_1) + \mu_h < 1$  In line with the argument leading to (4.3), we can also find constants  $\mathcal{K}_2 \coloneqq \mu_h + \chi, \mathcal{K}_3 \coloneqq \mu_h + \gamma + \tau, \mathcal{K}_4 \coloneqq$ 

 $\mu_h, \mathcal{K}_5 \coloneqq \beta_v \tau_1 + \mu_v, \mathcal{K}_6 \coloneqq \delta + \mu_v, and \mathcal{K}_7 \coloneqq \mu_v$  such that the remaining functions  $\mathbb{F}_i (i=2, 3, ..., 7)$  satisfy

$$\begin{split} \left\| \mathbb{F}_{2}(t, E_{h}^{*}(t)) - \mathbb{F}_{2}(t, E_{h}^{**}(t)) \right\| &\leq \mathcal{K}_{2} \|E_{h}^{*}(t) - E_{h}^{**}(t)\|, \\ \left\| \mathbb{F}_{3}(t, I_{h}^{*}(t)) - \mathbb{F}_{3}(t, I_{h}^{**}(t)) \right\| &\leq \mathcal{K}_{3} \|I_{h}^{*}(t) - I_{h}^{**}(t)\|, \\ \left\| \mathbb{F}_{4}(t, R_{h}^{*}(t)) - \mathbb{F}_{4}(t, R_{h}^{**}(t)) \right\| &\leq \mathcal{K}_{4} \|R_{h}^{*}(t) - I_{h}^{**}(t)\|, \\ \left\| \mathbb{F}_{5}(t, S_{v}^{*}(t)) - \mathbb{F}_{5}(t, S_{v}^{**}(t)) \right\| &\leq \mathcal{K}_{5} \|S_{v}^{*}(t) - S_{v}^{**}(t)\|, \\ \left\| \mathbb{F}_{6}(t, E_{v}^{*}(t)) - \mathbb{F}_{6}(t, E_{v}^{**}(t)) \right\| &\leq \mathcal{K}_{6} \|E_{v}^{*}(t) - E_{v}^{**}(t)\|, \\ \left\| \mathbb{F}_{7}(t, I_{v}^{*}(t)) - \mathbb{F}_{7}(t, I_{v}^{**}(t)) \right\| &\leq \mathcal{K}_{7} \|I_{v}^{*}(t) - I_{v}^{**}(t)\|, \end{split}$$

Furthermore, these functions are contractions if  $0 \le \mathcal{K}_i < 1$  (*i*= 2.3. · · ·, 7).

Next, in view of (4.1) we rewrite the fractional Zika virus model (3.2) as

$$\begin{cases} {}^{\mathbb{C}}_{0}D_{t}^{\sigma}\mathbf{X}(t) = g(t,\mathbf{X}(t)), 0 < t < T < \infty, \\ \mathbf{X}(0) = \mathbf{X}_{0}, \end{cases}$$
(4.4)

Where  $\mathbf{X}(t)$ ,  $\mathbf{X}(0)$  and  $g(t, \mathbf{X}(t))$  are vector functions defined as

$$\begin{cases} \mathbf{X}(t) = \begin{pmatrix} S_{h}(t) \\ E_{h}(t) \\ I_{h}(t) \\ R_{h}(t) \\ S_{v}(t) \\ E_{v}(t) \\ I_{v}(t) \end{pmatrix}, \quad \mathbf{X}(0) = \begin{pmatrix} S_{h}(0) \\ E_{h}(0) \\ I_{h}(0) \\ R_{h}(0) \\ S_{v}(0) \\ E_{v}(0) \\ I_{v}(0) \end{pmatrix}, \\ g(t, \mathbf{X}(t)) = \begin{pmatrix} \mathbb{F}_{1}(t, S_{h}(t)) \\ \mathbb{F}_{2}(t, E_{h}(t)) \\ \mathbb{F}_{3}(t, I_{h}(t)) \\ \mathbb{F}_{3}(t, I_{h}(t)) \\ \mathbb{F}_{5}(t, S_{v}(t)) \\ \mathbb{F}_{5}(t, S_{v}(t)) \\ \mathbb{F}_{6}(t, E_{v}(t)) \\ \mathbb{F}_{7}(t, I_{v}(t)) \end{pmatrix} = \begin{pmatrix} \Pi_{h} - \beta_{h} S_{h}(I_{v} + \rho I_{h}) - \mu_{h} S_{h} \\ \beta_{h} S_{h}(I_{v} + \rho I_{h}) - (\mu_{h} + \chi) E_{h} \\ \chi E_{h} - (\mu_{h} + \gamma + \tau) I_{h} \\ \gamma I_{h} - \mu_{h} R_{h} \\ \Pi_{v} - \beta_{v} S_{v} I_{h} - \mu_{v} S_{v} \\ \beta_{v} S_{v} I_{h} - (\mu_{v} + \delta) E_{v} \\ \delta E_{v} - \mu_{v} I_{v} \end{pmatrix} \end{cases}$$

