

A Mathematical Model Analysis of Age Structured Malaria Transmission

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Abstract

Mathematical models have been used to provide an explicit framework for understanding malaria transmission dynamics in human population for over 100 years. The equilibrium points of the model are found and their stability is investigated. By analyzing the model, a threshold parameter R_0 was found which the basic reproductive number is obtained. It is noted that when $R_0 < 1$ the disease will fail to spread and when $R_0 > 1$ the disease will persist in the population and become endemic. The stability of these two equilibrium points is controlled by the basic reproductive number R_0 which was determined using the next generation matrix approach. Our model divides the human population into three compartments: pre-school (0-5 years) and youngest age group (6 -25 years old) and the rest of the human population. The results of our mathematical analysis indicate that the disease-free equilibrium is asymptotically stable and occurs whenever the basic reproduction number R_0 is less than unity or $R_0 < 1$. For $R_0 > 1$, the disease-free equilibrium point is unstable.

Keywords: Mathematical model, equilibrium point, Basic Reproductive number, Stability

DOI: 10.7176/MTM/9-5-04

Publication date: May 31st 2019

INTRODUCTION

In order to reflect the age structure of a population affects the dynamics of disease transmission, we incorporate age into the model, and assume that the infection rate depend on the age. We model observations to be able to describe naturally occurring phenomena, to be able to predict future events and to be able to find optimal solutions to problems we encounter.

That first malaria model was expanded in several ways, for example by introducing the effects of human age [21], acquired immunity to malaria [17, 21] or genetic and special heterogeneity of parasite and host among these age and immunity play an important role in endemic malaria regions. In order to describe age-structured models for disease transmission we must first develop the theory of age-structured populations. In fact, the first models for age-structured populations were designed for the study of disease transmission in such populations. In this paper, we construct a new model to investigate the patterns of malaria transmission in the presence of age structure model.

Disease Free Equilibrium point

Disease-free equilibrium points (*DFE*) are steady state solutions where there is no malaria in the human population or Plasmodium parasite in the mosquito population. In absence of the disease, this implies that $i_{h1}, i_{h2}, i_{h3}, i_v = 0$. To determine the stability of the equilibrium points of the given model we evaluate the steady state of the given system with following conditions.

- Disease free equilibrium points $i_{h1}, i_{h2}, i_{h3}, i_v = 0$

1.1. Endemic disease equilibrium point

The equilibrium point at the endemic level has a strong influence in the behavior of the disease transmission for a given community. The endemic equilibrium point is the final reachable situation, even though there could be infinitely many different initial distributions of malaria in a community. Stability analysis is an important mathematical tool that provides the conditions that guarantee the stability of an endemic equilibrium point. In the presence of the disease, this implies that $i_{h1}, i_{h2}, i_{h3}, i_v \neq 0$

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1.2. Basic concept of reproductive number R_0

A quantity of central importance in epidemiology is the basic reproduction number denoted by R_0 . From time to time, people also call it the basic reproductive rate or ratio, or the basic reproduction ratio. R_0 is defined as the mean number of secondary infections produced when one infected individual is introduced into a host population where everyone is susceptible. As defined, a reproductive number less than one will cause the disease to die out; whereas, a reproductive number greater than one classifies the disease as endemic [2]. If $R_0 < 1$, then on average an infected individual produces less than one new infected individual over the course of its infectious period, and the infection cannot grow. Conversely, if $R_0 > 1$, then each infected individual produces, on average, more than one new infection, and the disease can invade the population.

1.3. Next generation matrices for compartmental epidemic model

The basic reproduction number R_0 is arguably the most important quantity in infectious disease epidemiology.

The next-generation matrix (NGM) is the natural basis for the definition and calculation of R_0 where finitely many different categories of individuals are recognized.

In epidemiology, the next-generation matrix is a method used to derive the basic reproduction number, for a compartmental model of the spread of infectious diseases. This method is given by Diekmann *et al.* (1990)[12] and Driessche and Watmough (2002).

To calculate the basic reproduction number by using a next-generation matrix, the whole population is divided into n compartments in which there are $m < n$ infected compartments. Let $x_i, i = 1, 2, 3, 4, \dots, m$ be the numbers of infected individuals in the i^{th} infected compartment at time t .

1.4. Routh-Hurwitz Criteria

1.4.1. General Procedure

The Routh-Hurwitz (RH) Criterion is a general mathematical technique that may be used to determine how many of the roots of a characteristic equation such as the one below have positive real parts, and are therefore unstable. Routh-Hurwitz is a method for determining whether a linear system is stable or not by examining the location of the roots of the characteristic equation of the system.

The Routh-Hurwitz Criteria do not yield information about the absolute values of the eigenvalues, but about their signs. For checking for stability, it is sufficient to know the sign of the real parts.

Consider the general form for a characteristic polynomial

$$p(\lambda) = \lambda^n + a_{n-1}\lambda^{n-1} + a_{n-2}\lambda^{n-2} + a_{n-3}\lambda^{n-3} + \dots + a_1\lambda + a_0$$

To determine whether the system is stable or not, check the following conditions.

1. Two necessary but not sufficient conditions that all the roots have negative real parts are:

- a) All the polynomial coefficients must have the same sign.
- b) All the polynomial coefficients must be nonzero.

