

Frictional Order Host-Vector Model for Transmission of Dengue Fever

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Abstract: Main purpose of this paper is to formulate an epidemiological model for dengue fever transmission using fractional order derivatives. Due to memory effect property, fractional order derivative has a benefit over the classical integer order models. This model for transmission of dengue fever of the non-integer order initial value problem will be based on the well-known fractional order Caputo derivative. Here our focus is on the existence of non-negative solutions of the frictional order dengue fever transmission model, furthermore, equilibria of the model and local asymptotic stability of model equilibria is investigated. In the end fractional order transmission model for dengue fever without immunity is presented.

Keywords: Dengue fever, Caputo derivative, Existence of positive solution, Model equilibria, Asymptotic stability.

I. INTRODUCTION

Mathematical models have been formulated, investigated and studied by many authors in engineering, finance, economics, science and in particular mathematical-biology (Infectious disease) using the classical integer order system of differential equations has received a lot of concentration in the past several years [1-14]. On the other hand due to the effective nature of fractional derivatives and integrals, many epidemiological models and other models in engineering and science have successful being originated and analyzed [15-26]. Fractional order derivatives has a significant characteristics called memory effect and this extraordinary property do not exist in the classical derivatives. These derivatives are nonlocal opposed to the local behavior of integer derivatives. It implies the next state of a fractional system depends not only upon its current state but also upon all of its historical states. Petras and Magin discussed in [27], "it is clear that the state of several systems (electrochemical, biological, etc.) at a given time depends on their configuration at previous times". The most important purpose of this paper is to originate and present a dengue fever transmission model with and without immunity using fractional order derivatives which has an advantage over the classical integer order models. Stability analysis of the model will also be a part of this paper.

II. MODEL FORMULATION

This part of paper deals with formulation of dengue disease transmission model of a non-integer order IVP (Initial Value Problem) using fractional order derivatives.

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Fractional calculus is an attractive and dominant tool for mathematical modeling. It has been applied in many areas of research such as economics, science and engineering. There are many interesting and attractive definitions of fractional derivatives in fractional calculus [26, 28], but here the famous Caputo derivatives is used due its advantage on initial value problems. Some important definitions related to frictional calculus are given below:

Definition 2.1. [26, 28], Frictional integral of order α is defined by

$$I^\alpha g(t) = \frac{1}{\Gamma(\alpha)} \int_0^t \frac{g(x)}{(t-x)^{1-\alpha}} dx$$

for $0 < t, 0 < \alpha < 1$.

Definition 2.2. [26, 28], Caputo Fractional derivative is defined by

$$D^\alpha g(t) = \frac{1}{\Gamma(p-\alpha)} \int_0^t \frac{g^p(x)}{(t-x)^{\alpha+1-p}} dx,$$

for $p-1 < \alpha < p$.

A Host-Vector dengue disease transmission model was developed by Esteva and Vargas in [29], and suppose that a recover individual from the disease will not be reinfected by the disease. They also assume that the host population H is constant with death and birth rate μ_h . Where S_h, I_h, R_h are susceptible, infective, and recover individuals in the host population and S_v, I_v are susceptible, Infective in the vector population V . Their model is given as follows:

$$\frac{dS_h}{dt} = \mu_h H - \frac{\beta_h b}{H} S_h I_v - \mu_h S_h$$

$$\frac{dI_h}{dt} = \frac{\beta_h b}{H} S_h I_v - (\mu_h + \gamma_h) I_h$$

$$\frac{dR_h}{dt} = \gamma_h I_h - \mu_h R_h \quad (1)$$

$$\frac{dS_v}{dt} = \mu_v - \frac{\beta_v b}{H} S_v I_h - \mu_v S_v$$

$$\frac{dI_v}{dt} = \frac{\beta_v b}{H} S_v I_h - \mu_v I_v,$$

where β_h, β_v are the transmission probability from vector to host and host to vector. γ_h represent the recovery rate in the host population and b is the

biting rate of the vector. Furthermore equation (1) can be reduced to three dimension dynamics with the condition $S_h + I_h + R_h = H$ and

$$S_v + I_v = \frac{\wedge}{\mu_v}$$

$$\frac{dS_h}{dt} = \mu_h H - \frac{\beta_h b}{H} S_h I_v - \mu_h S_h$$

$$\frac{dI_h}{dt} = \frac{\beta_h b}{H} S_h I_v - (\mu_h + \gamma_h) I_h \quad (2)$$

$$\frac{dI_v}{dt} = \frac{\beta_v b}{H} S_v I_h - \mu_v I_v$$

To normalize (2), we set $S = \frac{S_h}{H}, i = \frac{I_h}{H}, v = \frac{I_v}{\wedge / \mu_v}$

and get

$$\frac{d}{dt} s = \mu(1-s) - \rho sv$$

$$\frac{d}{dt} i = \rho sv - \beta i \quad (3)$$

$$\frac{d}{dt} v = \gamma(1-v)i - \delta v$$

Where

$$\rho = \frac{b\beta_h \wedge}{\mu_v H}, \beta = \gamma_h + \mu_h, \gamma = b\beta_v, \delta = \mu_v, \mu = \mu_h$$