Then the problem of investigating the existence and uniqueness of solutions for the considered fractional Zika virus model (3.2) is equivalent to that of investigating the existence and uniqueness of solutions

to the fractional IVP (4.4). In this direction, we establish the following theorem which is adapted from (Lyons, 2017).

**Theorem 4.1** Assume that the nonlinear functions  $\mathbb{F}_i(t)(i=1, 2, \dots, 7)$  satisfy the Lipschitz condition and that there exist constants  $M_i > 0$   $(i=1, 2, \dots, 7)$  such that  $||\mathbb{F}_i(t, \cdot)|| \le M_i$  hold on the rectangle  $\mathbb{B} = \{(t, X): |t| \le a, |X - X_0| \le b\}$  containing the point (0, X0). Then the

fractional IVP (4.4) admits a unique solution on the interval  $\mathcal{J} \coloneqq \{t: |t| \le h\}$  where  $h = \max\{h_1, h_2, \dots, h_r\} = \min\left\{a, \left(\frac{b}{M}\Gamma(\sigma+1)\right)^{\frac{1}{\sigma}}\right\}$  and  $a \coloneqq \max\{a_1, a_2, \dots, a_7\}, b \coloneqq$ 

 $\max\{b_1, b_2, \dots, b_7\}$  and  $M := \max\{M_1, M_2, \dots, M_7\}$  are positive constants.

**Proof:** In view of (4.2), we have the following equivalent Voltera-type integral equation:

$$\mathbf{X}(t) = \mathbf{X}_0 + \frac{1}{\Gamma(\sigma)} \int_0^t (t - \tau)^{\sigma - 1} g(\tau, \mathbf{X}(\tau)) d\tau.$$
(4.5)

**Existence of solutions:** We will consider only the interval [0, h] as a similar argument also holds in the interval [-h, 0]. The proof for the existence of solutions is established by constructing a sequence  $\{X_k(t)\}\ (k = 1,2,3,...\)$ of successive approximations (Picard's iterates)

$$\mathbf{X}_{0}(t) = \mathbf{X}_{0}, \mathbf{X}_{k+1}(t) = \mathbf{X}_{0} + \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t-\tau)^{\sigma-1} g(\tau, \mathbf{X}_{k}(\tau)) d\tau, k = 0, 1, 2, ..., t \epsilon[0, h], (4.6)$$
which converges uniformly to a function which is a solution of the integral equation (4.5) on [0, h]. This is accomplished in the following steps: properly defined for every  $t \epsilon[0, h]$ , and in

this interval the following inequality holds

$$\begin{cases} |X_{k}(t) - X_{0}| \leq \frac{Mh^{\sigma}}{\Gamma(\sigma+1)} \leq b \\ |X_{k+1}(t) - X_{k}(t)| \leq \frac{MK^{k}t^{(k+1)\sigma}}{\Gamma((k+1)\sigma+1)} \end{cases}$$

$$(4.7)$$

Note that  $\mathbf{X}_{k+1}(t)$  is well defined in the interval [0, h] if the point  $(t, \mathbf{X}_k(t))$  remains in the rectangle  $\mathbb{B}$ for every  $t \in [0, h]$ . Clearly, for k = 0,  $\mathbf{X}_0(t)$  is defined on [0, h] and satisfies (4.7) trivially on [0, h]. Now, assume that for  $k = n \ge 1$ ,  $\mathbf{X}_n$  is defined and satisfies (4.7) on [0, h], then the point  $(t, \mathbf{X}_n(t))$  remains in B for t in [0, h]. Moreover,  $g(t, \mathbf{X}_n(t))$  exists as a continuous function on [0, h]. Hence, by (4.6)  $\mathbf{X}_n+1(t)$  is defined on [0, h] and that concludes the induction argument. Furthermore, from (4.6), the induction hypothesis and (4.7) we have