2. If this condition is satisfied then compute the Routh-Hurwitz array as follows:

λ^n	a_n	a_{n-2}	a_{n-4}	a_{n-6}	\dots
λ^{n-1}	a_{n-1}	a_{n-3}	a_{n-5}	a_{n-7}	\dots
λ^{n-2}	b_1	b_2	b_3	b_4	\dots
λ^{n-3}	k_1	k_2	k_3	k_4	\dots
\vdots					
$\lambda^{0!}$					

Where the a_i coefficients come from the original polynomial. The other coefficients are computed in a fashion similar to a determinant, by selecting appropriate values in the array. The pattern of values is illustrated via the following cases:

The associated Routh Array is then:

λ^n	a_n	a_{n-2}	a_{n-4}	...
λ^{n-1}	a_{n-1}	a_{n-3}	a_{n-5}	...
λ^{n-2}	b_1	b_2	b_3	...
λ^{n-3}	k_1	k_2	k_3	
...				
λ^0				

Where

$$b_1 = \frac{-1}{a_{n-1}} \begin{vmatrix} a_n & a_{n-2} \\ a_{n-1} & a_{n-3} \end{vmatrix} = \frac{-1}{a_{n-1}} (a_n a_{n-3} - a_{n-2} a_{n-1})$$

$$b_2 = \frac{-1}{a_{n-1}} \begin{vmatrix} a_n & a_{n-4} \\ a_{n-1} & a_{n-5} \end{vmatrix} = \frac{-1}{a_{n-1}} (a_n a_{n-5} - a_{n-4} a_{n-1})$$

$$b_3 = \frac{-1}{a_{n-1}} \begin{vmatrix} a_n & a_{n-6} \\ a_{n-1} & a_{n-7} \end{vmatrix} = \frac{-1}{a_{n-1}} (a_n a_{n-7} - a_{n-6} a_{n-1})$$

$$k_1 = \frac{-1}{b_1} \begin{vmatrix} a_{n-1} & a_{n-3} \\ b_1 & b_2 \end{vmatrix} = \frac{-1}{b_1} (a_{n-1} b_2 - b_1 a_{n-3})$$

$$k_2 = \frac{-1}{b_1} \begin{vmatrix} a_{n-1} & a_{n-5} \\ b_1 & b_3 \end{vmatrix} = \frac{-1}{b_1} (a_{n-1} b_3 - b_1 a_{n-5})$$

THE BASICS MODEL

These categories or compartments are represented by standard notation of *S-E-I-R* after the pioneering work of Kermack and McKendrick. There may be variations in the compartment structure depending on the type of disease. For example, the *I* class of individuals may not recover at all and die; *R* can consist of individuals, who recover with temporary or permanent immunity, thereby further subdividing the epidemiological compartments. Using these notations, eight classes of compartmental models are possible - *SI*, *SIS*, *SEI*, *SEIS*, *SIR*, *SIRS*, *SEIR* and *SEIRS*. For example, in an *SEIRS* model, a fraction of the susceptible (*S*) population gets exposed (*E*) to infection, a part of which then becomes infectious (*I*). Some from the *I* class recover from the disease, and become part of the *R* class with temporary immunity. When immunity is lost, they become susceptible to pathogen attack again, and enter the *S* class. But in this paper we consider the standard *SIR* model where the individuals in the population are divided into three compartments. The susceptible (*S*) which refers to the healthy individuals that has not yet come into contact with malaria disease. The infective (*I*) are those who have become infected with malaria disease and are able to transmit the disease and the recovered (*R*) are individuals who have recovered from malaria disease. The proportions of the individuals in the compartment of the population, i.e. *S*, *I* and *R*, at time *t* is denoted as *S*(*t*), *I*(*t*) and *R*(*t*) respectively.

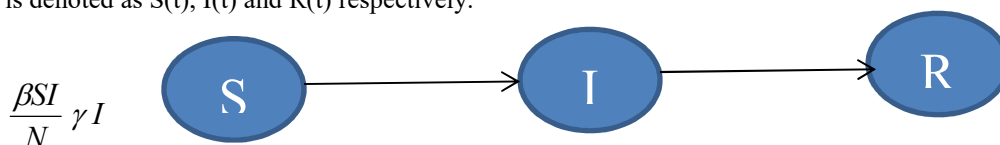


Figure 3 Flow chart showing the SIR model

β = The per-capita contact rate between any two individuals.

$\frac{1}{N}$ = The proportion of infectious individuals in the population.

γ = The per-capita rate of recovery once infectious.

$\gamma \times I$ = The rate at which infectious individuals are removed from the infectious class.

$\beta \times S \times \frac{1}{N}$ = The rate at which susceptible individuals come across infectious individuals and then become infectious.

3.1. Model description and formulation

We formulate a mathematical model for the transmission and spread of malaria in the human and mosquito population. We formulate a model for the spread of malaria in the human and mosquito population with the total population size at time t given is by is given by $N_H(t)$ and $N_v(t)$ respectively. These are further compartmentalized into epidemiological classes shown below. The vector component of the model does not include immune class as mosquito never recover from infection, that is, their infective period ends with their death due to their relatively short life-cycle. Thus the immune class in the mosquito population is negligible and death occurs equally in all groups. Our model also excludes the immature mosquitoes since they do not participate in the infection cycle and are, thus, in the waiting period, which limits the vector population growth. Our model has the following variables and parameters:

$S_H(t)$ the number of susceptible human hosts at time t , $I_H(t)$ the number of infected human hosts at time t , $R_H(t)$ the number of partially immune human hosts at time t , $S_v(t)$ the number of susceptible mosquito vectors at time t , $I_v(t)$ the number of infected mosquito vectors at time t , Given that the total number of bites made by the mosquitoes must equal the number of bites received by the humans, $(N_H, N_v) = \frac{N_v}{N_H}$ is a constant. Thus

$$m = \frac{N_v}{N_H} \text{ is number of female mosquitoes per human host.}$$

Where $m = \frac{N_v}{N_H}$ is the female vector – host ratio, defined as the number of female mosquitoes per human host

[2].

Our model, illustrated in **Figure 3**, seeks to look into the dynamics of the Susceptible-Infectious-Recovered and Susceptible Infectious (SIR-SI) malaria transmission, taking into account the three human classes, pre-school children (H_1 – aged five years below), age from six to 25 (H_2 5–25 years old) and the rest of the humans (H_3 – older than twenty five years).