Replacing the integer derivatives in system (3) by Caputo derivatives of order α we obtain:

$$D^\alpha s = \mu(1-s) - \rho sv$$

$$D^\alpha i = \rho sv - \beta i \quad (4)$$

$$D^\alpha z = \gamma(1-v)i - \delta v$$

If we follow the method used by Diethelm [32], system (4) becomes

$$D^\alpha s = \mu^\alpha (1-s) - \rho^\alpha sv$$

$$D^\alpha i = \rho^\alpha sv - \beta^\alpha i \quad (5)$$

$$D^\alpha v = \gamma^\alpha (1-v)i - \delta^\alpha v$$

Now, it is understandable that the dimension on both sides of the system(5) correspond to $(times)^\alpha$. It is noticeable that when the fractional order α approaches to 1, then the model problem (5), becomes the classical endemic model (3).

III. NON-NEGATIVE SOLUTION

Let us assume $R^3 = \{Z \in R^3 : Z \geq 0\}$, where $Z = (s, i, v)^T$. To show the non-negative solution of the model we will apply the following lemma presented in [30].

Lemma: 3.1. [30], Generalized Mean Value Theorem: Let $g(x) \in C[c, d]$ and $D^\alpha g(x) \in C[c, d]$ for $0 \leq \alpha \leq 1$, then we have

$$g(x) - g(c) + \frac{1}{\Gamma(\alpha)} D^\alpha g(\xi)(x-c)^\alpha,$$

with the condition $c \leq \xi \leq x$, for all $x \in [c, d]$.

Remark 3.1: Assume that $g(x) \in C[c, d]$ and $D^\alpha g(x) \in C[c, d]$, for $0 \leq \alpha \leq 1$. It follows from lemma 3.1 that $g(x)$ is non-decreasing if $D^\alpha g(x) \geq 0$, for all $x \in [c, d]$ and $g(x)$ non increasing if $D^\alpha g(x) \leq 0$ for all $x \in [c, d]$.

Theorem 3.1 A unique solution of the fractional order initial value problem (5) exists and it remains in R^3_+ . on each hyperplane bounding the non negative orthant, the vector field points into R^3_+ .

Proof 3.1 Existence and uniqueness of the solution of model problem (5) in $(0, \infty)$ follows by the use of theorem 3.1 and remark 3.2 in [31]. The domain R^3_+ is positively invariant for the model problem, because

$$D^\alpha s|_{s=0} = \mu \geq 0$$

$$D^\alpha i|_{i=0} = 0$$

$$D^\alpha v|_{v=0} = \gamma \geq 0,$$

on each hyper plane bounding the non negative outhunt, the vector field points into R^3_+ .

IV. EQUILIBRIA OF THE MODEL

The frictional order model (5) has two biological meaningful equilibrium points i.e. disease free and endemic equilibrium. These two equilibria depending on i and v : if there is no disease for host and mosquitoes i.e. if $i = v = 0$, then the equilibrium point is said to be disease free equilibrium, and if $i \neq 0, v \neq 0$, then the equilibrium point is called endemic. To determine the equilibria of the frictional order model (5). Assume that

$$D^\alpha s = 0, D^\alpha i = 0, D^\alpha v = 0, \text{ we get}$$

$$0 = \mu^\alpha (1-s) - \rho^\alpha sv$$

$$0 = \rho^\alpha sv - \beta^\alpha i$$

$$0 = \gamma^\alpha (1-v)i - \delta^\alpha v,$$

then the disease free equilibrium $E_f(1, 0, 0)$ and endemic equilibrium $E_e(s^*, i^*, v^*)$, where

$$s^* = \frac{\mu^\alpha \gamma^\alpha + \beta^\alpha \delta^\alpha}{\gamma^\alpha (\mu^\alpha + \rho^\alpha)}, i^* = \frac{\mu^\alpha (\rho^\alpha \gamma^\alpha - \beta^\alpha \delta^\alpha)}{\beta^\alpha \gamma^\alpha (\mu^\alpha + \rho^\alpha)},$$

and $v^* = \frac{\mu^\alpha (\rho^\alpha \gamma^\alpha - \beta^\alpha \delta^\alpha)}{\rho^\alpha (\mu^\alpha \gamma^\alpha + \beta^\alpha \delta^\alpha)}$. by and

The second fixed point exists only if the threshold parameter

$$R = \frac{\rho^\alpha \gamma^\alpha}{\beta^\alpha \delta^\alpha} > 0,$$

the basic reproductive number is denoted by R_0 and is equal to \sqrt{R} .