$$|\mathbf{X}_{k+1}(t) - \mathbf{X}_0| \leq \frac{1}{\Gamma(\sigma)} \int_0^t (t-\tau)^{\sigma-1} |g(\tau, \mathbf{X}_k(\tau))| d\tau \leq \frac{Mt^{\sigma}}{\Gamma(\sigma+1)} \leq \frac{Mh^{\sigma}}{\Gamma(\sigma+1)} \leq b.$$

Hence, the property (4.7) is satisfied by  $X_{k+1}$  and the induction argument is complete. **STEP II:** In this step, we show via an induction argument that the inequality

$$|\mathbf{X}_{k+1}(t) - \mathbf{X}_{k}(t)| \le \frac{M}{\mathcal{K}} \frac{(\mathcal{K}t^{\sigma})^{k+1}}{\Gamma((k+1)\sigma+1)}.$$
(4.8)

Holds on [0, h]. For the case k = 0, we have

$$\begin{split} |\mathbf{X}_{k+1}(t) - \mathbf{X}_{k}(t)| &= \left|\frac{1}{\Gamma(\sigma)}\right| \int_{0}^{t} (t-\tau)^{\sigma-1} |g(\tau, \mathbf{X}_{k}(\tau))| d\tau \\ &\leq \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t-\tau)^{\sigma-1} |g(\tau, \mathbf{X}_{0}(\tau))| d\tau \leq \frac{Mt^{\sigma}}{\Gamma(\sigma+1)}. \end{split}$$

Assume that (4.8) is true for  $\mathcal{K} \geq 1$ , that is

$$|\mathbf{X}_{k+1}(t) - \mathbf{X}_{k}(t)| \le \frac{Mt^{\sigma}}{\Gamma(k_{\sigma} + 1)} t^{k\sigma}$$

on [0, h] where  $\mathcal{K}$  is the Lipschetz constant. Then

$$\begin{split} |\mathbf{X}_{k+1}(t) - \mathbf{X}_{k}(t)| \\ &\leq \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t-\tau)^{\sigma-1} |g(\tau, \mathbf{X}_{k}(\tau)) - g(\tau, \mathbf{X}_{k-1}(\tau))| d\tau \\ &\leq \frac{1}{\Gamma(\sigma)} \int_{0}^{t} (t-\tau)^{\sigma-1} |\mathbf{X}_{k}(\tau) - \mathbf{X}_{k-1}(\tau)| d\tau \\ &\leq \frac{M\mathcal{K}^{k}}{\Gamma(\sigma)\Gamma(k\sigma+1)} \int_{0}^{t} (t-\tau)^{\sigma-1} \tau^{k\sigma} d\tau. \end{split}$$

Using the substitution  $\tau = \omega t$ , we have

$$\begin{split} \frac{M\mathcal{K}^{k}}{\Gamma(\sigma)\Gamma(k\sigma+1)} &\int_{0}^{t} (t-\tau)^{\sigma-1}(\tau-0)^{k\sigma} d\tau \\ &= \frac{M\mathcal{K}^{k}t^{(k+1)\sigma}}{\Gamma(\sigma)\Gamma(k\sigma+1)} \int_{0}^{t} (t-\omega)^{\sigma-1}\omega^{k\sigma} d\tau \\ &= \frac{M\mathcal{K}^{k}t^{(k+1)\sigma}}{\Gamma(\sigma)\Gamma(k\sigma+1)} B(\sigma,k\sigma+1) \\ &= \frac{M\mathcal{K}^{k}t^{(k+1)\sigma}}{\Gamma(\sigma)\Gamma(k\sigma+1)} \frac{M\mathcal{K}^{k}t^{(k+1)\sigma}}{\Gamma((k+1)\sigma+1)} \\ &= \frac{M\mathcal{K}^{k}t^{(k+1)\sigma}}{\Gamma((k+1)\sigma+1)}. \end{split}$$

Here B(.,.) denotes the Beta function. Hence, we have

$$|\mathbf{X}_{k+1}(t) - \mathbf{X}_{k}(t)| \leq \frac{M\mathcal{K}^{k}t^{(k+1)\sigma}}{\Gamma((k+1)\sigma+1)}$$

Therefore, the inequality (4.8) is true for all *k*.