The human population N_H is divided into nine compartments denoted by the following variables:

S_{H1} is the number of susceptible humans of age five (5) years and below;

I_{H1} is the number of infectious humans of age five (5) years and below;

R_{H1} is the number of recovered humans (with temporary immunity) of age five (5) years and below;

S_{H2} is the number of susceptible humans older than five (5) years and below 25 years old ;

I_{H2} is the number of infectious humans older than five (5) years and below 25 years old;

R_{H2} is the number of recovered humans (with temporary immunity) older than five (5) years and below 26 years old

S_{H3} is the number of susceptible humans of age greater than or equal to 25 years

I_{H3} is the number of infectious humans of age greater than or equal to 25 years

R_{H3} is the number of recovered humans of age greater than or equal to 25 years

In general, any member of the human population (pre-school or older) remains in the susceptible class S for a certain period unless bitten and becomes infectious. Once in the infective class I , an individual may recover without acquiring immunity and be transferred back into the susceptible class S , or stays in class I while infectious before moving to the recovered class R with an acquired temporary immunity. Humans in the recovered class may lose immunity and move back to the susceptible class.

Humans leave the population through natural death rate and through per-capita death rate due to infection.

In particular, members of the susceptible pre-school children S_{H1} leave the population and move to infectious I_{H1} or grow older to susceptible S_{H2} . Infectious I_{H1} leave the population and transfer to the infectious I_{H1} , or recovered R_{H1} . For the recovered R_{H1} population, members may grow to the recovered R_{H2} . Similarly, for the rest of the human population, susceptible S_{H2} members may move to the infectious class I_{H2} . Infectious I_{H2} members can transfer to the recovered R_{H2} . Recovered R_{H2} members can lose immunity and assume that not move back to the susceptible S_{H2} group.

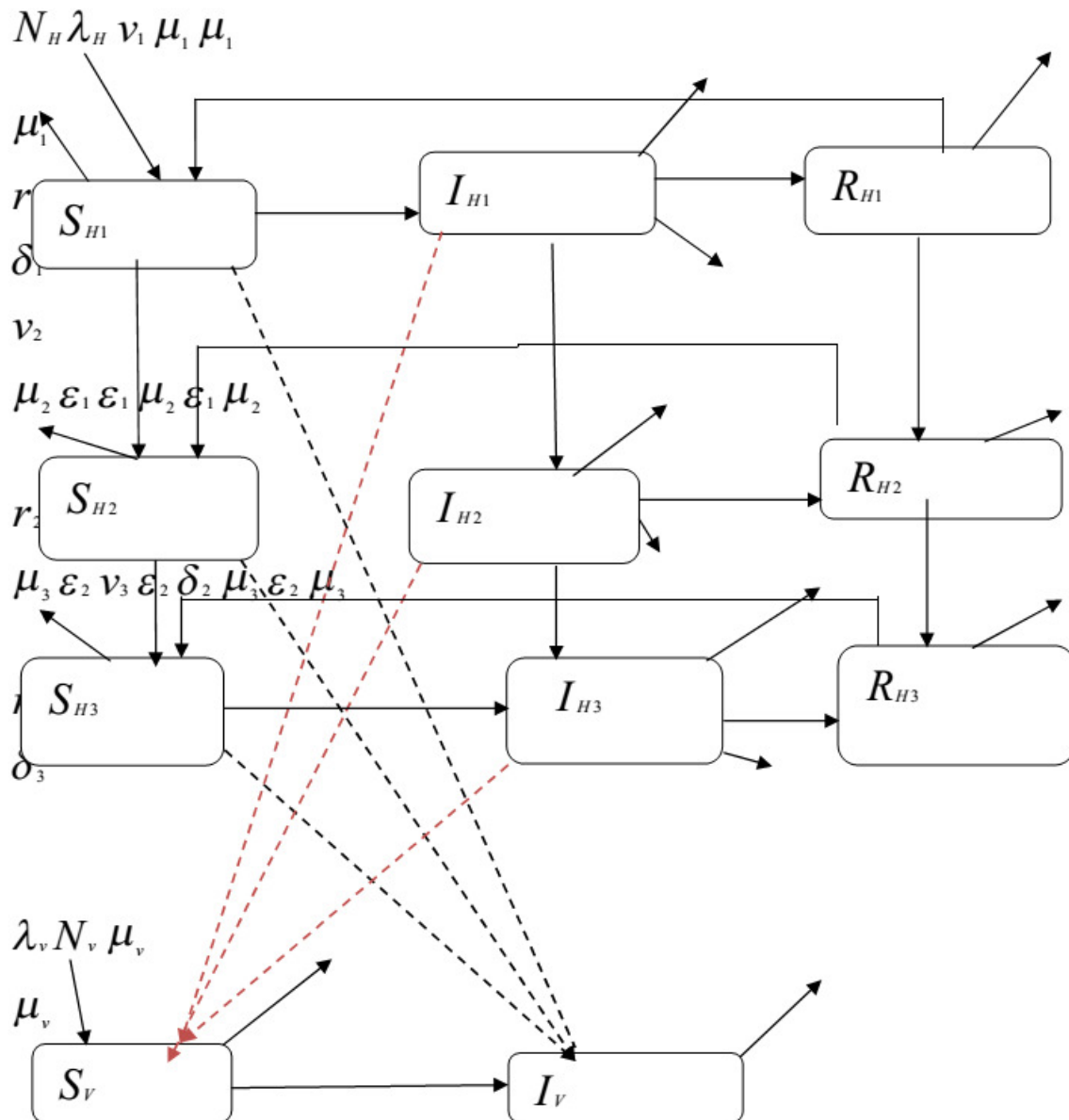


Figure 4: transmission pattern of SIR – SI with age-structured human population susceptible mosquitoes S_v may transfer into the infective class I_v . Due to short life cycle of mosquitoes, the infective period ends with their death.