Theorem 4.1 *The disease free equilibrium E_f is globally stable if $R < 1$. If $R > 1$, then the endemic fixed point E_e is globally asymptotically stable and E_f is unstable.*

It is proved in [29] that disease free equilibrium point is globally stable if $R < 1$. For $R > 1$ the fixed point E_e becomes locally asymptotically stable and the fixed point E_f becomes unstable. The global stability is shown using the property of stability of periodic orbits.

V. MODEL WITHOUT IMMUNITY

There is no enough information about the immunity after recovery for Dengue Fever, that's why we suppose that the immune individuals in the host population is negligible. In this case the Host-Vector Model (1) for Dengue Transmission becomes as:

$$\begin{aligned} \frac{ds_h}{dt} &= \mu_h H - \frac{\beta_h b}{H} S_h I_v - \mu_h S_h \\ \frac{dI_h}{dt} &= \frac{\beta_h b}{H} S_h I_v - (\mu_h + \gamma_h) I_h \\ \frac{dS_v}{dt} &= \wedge - \frac{\beta_v b}{H} S_v I_h - \mu_v S_v \\ \frac{dI_v}{dt} &= \frac{\beta_v b}{H} S_v I_h - \mu_u I_v, \end{aligned} \quad (6)$$

where $S_h + I_h = H$ and $S_v + I_v = \wedge$ Simplifying in the

same way, we get

$$\begin{aligned} \frac{d}{dt} i &= \rho(1-i)v - \beta i \\ \frac{d}{dt} v &= \gamma(1-v)i - \delta v. \end{aligned} \quad (7)$$

Replacing the integer derivatives in system (7) by Caputo derivatives of order α then the system can be written as:

$$\begin{aligned} D^\alpha i &= \rho(1-i)v - \beta i \\ D^\alpha v &= \gamma(1-v)i - \delta v, \end{aligned} \quad (8)$$

Using the same technique used by Diethelm [32], system (8) becomes

$$\begin{aligned} D^\alpha i &= \rho^\alpha (1-i)v - \beta^\alpha i \\ D^\alpha v &= \gamma^\alpha (1-v)i - \delta^\alpha v \end{aligned} \quad (9)$$

In this case we have two fixed point (0,0) and

$$(i^*, v^*), \text{ Where } i^* = \frac{\rho^\alpha \gamma^\alpha - \beta^\alpha \delta^\alpha}{\gamma^\alpha (\rho^\alpha + \beta^\alpha)} \text{ and } v^* = \frac{\rho^\alpha \gamma^\alpha - \beta^\alpha \delta^\alpha}{\rho^\alpha (\gamma^\alpha + \delta^\alpha)}$$

It is simple and easy to and the basic reproduction number for system (9). The origin is locally stable if the basic reproduction number is less than one and unstable if the basic reproduction number greater than one.

VI. CONCLUSION

In this research we formulated a frictional order Host-vector model for dengue fever. In our work we established the existence and uniqueness of non-negative solutions of the frictional order model. We have shown that there are two biological meaningful equilibria of the system and proved that the disease free equilibrium E_f is globally stable if the basic reproduction number less than one. If the basic reproduction number greater than one, then the endemic fixed point E_e is globally asymptotically stable and E_f is unstable. In last section the model is analyzed and formulated without immunity.