**STEP III.** We show that the sequence of functions  $\{\mathbf{X}_k(t)\}$  defined in (4.6) converge

uniformly on [0, h]. To this end, we observe that the sequence of successive approximations consists of the sequence of partial sum of the series

$$\mathbf{X}_{0}(t) + \sum_{k=0}^{\infty} [\mathbf{X}_{k+1}(t) - \mathbf{X}_{k}(t)].$$
(4.9)

To prove uniform convergence of  $\{\mathbf{X}_k(t)\}$ , it is enough to establish that the partial sums of the series (4.9) are uniformly convergent. From (4.8) we have

$$\sum_{k=0}^{\infty} |\mathbf{X}_{k+1}(t) - \mathbf{X}_k(t)| \le \frac{M}{\mathcal{K}} \sum_{k=0}^{\infty} \frac{(\mathcal{K}t^{\sigma})^{(k+1)}}{\Gamma((k+1)\sigma+1)}.$$
(4.10)

In other words, the terms in (4.9) are bounded in absolute value on the interval [0, h] by the terms in the positive series

$$\frac{M}{\mathcal{K}}\sum_{k=0}^{\infty}\frac{(\mathcal{K}t^{\sigma})^{k+1}}{\Gamma((k+1)\sigma+1)}$$

which converges to  $\frac{M}{\pi}E_{\sigma,1}(\mathcal{K}t^{\sigma})$ . Hence, by Weierstrass M-test, the series in (4.9) converges uniformly on [0, h]. Therefore, the sequence  $\{\mathbf{X}_k\}$  is uniformly convergent on [0, h] to continuous function say  $\mathbf{X}(t)$ .

**STEP IV.** In this step, we show that  $\mathbf{X}(t)$ is the solution of the integral equation

(4.5). Since the sequence 
$$\{\mathbf{X}_k\}$$
 is uniformly convergent to some continuous function  $\mathbf{X}(t)$ , then from (4.6) we have

sequence

the

Since

$$\begin{aligned} \mathbf{X}(t) &= \lim_{k \to \infty} \mathbf{X}_{k+1}(t) \\ &= \lim_{k \to \infty} \left[ X_0 + \frac{1}{\Gamma(\sigma)} \int_0^t (t - \tau)^{\sigma - 1} g(\tau, \mathbf{X}_k(\tau)) d\tau \right] \quad (4.11) \\ &= X_0 + \frac{1}{\Gamma(\sigma)} \int_0^t (t - \tau)^{\sigma - 1} g(\tau, \mathbf{X}(\tau)) d\tau \end{aligned}$$

Since  $\mathbf{X}(t)$  satisfies the equivalent integral equation, then it must satisfy the initial value problem (4.4).

Uniqueness: To complete the proof of the theorem, we establish uniqueness of solution to the fractional IVP (4.4) with Caputo derivative by claiming that there

exists another solution  $\widetilde{\mathbf{X}}(t) := (\widetilde{S}_h(t), \widetilde{E}_h(t))$  $\tilde{I}_h(t), \ \tilde{R}_h(t), \ \tilde{S}_v(t), \ \tilde{E}_v(t), \ \tilde{I}_v(t))^{\tau}.$ 

Now, consider the function  $\theta(t) = |\mathbf{X}(t)|$  –  $\widetilde{\mathbf{X}}(t)$ . Since both  $\mathbf{X}(t)$  and  $\widetilde{X}(t)$  are solutions to the IVP (4.4), we see that  $\boldsymbol{\theta}_0 = |\mathbf{X}_0 - \widetilde{\mathbf{X}}_0|$ = 0. Furthermore, from Theorem 4.1we have