3.2. Model Equations

Applying the assumptions, definitions of variables and parameters and description of terms above, the differential equations which describe the dynamics of malaria in the human and mosquito populations are formulated as shown below:

For first age group (0-5 years) we formulate:

$$\frac{ds_{H1}}{dt} = \lambda_H N_H - \mu_1 S_{H1} - \frac{a_1^b s_{H1} I_V}{N_H} + V_1 R_{H1} - \varepsilon_1 S_{H1} \quad (3.5.1)$$

$$\frac{dI_{H1}}{dt} = \frac{a_1^b S_{H1} I_V}{N_H} - \mu_1 I_{H1} - r_1 I_{H1} - \delta_1 I_{H1} - \varepsilon_1 I_{H1} \quad (3.5.2)$$

$$\frac{dR_{H1}}{dt} = -\mu_1 R_{H1} + r_1 R_{H1} - \varepsilon_1 R_{H1} - v_1 R_{H1} \quad (3.5.3)$$

For second age group (5-25 years) we formulated

$$\frac{ds_{H2}}{dt} = -\mu_2 s_{H2} - \frac{a_2^b S_{H2} I_V}{N_H} + V_2 R_{H2} + \varepsilon_1 S_{H1} - \varepsilon_2 S_{H2} \quad (3.5.4)$$

$$\frac{dI_{H2}}{dt} = \frac{a_2^b S_{H2} I_V}{N_H} - \mu_2 I_{H2} - r_2 I_{H2} - \delta_2 I_{H2} + \varepsilon_1 I_{H1} - \varepsilon_2 I_{H2} \quad (3.5.5)$$

$$\frac{dR_{H2}}{dt} = -\mu_2 R_{H2} + r_1 I_{H2} + \varepsilon_1 R_{H1} - \varepsilon_2 R_{H2} - v_2 R_{H2} \quad (3.5.6)$$

For the remaining age group (5-25 years) we formulate

$$\frac{ds_{H3}}{dt} = -\mu_3 s_{H3} - \frac{a_3^b S_{H3} I_V}{N_H} + V_3 R_{H3} + \varepsilon_2 S_{H2} \quad (3.5.7)$$

$$\frac{dI_{H3}}{dt} = \frac{a_3^b S_{H3} I_V}{N_H} - \mu_3 I_{H3} - r_3 I_{H3} - \delta_3 I_{H3} + \varepsilon_2 I_{H2} \quad (3.5.8)$$

$$\frac{dR_{H3}}{dt} = -\mu_3 I_{H3} + r_3 I_{H3} + \varepsilon_2 I_{H2} - v_3 R_{H3} \quad (3.5.9)$$

$$\frac{dS_V}{dt} = \lambda_V N_V - \frac{a_1^c S_V I_{H1}}{N_H} - \frac{a_2^c S_V I_{H2}}{N_H} - \frac{a_3^c S_V I_{H3}}{N_H} - \mu_V S_V \quad (3.5.10)$$

$$\frac{dI_V}{dt} = \frac{a_1^c S_V I_{H1}}{N_H} + \frac{a_2^c S_V I_{H2}}{N_H} + \frac{a_3^c S_V I_{H3}}{N_H} - \mu_V S_V \quad (3.5.11)$$

The total population size N_{H1} , N_{H2} , N_{H3} and N_V can be determined by

$$N_{H1} = S_{H1} + I_{H1} + R_{H1} = 1$$

$$N_{H2} = S_{H2} + I_{H2} + R_{H2} = 1$$

$$N_{H3} = S_{H3} + I_{H3} + R_{H3} = 1$$

The total population of the first age group is derived by adding equations (3.5.1)-(3.5.3) to obtain

$$\frac{dN_{H1}}{dt} = \lambda_H N_{H1} - (\mu_1 + \varepsilon_1) S_{H1} - (\mu_1 + \varepsilon_1 + \delta_1) I_{H1} - (\mu_1 + \varepsilon_1) R_{H1}$$

The total population of the second age group is derived by adding equations (3.5.4)-(3.5.6) to obtain

$$\frac{dN_{H2}}{dt} = (\mu_2 + \varepsilon_2) S_{H2} - (\mu_2 + \varepsilon_2 + \delta_2) I_{H2} - (\mu_2 + \varepsilon_2) R_{H2} + \varepsilon_1 (S_{H1} + I_{H1} + R_{H1})$$

And the total population of the second age group is derived by adding equations (3.5.7)-(3.5.9) to obtain

$$\frac{dN_{H3}}{dt} = -\mu_3 S_{H3} - (\mu_3 + \delta_3) I_{H3} + \varepsilon_2 (S_{H2} + I_{H2} + R_{H2}) - \mu_3 R_{H3}$$

In the model, the term $\frac{a_i b S_{H1} I_v}{N_H}$ denotes the rate at which the human hosts S_{H1} get infected by infected

mosquitoes I_v for $i = 1, 2, 3$ and $\frac{a_j c S_v I_H}{N_H}$ refers to the rate at which the susceptible mosquitoes S_v are infected

by the infected human hosts I_H for $j = 1, 2, 3$. It is important to note that the rate of infection of human host S_H by infected vector I_v is dependent on the total number of humans N_H available per infected vector [19] and [36]. The model assumes that all newborns are susceptible in both human and mosquito populations.

The per capita birth rate for humans and mosquito populations, λ_H and λ_v respectively, are both positive. The human and mosquito populations experience per capita natural death rate μ_1, μ_2, μ_3 and μ_v respectively. Infected humans die due to infection at per capita death rate δ_1, δ_2 and δ_3 respectively. We can consider recovered human individuals lose their immunity at per capital rates. To ensure that we have a stable positive human population, we assume that $\lambda_H > \delta_1 + \delta_2 + \delta_3 + \mu_1 + \mu_2 + \mu_3$.

Moreover, it is clear that the birth rate should be greater than the per capita rate ε_1 and ε_2 of growing up.