REFERENCES

- Xue Y, Yuan X, Liu M. Global stability of a multi-group SEI model, Applied Mathematics and Computation. 2014;226:51-60.
- Magal P, Ruan S. Susceptible-infectious-recovered models revisited: From the individual level to the population level. Mathematical , Mathematical Biosciences. 2014;250:26-40.
- Hethcote HW, van den Driessche. Two SIS epidemiologic models with delays, J. Math. Biol. 2000;40:3-26.
- J. Dushoff J, Huang W, Castillo-Chavez C. Backwards bifurcations and catastrophe in simple models of fatal diseases, Journal of Mathematical Biology. 1998;36:227-248.
- Castillo-Chavez C, Thieme HR. Asymptotically autonomous epidemic models, Mathematical Population Dynamics: Analysis of Heterogeneity. 1995;1:33-50.
- Chitnis N, Cushing JM, Hyman JM. Bifurcation analysis of a mathematical model for malaria transmission, SIAM Journal on Applied Mathematics. 2006;67:24-45.
- Okosun KO, Makinde OD. Optimal control analysis of malaria in the presence of non-linear incidence rate, Appl. Comput. Math. 2013;12:20-32.
- Koella JC, Anita R. Epidemiological models for the spread of anti-malarial resistance, Malaria Journal. 2003;2:3.
- Mukandavire Z, Gumel AB, Garira W, Tchuente JM. Mathematical analysis of a model for HIV-malaria co infection, Math Biosci Eng. 2009;6:333-362.
- Brauer F, Sanchez DA. Constant rate population harvesting: Equilibrium and stability, Theoretical Population Biology. 1975;8(1):12-30.
- Feng Z, Velasco-Hernandez JX. Competitive exclusion in a vector-host model for the dengue fever, Journal of Mathematical Biology. 1997;35(5):523-544.
- Hethcote HW. Note on determining the limiting susceptible population in an epidemic model., Mathematical Biosciences. 1970;9:161163.

13. Ngwa GA, Shu WS. A. A mathematical model for endemic malaria with variable human and mosquito populations, *Mathematical and Computer Modelling*. 2000;32(7-8):747-763.
14. Hethcote HW. Qualitative analyses of communicable disease models, *Mathematical Bio- sciences*. 1976;28(3-4):335-356.
15. Ahmed A, El-Sayed AMA, El-Saka HA. Equilibrium points, stability and numerical solutions of fractional-order predator-prey and rabies models, *Journal of Mathematical Analysis and Applications*. 2007;325:542-553.
16. Arafa AAM, Rida SZ, Khalil M. Solution of fractional order model of childhood diseases with constant vaccination strategy, *Mathematical Sciences Letters*. 2012;1:17-23.
17. Javidi M, Ahmad B. A study of fractional-order cholera mode, *Appl: Math. Inf. Sci*. 2014; 8:2195-2206.
18. Rocco A, West BJ. Fractional calculus and the evolution of fractional phenomena, *Physica A: Statistical Mechanics and Its Applications*. 1999;265(3-4):535-546.
19. Ding Y, Ye H. A fractional-order differential equation model of HIV infection of CD4+T- cells, *Mathematical and Computer Modelling*. 2009;50(3-4):386-392.
20. Ahmed E, El-Saka HA. On fractional order models for hepatitis C, *Nonlinear Biomedical Physics*. 2010;4:1-3.
21. Pinto CMA, Machado JAT. Fractional model for malaria transmission under control strategies, *Computers and Mathematics with Applications*. 2013;66(5):908-916.
22. Liu Y, Lu P, Szanton I. Numerical analysis for a fractional differential time-delay model of HIV infection of CD4 + T-cell proliferation under antiretroviral therapy, *Abstract and Applied Analysis*. 2014;Vol. 2014, Article ID 291614, 13 pages, doi:10.1155/2014/291614.
23. Ye H, Ding Y. Nonlinear dynamics and chaos in a fractional-order HIV model, *Mathematical Problems in Engineering*. 2009; vol. 2009, Article ID 378614, 12 pages. doi:10.1155/2009/378614.
24. Gokdogan A, Yildirim A. Merdan M. Solving a fractional order model of HIV infection of CD4+T cells, *Mathematical and Computer Modelling*. 2011;54:21322138.
25. El-Saka HAA. Backward bifurcations in fractional-order vaccination model, *Journal of the Egyptian Mathematical Society*. 2015;23(1):49-55.
26. Petras I. Fractional-order nonlinear systems: Modeling, analysis and simulation, Springer- Verlag; 2011.
27. Petras I, Magin IR. Simulation of drug uptake in a two compartmental fractional model for a biological system, *Commun Nonlinear Sci Numer Simul*. 2011;16(12):4588-4595.
28. Podlubny I. Fractional differential equations, Academic Press, New York; 1999.
29. Esteva L and Vargas C. Analysis of a Dengue Fever Transmission Model, *Journal of Bioscience* 150(1998), 131-151.
30. Odibat ZM, Shawagfeh NT. Generalized Taylor's formula, *Applied Mathematics and Computation*. 2007;186(1):286-293.
31. Lin W. Global existence theory and chaos control of fractional differential equations, *Journal of Mathematical Analysis and Applications*. 2007;332(1):709-726.
32. Diethelm K. A fractional calculus based model for the simulation of an outbreak of dengue fever, *Nonlinear Dynamics*. 2013;71(4):613-619.