$$\begin{aligned} \theta(t) &= \left| \mathbf{X}(t) - \widetilde{\mathbf{X}}(t) \right| \\ &= \left| \frac{1}{\Gamma(\sigma)} \int_0^t (t - \tau)^{\sigma - 1} g(\tau, \mathbf{X}(\tau)) - g(\tau, \widetilde{\mathbf{X}}(\tau)) d\tau \right| \\ &\leq \frac{1}{\Gamma(\sigma)} \int_0^t (t - \tau)^{\sigma - 1} \left| g(\tau, \mathbf{X}(\tau)) - g(\tau, \widetilde{\mathbf{X}}(\tau)) \right| d\tau \qquad (4.12) \\ &\leq \frac{\mathcal{K}}{\Gamma(\sigma)} \int_0^t (t - \tau)^{\sigma - 1} \theta(\tau) d\tau. \end{aligned}$$

Since  $\mathcal{K} > 0$  and  $\theta(t) \ge 0$ , the inequality in (4.12) satisfies all conditions of the Gronwall-type inequality (Denton and Vatsala, 2010). Thus, we have

$$\theta(t) \le \theta_0 E_{\sigma,1}(\mathcal{K}t^{\sigma}) = 0$$

This proves that  $X = \tilde{X}$  and hence, the solution of the fractional IVP (4.4) exists on [0, *h*] and is unique. Consequently, the fractional Zika virus model (3.2) in Caputo derivative admits unique solution.

#### 5 Numerical Schemes

We present corresponding FAB schemes for the fractional Zika virus models (4.1) with fractional derivative in the Caputo sense. We furnish graphical visualizations comparing the behaviours of each state variables of the fractional Zika virus models (4.1) for distinct values of the fractional order parameter  $\sigma$ . The simulation parameter values are taken as  $\Pi$  $h = 0.8, \Pi_V = 0.08, \beta_h = 0.007, \beta_V = 0.009, \mu_h =$  *Remark* 4.5. It is easy to see that if  $\sigma = 1$ , the proof Theorem 4.4 is similar to Picard's original proof for systems of ordinary differential equations with integer order derivatives.

0.0028,  $\mu_{\nu}$ = 0.071,  $\rho$ = 0.05, *X*= 0.7,  $\gamma$ = 0.05,  $\tau$ = 0.08,  $\delta$ = 0.5 (Bonyah et al, 2017) while the initial values used are  $S_{h0}$  = 100,  $E_{h0}$  = 10,  $I_{h0}$  = 30,  $R_{h0}$  = 20,  $S_{\nu0}$  = 10,  $E_{\nu0}$  = 50,  $I_{\nu0}$  = 10.

# 5.1 FAB scheme for model in Caputo derivative

Using the fundamental theorem of integral calculus, we obtain the following corresponding nonlinear fractional Volterrra-type integral equation

$$S_h(t) - S_h(0) = \frac{1}{\Gamma(\sigma)} \int_0^t (t - \tau)^{\sigma - 1} \mathbb{F}_1(\tau, S_h(\tau)) d\tau.$$
(5.1)

for the  $S_h$ -equation (4.5) in Caputo derivative. At  $t = t_{k+1}$  and  $t = t_k$ , k = 0, 1, 2, ..., (5.1) can be read as

$$S_h(t_{k+1}) - S_h(0) = \frac{1}{\Gamma(\sigma)} \int_0^{t_{k+1}} (t_{k+1} - t)^{\sigma - 1} \mathbb{F}_1(t, S_h(t)) dt$$

and

$$S_h(t_k) - S_h(0) = \frac{1}{\Gamma(\sigma)} \int_0^{t_k} (t_k - t)^{\sigma - 1} \mathbb{F}_1(t, S_h(t)) dt.$$

respectively. We easily see that,

$$S_h(t_{k+1}) - S_h(t_k) = X_{\sigma,1} - X_{\sigma,2},$$
 (5.2)

where:

$$X_{\sigma,1} = \frac{1}{\Gamma(\sigma)} \int_0^{t_{k+1}} (t_{k+1} - t)^{\sigma-1} \mathbb{F}_1(t, S_h(t)) dt,$$
(5.3)

and

$$X_{\sigma,2} = \frac{1}{\Gamma(\sigma)} \int_0^{t_k} (t_k - t)^{\sigma - 1} \mathbb{F}_1(t, S_h(t)) dt, \qquad (5,4)$$