Consider the interaction between the two populations shown in **Figure 3**. If a_1 is the average number of bites per mosquito per unit time, then there are $\frac{a_1 N_v}{N_H}$ bites per human per time. For the pre-school humans H_1 , note that

there are S_{H1} susceptible humans and the proportion of the total number of bites that are possibly infectious to humans is $\frac{I_v}{N_v}$. Then the number of potentially infectious bites given to susceptible pre-school children is

$\frac{a_1 I_v S_{H1}}{N_H}$ bites per time. However, only a fraction of these bites, namely b , successfully infect the pre-school

children. Hence we have:

$$\frac{\text{(pre-school humans infected)}}{\text{(unit time)}} = \frac{a_1 b I_v S_{H1}}{N_H}$$

The same formulation is constructed for the rest of the human group H_2 and H_3 . Similarly for the mosquito population, we have

$$\frac{\text{(mosquitoes infected)}}{\text{(unit time)}} = \frac{(a_1 c I_{H1} S_v)}{N_H} + \frac{(a_2 c I_{H2} S_v)}{N_H} + \frac{a_3 c S_v I_{H3}}{N_H}$$

To analyze the age-structured malaria model given by Equations (3.5.1)–(3.5.11), we construct the fractional quantities by scaling the population of each class by the total species population (We normalize our dynamical

equations by letting). The proportions for the system are $S_{h1} = \frac{S_{H1}}{N_H}, S_{h2} = \frac{S_{H2}}{N_H}, S_{h3} = \frac{S_{H3}}{N_H}, i_{h1} = \frac{I_{H1}}{N_H},$

$$i_{h2} = \frac{I_{H2}}{N_H}, i_{h3} = \frac{I_{H3}}{N_H}, r_{h1} = \frac{R_{H1}}{N_H}, r_{h2} = \frac{R_{H2}}{N_H}, r_{h3} = \frac{R_{H3}}{N_H}, s_v = \frac{S_v}{N_v} \text{ and } i_v = \frac{I_v}{N_v}$$

$$\text{Let } r_{h1} = 1 - (i_{h1} - s_{h1})$$

$$r_{h2} = 1 - (i_{h2} - s_{h2})$$

$$r_{h3} = 1 - (i_{h3} - s_{h3})$$

Differentiating the scaling equations (3.5.1)–(3.5.11) and solving for the derivatives of the scaled variables, we obtain the following reduced system of nine DE

$$\frac{di_{h1}}{dt} = \frac{1}{N_H} \left(\frac{dI_{H1}}{dt} - i_{h1} \frac{dN_H}{dt} \right) = abms_{h1}i_v - (r_1 + \delta_1 + \lambda_h)i_{h1} + \delta_1 i_{h1}^2 = 0 \quad (3.5.12)$$

$$\frac{ds_{h1}}{dt} = \frac{1}{N_{H1}} \left(\frac{ds_{H1}}{dt} - s_{h1} \frac{dN_{H1}}{dt} \right) = \lambda_h(1 - s_{h1}) - abms_{h1}i_v + v_1(1 - s_{h1} - i_{h1}) + \delta_1 s_{h1}i_{h1} = 0 \quad (3.5.13)$$

$$\frac{di_v}{dt} = \frac{1}{N_v} \left(\frac{dI_v}{dt} - i_v \frac{dN_v}{dt} \right) = ca_1(1 - i_v)i_{h1} - \lambda_v i_v = 0 \quad (3.5.14)$$

$$\frac{di_{h2}}{dt} = \frac{1}{N_{H2}} \left(\frac{dI_{H2}}{dt} - i_{h2} \frac{dN_{H2}}{dt} \right) = a_2 bm s_{h2} i_v - (r_2 + \delta_2 - \epsilon_1)i_{h2} - \epsilon_1 i_{h1} + \delta_2 i_{h2}^2 \quad (3.5.15)$$

$$\frac{ds_{h2}}{dt} = \frac{1}{N_{H2}} \left(\frac{ds_{H2}}{dt} - s_{h2} \frac{dN_{H2}}{dt} \right) = -a_2 bm s_{h2} i_{v2} - \epsilon_1 s_{h1} + \delta_2 s_{h2} i_{h2} - \epsilon_2 s_{h2} + v_2(1 - s_{h2} - i_{h2}) \quad (3.5.17)$$

$$\frac{di_{v2}}{dt} = \frac{1}{N_{v2}} \left(\frac{dI_{v2}}{dt} - i_{v2} \frac{dN_{v2}}{dt} \right) = a_2 c(1 - i_{v2})i_{h2} - \lambda_v i_v$$

$$\frac{di_{h3}}{dt} = \frac{1}{N_{H3}} \left(\frac{dI_{H3}}{dt} - i_{h3} \frac{dN_{H3}}{dt} \right)$$

$$= a_3 bm s_{h3} i_{v2} - (r_3 + \delta_3 + \epsilon_2)i_{h3} + \epsilon_2 i_{h2} + \delta_3 i_{h3}^2 \quad (3.5.18)$$

$$\frac{ds_{h3}}{dt} = \frac{1}{N_{H3}} \left(\frac{ds_{H3}}{dt} - s_{h3} \frac{dN_{H3}}{dt} \right) = -a_3 bm s_{h3} i_{v3} - \epsilon_2 s_{h3} + \epsilon_2 s_{h2} + \epsilon_2 s_{h2} + v_3(1 - s_{h3} - i_{h3}) + \delta_3 s_{h3} i_{h3}$$

$$\frac{di_{v1}}{dt} = \frac{1}{N_{v3}} \left(\frac{dI_{v1}}{dt} - i_{v1} \frac{dN_{v1}}{dt} \right) = ca_3(1 - i_v)i_{h3} - \lambda_v i_v = 0 \quad (3.5.20)$$

3.3. Existence and Stability of Equilibrium Points

3.3.1. Equilibrium points of the model

The equilibrium analysis helps to achieve this. Thus, we shall consider two equilibriums the diseases-free equilibrium points and endemic equilibrium points. The points at which the differential equations of the system (3.5.12)-(3.5.20) equal to zero are referred to as equilibrium points or steady-state solutions. It is important to note that there are no trivial equilibrium points as long as the recruitment terms N_{H1}, N_{H2}, N_{H3} and N_{v1}, N_{v2}, N_{v3} are not zero.

3.3.2. Existence of Disease-free Equilibrium Point

Disease-free equilibrium points are steady-state solutions where there is no malaria infection (no disease). We define the “diseased” classes as the human or mosquito populations that are either infectious, or recovered, that is $i_{h1}, i_{h2}, i_{h3}, i_v = 0$.

Thus, the disease-free equilibrium point, E_0 , for the malaria model (3.5.12)-(3.5.14) in the absence of disease implies that $i_{h1} = i_v = 0$.

From equation (3.5.12)-(3.5.14) at $i_{h1} = i_v = 0$ we have

$$\lambda_h(1 - s_{h1}) - v_1(1 - s_{h1}) = 0$$

$$\Rightarrow \lambda_h - \lambda_h s_{h1} + v_1(1 - s_{h1}) = 0$$

$$\Rightarrow s_{h1} = \frac{v_1 + \lambda_h}{v_1 + \lambda_h} = 1$$

This implies that the disease free equilibrium point is given by $(s_{h1}^*, i_{h1}, i_{v1}) = (1, 0, 0)$

3.3.3. Calculating Reproductive Number R_0 for the first age group

To calculate R_0 , we use the compartmental model approach presented by VandenDriessche and Watmough and Diekmann et al [12].

$$F = \left[\frac{\partial F_i(x_0)}{\partial x_j} \right] \text{ and } v = \left[\frac{\partial v_i(x_0)}{\partial x_j} \right]$$

where F_i = be the rate of appearance of new infections in compartments i

v_i = be the transfer of individuals out of compartments i by all other means

v_i^- = be the rate transfer of individuals out of compartment i

v_i^+ = be the rate transfer of individuals in to compartment i by all other means

x_0 = be the disease free equilibrium

R_0 = is the largest eigenvalues of Matrix of partial derivatives

$$\text{Then } Fv^{-1} = \left[\frac{\partial F_i(x_0)}{\partial x_j} \right] \times \left[\frac{\partial v_i(x_0)}{\partial x_j} \right]$$

To find reproduction number R_0 for the first age group will refer to the ODEs (3.5.12)-(3.5.14) and it is important to distinguish new infections from all other changes in population

Gains terms Loss terms

$$F_1 = a_1 b m i_v s_{h1} \quad v_1 = (r_1 + \delta_1 + \lambda_h) i_{h1} + \delta_1 i_{h1}^2$$

$$F_2 = 0 \quad v_2 = \lambda_h (1 - s_{h1}) + a_1 b m i_v s_{h1} + v_1 (1 - s_{h1} - i_{h1}) + \delta_1 s_{h1} i_{h1}$$

$$F_3 = ac(1 - i_v) \quad v_3 = \lambda_v i_v$$

Then the Jacobian matrix of F (gain terms) is given by

$$F(i_{h1}^*, s_{h1}^*, i_v^*) = \begin{bmatrix} 0 & a_1 b m i_v & a_1 b m s_{h1} \\ 0 & 0 & 0 \\ ca_1(1 - i_v) & 0 & -a_1 c i_{h1} \end{bmatrix}$$

Then the Jacobian of F at equilibrium point $E_0 = (s_{h1}^*, i_{h1}^*, i_v^*) = (1, 0, 0)$ is given by

$$F(0, 1, 0) = \begin{bmatrix} 0 & 0 & abms_{h1} \\ 0 & 0 & 0 \\ ca_1 & 0 & 0 \end{bmatrix}$$

And the Jacobian matrix of V (loss terms) is given by

$$v(i_{h1}^*, s_{h1}^*, i_v^*) = \begin{bmatrix} k_1 & 0 & 0 \\ k_2 & k_3 & k_4 \\ 0 & 0 & \lambda_v \end{bmatrix}$$

where

$$k_1 = r_1 + \delta_1 + \lambda_h + 2\delta_1 i_{h1}$$

$$k_2 = v_1 + \delta_1 s_{h1}$$

$$k_3 = v_1 + \lambda_h + \delta_1 i_{h1} + a_1 b m i_v$$

$$k_4 = a_1 b m s_{h1}$$

Then the Jacobian matrix of loss terms V at DFE (0, 1, 0) is given by

$$v(0,1,0) = \begin{bmatrix} k_{11} & 0 & 0 \\ k_{22} & k_{33} & k_{44} \\ 0 & 0 & \lambda_v \end{bmatrix}$$

where $k_{11} = r_1 + \delta_1 + \lambda_h$, $k_{22} = v_1 + \delta_1$, $k_{33} = v_1 + \lambda_h$, $k_{44} = a_1 b m$

The matrix Fv^{-1} is called the next generation matrix of the epidemiological model.