Over the interval [ $t_k$ ,  $t_{k+1}$ ], the function  $F_1(t, S_h(t))$  can be approximated by the two-point Lagrange interpolation polynomial of the form

$$\mathbb{F}_{1}(t, S_{h}(t)) \simeq \frac{t - t_{k-1}}{t_{k} - t_{k-1}} \mathbb{F}_{1}(t_{k}, S_{h}(t_{k})) + \frac{t - t_{k}}{t_{k-1} - t_{k}} \mathbb{F}_{1}(t_{k-1}, S_{h}(t_{k-1}))$$
$$= \frac{t - t_{k-1}}{h} \mathbb{F}_{1}(t_{k}, S_{h}(t_{k})) - \frac{t - t_{k}}{h} \mathbb{F}_{1}(t_{k-1}, S_{h}(t_{k-1})),$$

Where  $h = t_k - t_{k-1}$  is the step-size. Substituting (5.5) into the first and second integrals in (5.3) yield

$$X_{\sigma,1} = \frac{\mathbb{F}_{1}(t_{k}, S_{h}(t_{k}))}{h\Gamma(\sigma)} \left[ \frac{2ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} \right] - \frac{\mathbb{F}_{1}(t_{k-1}, S_{h}(t_{k-1}))}{h\Gamma(\sigma)} \left[ \frac{ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} \right], (5.5)$$

And

$$X_{\sigma,2} = \frac{\mathbb{F}_1(t_k, S_h(t_k))}{h\Gamma(\sigma)} \left[ \frac{ht_k^{\sigma}}{\sigma} - \frac{t_k^{\sigma+1}}{\sigma+1} \right] - \frac{\mathbb{F}_1(t_k, S_h(t_{k-1}))}{h\Gamma(\sigma)} \frac{t_k^{\sigma+1}}{\sigma+1} , (5.6)$$

respectively, after some manipulations. By inserting (5.5) and (5.6) into (5.2), we obtain

$$S_{h}(t_{k+1}) = S_{h}(t_{k}) + \frac{\mathbb{F}_{1}(t_{k}, S_{h}(t_{k}))}{h\Gamma(\sigma)} \left[ \frac{2ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} - \frac{ht_{k}^{\sigma}}{\sigma} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \right] - \frac{\mathbb{F}_{1}(t_{k-1}, S_{h}(t_{k-1}))}{h\Gamma(\sigma)} \left[ \frac{ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \right]$$
(5.7)

as the final two-step FAB scheme for the  $S_h$ -equation (3.2) with Caputo derivative. In the same way, we can obtain a similar scheme for each of the remaining equations in (3.2). In particular, we have

$$\begin{split} E_{h}(t_{k+1}) &= E_{h}(t_{k}) + \frac{\mathbb{F}_{1}(t_{k}, E_{h}(t_{k}))}{h\Gamma(\sigma)} \bigg[ \frac{2ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} - \frac{ht_{k}^{\sigma}}{\sigma} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \bigg] \\ &- \frac{\mathbb{F}_{1}(t_{k-1}, E_{h}(t_{k-1}))}{h\Gamma(\sigma)} \bigg[ \frac{ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \bigg], \\ I_{h}(t_{k+1}) &= I_{h}(t_{k}) + \frac{\mathbb{F}_{1}(t_{k}, I_{h}(t_{k}))}{h\Gamma(\sigma)} \bigg[ \frac{2ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} - \frac{ht_{k}^{\sigma}}{\sigma} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \bigg] \\ &- \frac{\mathbb{F}_{1}(t_{k-1}, I_{h}(t_{k-1}))}{h\Gamma(\sigma)} \bigg[ \frac{ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \bigg], \end{split}$$

$$\begin{split} R_{h}(t_{k+1}) &= R_{h}(t_{k}) + \frac{\mathbb{F}_{1}(t_{k}, R_{h}(t_{k}))}{h\Gamma(\sigma)} \left[ \frac{2ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} - \frac{ht_{k}^{\sigma}}{\sigma} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \right] \\ &- \frac{\mathbb{F}_{1}(t_{k-1}, R_{h}(t_{k-1}))}{h\Gamma(\sigma)} \left[ \frac{ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \right], \\ S_{v}(t_{k+1}) &= S_{v}(t_{k}) + \frac{\mathbb{F}_{1}(t_{k}, S_{v}(t_{k}))}{h\Gamma(\sigma)} \left[ \frac{2ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} - \frac{ht_{k}^{\sigma}}{\sigma} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \right] \\ &- \frac{\mathbb{F}_{1}(t_{k-1}, S_{v}(t_{k-1}))}{h\Gamma(\sigma)} \left[ \frac{ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \right], \\ E_{v}(t_{k+1}) &= E_{v}(t_{k}) + \frac{\mathbb{F}_{1}(t_{k}, E_{v}(t_{k}))}{h\Gamma(\sigma)} \left[ \frac{2ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} + \frac{ht_{k}^{\sigma}}{\sigma} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \right] \\ &- \frac{\mathbb{F}_{1}(t_{k-1}, E_{v}(t_{k-1}))}{h\Gamma(\sigma)} \left[ \frac{ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} - \frac{ht_{k}^{\sigma}}{\sigma} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \right] \\ I_{v}(t_{k+1}) &= I_{v}(t_{k}) + \frac{\mathbb{F}_{1}(t_{k}, I_{v}(t_{k}))}{h\Gamma(\sigma)} \left[ \frac{2ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} + \frac{ht_{k}^{\sigma}}{\sigma} + \frac{t_{k}^{\sigma+1}}{\sigma+1} \right] \\ &- \frac{\mathbb{F}_{1}(t_{k-1}, I_{v}(t_{k-1}))}{h\Gamma(\sigma)} \left[ \frac{ht_{k+1}^{\sigma}}{\sigma} - \frac{t_{k+1}^{\sigma+1}}{\sigma+1} + \frac{ht_{k}^{\sigma+1}}{\sigma+1} \right]. \quad (5.8) \end{split}$$

as the two-step FAB scheme for the fractional Zika virus model (3.2) with Caputo derivative.

## 5.2 Simulations and discussion of results

Using the above two-step FAB schemes (5.8), we present graphical visualizations to demonstrate the behaviour of the approximate solutions to the fractional Zika virus model (3.2) with Caputo derivatives for each system variable. The plots in each of the graphs are with respect to distinct values of the fractional order parameter  $\sigma$ with  $\sigma$ = 1.0; 0.9; 0.8; 0.7. The time level up to 100 days and the step size used for evaluating the approximate solutions is h = 0.002. The graphs for susceptible individuals  $S_h(t)$ , exposed individuals  $E_h(t)$ , infected individuals  $I_h(t)$ , recovered individuals  $R_h(t)$ , susceptible

vectors  $S_{\nu}(t)$ , exposed vectors  $E_{\nu}(t)$  and infected vectors  $I_{\nu}(t)$  are presented in Figure 1-7, respectively. In each of the plots, it is observed that the magnitude of  $\sigma$ continuously affects the trend of each state variable for both the human and vector populations.

Figure 1- 4: plots shows the dynamics of individuals, the susceptible exposed individual. Infected individual and Recovered individual respectively using above scheme for the Caputo the derivatives. Figure 5-7 demonstrates the dynamics of susceptible vectors, exposed vectors and infected vectors for different values of  $\sigma$  in each of the plots.

Figure 1: As the value of  $\sigma$  increases from 0.7 to 1, there is decrease in the number of susceptible individuals and then a gradual increase after some time until it steadies at equilibrium

Figure 2 shows the behaviour of the exposed individual using the corresponding  $E_h(t)$ -schemes of (5.8) for the Caputo derivatives for distinct values of the fractional parameter

Figure 3: It is observed that as the fractional order parameter increases from 0.7 to 1 the number of infected individual decreases after some time

Figure 4 shows the behaviour of the recovered individual using the corresponding  $R_h(t)$ -schemes of (5.8) for the Caputo derivatives for distinct values of the fractional parameter

Using the corresponding  $S_{\nu}(t)$ -schemes in (5.8) for the Caputo operators, Figure 5 demonstrates the dynamics of susceptible vectors



Figure 6: The plots demonstrate the dynamics of the exposed vectors for the case of the  $E_{\nu}(t)$ -equation of (3.2) using the corresponding  $E_{\nu}(t)$ -schemes in (5.8) for the Caputo

we compare the dynamics of Zika infected vectors by presenting plots for the approximate  $I_{\nu}(t)$ -solution using the corresponding  $I_{\nu}(t)$ -schemes of (5.8) for the Caputo for different values of  $\sigma$ . It is observed that as the fractional order parameter increases from 0.7 to 1 the population of infected vectors decreases after some time