To find v^{-1} let $A = \begin{pmatrix} a & b & c \\ d & e & f \\ g & h & i \end{pmatrix}$. Then $\begin{pmatrix} a & b & c \\ d & e & f \\ g & h & i \end{pmatrix} \times \begin{bmatrix} k_{11} & 0 & 0 \\ k_{22} & k_{33} & k_{44} \\ 0 & 0 & \lambda_v \end{bmatrix} = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{pmatrix}$

$$\begin{pmatrix} ak11+bk22 & bk33 & bk44+chv \\ dk11+ek22 & ek33 & ek44+fhv \\ gk11+ek22 & hk33 & hk44+ihv \end{pmatrix} = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{pmatrix}$$

$$\Rightarrow \begin{cases} ak11+bk22=1 \\ bk33=0 \\ bk44+chv=0 \end{cases} \Rightarrow b=c=0 \text{ and } a = \frac{1}{k11}$$

$$\begin{cases} dk11+ek22=0 \\ ek33=1 \\ ek44+fhv=0 \end{cases} \Rightarrow d = \frac{-k22}{k11k33}, f = \frac{-k44}{k33hv} \text{ and } e = \frac{1}{k33}$$

$$\begin{cases} gk11+hk22=0 \\ hk33=0 \\ hk44+ihv=1 \end{cases} \Rightarrow h=g=0 \text{ and } i = \frac{1}{hv}$$

$$\Rightarrow v^{-1} = \begin{pmatrix} \frac{1}{k11} & 0 & 0 \\ -\frac{k22}{k11k33} & \frac{1}{k33} & \frac{-k44}{k33hv} \\ 0 & 0 & \frac{1}{hv} \end{pmatrix}$$

Hence $Fv^{-1} = \begin{bmatrix} 0 & 0 & a_1 b m s_{h1} \\ 0 & 0 & 0 \\ ca_1 & 0 & 0 \end{bmatrix} \times \begin{pmatrix} \frac{1}{k11} & 0 & 0 \\ -\frac{k22}{k11k33} & \frac{1}{k33} & \frac{-k44}{k33hv} \\ 0 & 0 & \frac{1}{hv} \end{pmatrix}$

$$= \begin{pmatrix} 0 & 0 & \frac{a_1 b m s_{h1}}{\lambda_v} \\ 0 & 0 & 0 \\ \frac{c a_1}{k_{11}} & 0 & 0 \end{pmatrix}$$

Then the spectral radius (the dominant eigen value) Fv^{-1} is equal to R_0 .

That is $|J - \lambda I| = 0$

$$\begin{vmatrix} -\lambda & 0 & \frac{a_1 b m}{\lambda_v} \\ 0 & -\lambda & 0 \\ \frac{c a_1}{k_{11}} & 0 & -\lambda \end{vmatrix} = 0$$

which forms the characteristics equation

$$\begin{aligned} \Rightarrow -\lambda^3 + \frac{a_1^2 b m c \lambda}{k_{11} \lambda_v} &= 0 \\ = -\lambda(\lambda^2 - \frac{a_1^2 b m c}{k_{11} \lambda_v}) &= 0 \\ \Rightarrow \lambda = 0 \text{ or } \lambda^2 = \frac{a_1^2 b m c}{k_{11} \lambda_v} \end{aligned}$$

Therefore the dominant Eigen values of FV^{-1} is $\lambda^2 = \frac{a_1^2 b m c}{k_{11} \lambda_v}$

$$\Rightarrow R_0^2 = \frac{a_1^2 b m c}{k_{11} \lambda_v}, \text{ then } R_0 = \sqrt{\frac{a_1^2 b m c}{k_{11} \lambda_v}} \quad (3.5.21)$$

This is the required reproduction number for the first age group.

From (3.5.21) if $a_1^2 b m c > k_{11} \lambda_v$ then $R_0 > 1$, this means that the equilibrium point is unstable (i.e the malaria disease is highly persistence to the population). If $a_1^2 b m c < k_{11} \lambda_v$ then $R_0 < 1$ and the equilibrium point is stable (i.e the disease is die out).

3.1.1 Equilibrium points and stability analysis

To determine the stability of the equilibrium points of the given model we evaluate the steady state of the given system with following conditions.

- Disease free equilibrium points (in the absence of infection) $i_{h1}, i_{v1} = 0$
- Endemic equilibrium points, that is $i_{h1}, i_{v1} \neq 0$

3.3.4. Local Stability analysis (sh1*,ih1*,iv1*)=(0,1,0)

To determine the stability of the system, we will consider linearizing the systems of equations (3.5.12)– (3.5.14) about the equilibrium points by taking the Jacobian of them.

The Jacobian matrix at any point $(i_{h1}^*, s_{h1}^*, i_v^*)$ is given by

$$J(i_{h1}^*, s_{h1}^*, i_v^*) = \begin{bmatrix} -(\lambda_h + r_1 + \delta_1) + 2\delta_1 i_{h1} & -a_1 b m i_v^* & a_1 b m s_{h1}^* \\ -v_1 + \delta_1 s_{h1}^* & -(\lambda_h + v_1 + a_1 b m i_v^* - \delta_1 i_{h1}) & -a_1 b m s_{h1}^* \\ a_1 c (1 - i_v) & 0 & -a_1 c i_{h1} - \lambda_v \end{bmatrix}$$

For the disease-free equilibrium, we evaluate the Jacobian matrix at the equilibrium points

$(i_{h1}^*, s_{h1}^*, i_v^*) = (0, 1, 0)$ and hence we obtain

$$J(0,1,0) = \begin{bmatrix} -(\lambda_h + r_1 + \delta_1) & 0 & a_1bm \\ -v_1 + \delta_1 & -(\lambda_h + v_1) & -a_1bm \\ a_1c & 0 & -\lambda_v \end{bmatrix} \quad (3.5.22)$$

The characteristic polynomial of (3.5.22) is

$$J(0,1,0) = \begin{bmatrix} -(\lambda_h + r_1 + \delta_1) - \lambda & 0 & a_1bm \\ -v_1 + \delta_1 & -(\lambda_h + v_1) - \lambda & -a_1bm \\ a_1c & 0 & -\lambda_v - \lambda \end{bmatrix} \quad (3.5.23)$$

Now applying $|J - \lambda I_3| = 0$ on (ii) we have

$$\begin{vmatrix} -(\lambda_h + r_1 + \delta_1) - \lambda & 0 & a_1bm \\ -v_1 + \delta_1 & -(\lambda_h + v_1) - \lambda & -a_1bm \\ a_1c & 0 & -\lambda_v - \lambda \end{vmatrix} = 0 \quad (3.5.24)$$