#### 6 Conclusions

We analyzed a fractional mathematical model for the transmission dynamics of Zika virus under the framework of singular kernels. The solution set of the classical model is shown to be non-negative and positively invariant. We then determine the equilibrium points of the model and the basic reproduction number is determined via the next generation matrix technique. Existence and uniqueness of solutions to the fractional model with respect to Caputo derivatives are established via a fixedpoint technique. Numerical investigations using the two-step Adams-Bashforth method for the fractional Zika virus model with respect to the Caputo fractional differential operators are then carried out



with the purpose of demonstrating the dynamics of each of the system variables for different values of the fractional order parameter. We made comparison of the obtained results for each compartment with respect to the Caputo derivative.

#### Acknowledgments

The authors express their gratitude to the reviewers for their helpful suggestions.

#### **Conflict of interest**

The authors declare no potential conflict of interests regarding the publication of this paper.

#### References

Agusto FB, Bewick S, Fagan WF. (2017). Mathematical model of Zika virus with vertical transmission. Infectious Disease modelling; 2: 244-267

- Ali A., Iqbal Q., Asamoah J. K. K., Islam S. (2022). Mathematical modeling for the transmission potential of Zika virus with optimal control strategies. Eur. Phys. J.Plus 137:146
- Atangana A, Owolwabi K. M. (2018) New numerical approach for fractional differential equations. Mathemat. Model Nat. Phenomena. 2018; 13.
- Bonyah E., Khan M. A., Okosun K. O., Islam S. (2017) A theoretical model for Zika virus transmission. PLoS ONE 12 (10) :e0185540.
- Barrett, A. D. (2018). Current status of zika vaccine development: Zika vaccines advance into clinical evaluation. npj Vaccines, 3(1):1-4.
- Chikaki, E. and Ishikawa, H. (2009). A dengue transmission model in Thailand considering sequential infections with all four serotypes. The Journal of Infection in Developing Countries, 3(09):711-722.
- Denton, Z., Vatsala, A. S. (2010) Fractional Integral Inequalities and Applications. Comput. Math. Appl. 59, 1087-1094.
- Dick, G., Kitchen, S., Haddow, A., et al. (1952). Zika virus (ii). Pathogenicity and physical properties. Transactions of the royal society of tropical medicine and hygiene, 46(5).
- Farman M, Ahmad A, Akg<sup>\*</sup>ul A, Saleem MU, Rizwan M, Ahmad MO. (2020). A mathematical analysis and simulation for Zika virus model with time fractional derivative. Math Meth Appl Sci.;1-12. <u>https://doi.org/10.1002/mma.6891</u>
- Lloyd, A. (2009). Sensitivity of modelbased epidemiological parameter estimation to model assumptions. In Mathematical and statistical estimation approaches in

epidemiology (pp. 123-141). Springer.

Owolabi KM. (2016) Numerical solution of diffusive HBV model in a fractional medium. Springer Plus.; 5: 1643.

https://doi.org/10.1186/s40064-016-3295-x

- Podlubny I. (1999). Fractional Differential Equations. San Diego,
- Rezapour S., Mohammad H., Jajarmi A. (2020). A new mathematical model for Zika virus transmission. Advances in Difference Equations (2020) 2020:589. <u>https://doi.org/10.1186/s13662-020-03044-7</u>.
- World Health Organization (2016-2017), <u>http://www.who.int/mediacentre/fact</u> <u>sheets/zika/en/</u>.
- World Health Organization (WHO) 2021, Infectious diseases, <u>https://www.who.int/topics/infectiou</u> <u>sdiseases/(Accessed 10-10-2022)</u>
- World Health Organization (WHO), A global brief on vector-borne diseases. "<u>https://apps.who.int/iris/bitstream/h</u> andle/10665/111008/WHODCOWH D2014.1eng.pdf?sequence=2014"
- World Health Organization (WHO), WHO statement on the first meeting of the International Health Regulations (2005) Emergency Committee on Zika virus and observed increase in neurological disorder and neonatal malformations, February 1, 2016.