Solving for roots of the characteristic polynomial (with real coefficient) given in the Jacobian matrix leads to the characteristic equation

$$\lambda^3 + k_1\lambda^2 + k_2\lambda + k_3 = 0 \quad (3.5.25)$$

Where k_1 , k_2 , and k_3 are the coefficients given by the expressions:

$$\begin{aligned} k_3 &= -(\lambda_v + v_1)a_1^2bcm + \delta_1v_1\lambda_v + \lambda_h^2\lambda_v + r_1v_1\lambda_v + r_1\lambda_h\lambda_v + \delta_1\lambda_h\lambda_v + \lambda_vv_1\lambda_h \\ k_2 &= (\delta_1 + r_1 + \lambda_h)\lambda_v - a_1^2bcm + \delta_1\lambda_h + r_1v_1 + \lambda_h^2 + \delta_1v_1 + \lambda_vv_1 + \lambda_h\lambda_v + r_1\lambda_h + \lambda_h\lambda_v \\ &= \left(\frac{1}{R_0^2} - 1\right) + (\delta_1 + r_1 + \lambda_h)\lambda_h + (r_1 + \delta_1 + \lambda_v)v_1 + 2\lambda_h\lambda_v \end{aligned}$$

$$k_1 = v_1 + \lambda_v + 2\lambda_h + \delta_1 + r_1$$

Using Routh-Hurwitz criteria (Edelstein-Keshet 1988), the equilibrium is locally stable if the following conditions are satisfied.

The conditions are:

- (i) $k_1 > 0$
- (ii) $k_3 > 0$
- (iii) $k_1k_2 - k_3 > 0$

If these conditions are true, then all roots of the characteristic polynomial equation have negative real part, which concludes that there is a stable equilibrium. To show the three condition (i),(ii) and (iii) let:

$$(i) \quad k_1 > 0$$

Clearly $k_1 > 0$ since all parameters are positive.

$$(ii) \quad k_3 > 0$$

$$\begin{aligned} k_3 &= \delta_1v_1\lambda_v + \lambda_v\lambda_h^2 + r_1v_1\lambda_v + r_1\lambda_v\lambda_h + \delta_1\lambda_h\lambda_v + \lambda_hv_1\lambda_v - a_1^2bcm(\lambda_v + v_1) \\ &= ((\delta_1 + r_1 + \lambda_h)\lambda_v - a_1^2bcm)v_1 + (\delta_1 + r_1 + \lambda_h)\lambda_v\lambda_h - a_1^2bcm\lambda_h \\ &= \left(\frac{1}{R_0^2} - 1\right)v_1 + (\delta_1 + r_1 + \lambda_h)\lambda_v\lambda_h - a_1^2bcm\lambda_h \\ &= \left(\frac{1}{R_0^2} - 1\right)v_1 + \left(\frac{1}{R_0^2} - 1\right)\lambda_h \\ &= \left(\frac{1}{R_0^2} - 1\right)(\lambda_h + v_1) \end{aligned}$$

Clearly the disease free equilibrium point is locally asymptotically stable if $R_0^2 < 1$. If $R_0^2 > 1$ then, $k_3 < 0$. Hence, there is only one sign change in the sequence, k_1, k_2, k_3 and so by Descartes rule of sign, there is one eigenvalue with positive real part. This implies that the disease-free equilibrium point $(0, 1, 0)$ is unstable.

$$\Rightarrow k_3 > 0 \text{ if } R_0^2 < 1 \text{ and } k_3 < 0 \text{ if } R_0^2 > 1$$

$$(iii) k_1 k_2 - k_3 > 0$$

$$\Rightarrow \left(\frac{1}{R_0^2} - 1 \right) + (\delta_1 + r_1 + \lambda_h) \lambda_h + (r_1 + \delta_1 + \lambda_v) v_1 + 2\lambda_h \lambda_v \quad (v_1 + \lambda_v + 2\lambda_h + \delta_1 + r_1) - \left(\frac{1}{R_0^2} - 1 \right) (\lambda_h + v_1)$$

Hence the DFE point $(0, 1, 0)$ are unstable by Routh-Hurwitz stability criteria.

But if $R_0 < 1$ the DFE point $(0, 1, 0)$ are stable.

3.3.5. Existence and uniqueness of endemic equilibrium $(i_{h1}^*, s_{h1}^*, i_{v1}^*)$

Here, the condition for the existence and uniqueness of the endemic equilibrium of model system Eq. (3.5.12)-(3.5.14) is determined. Let $X^* = (i_{h1}^*, s_{h1}^*, i_{v1}^*)$ be the endemic equilibrium. To find the endemic equilibrium, we equate all equations in model system Eq. (3.5.12)-(3.5.14) to zero and rewrite it as follows:

$$\begin{aligned} a_1 b m s_{h1} i_v - (r_1 + \delta_1 + \lambda_h) i_{h1} + \delta_1 i_{h1}^2 &= 0 \\ \lambda_h (1 - s_{h1}) - a_1 b m s_{h1} i_v + v_1 (1 - s_{h1} - i_{h1}) + \delta_1 s_{h1} i_{h1} &= 0 \\ c a_1 (1 - i_{v1}) i_{h1} - \lambda_v i_{v1} &= 0 \end{aligned}$$

In the presence of infection $(i_{h1}, i_{v1} \neq 0)$, the mathematical model (3.5.12)-(3.5.14) has a unique endemic equilibrium given by $X^* = (i_{h1}^*, s_{h1}^*, i_{v1}^*)$ which are determined from the system as follows. The associated Jacobian matrix is obtained as:

$$J(i_{h1}^*, s_{h1}^*, i_{v1}^*) = \begin{bmatrix} -(\lambda_h + r_1 + \delta_1) + 2\delta_1 i_{h1}^* & -a_1 b m i_v^* & a_1 b m s_{h1}^* \\ -v_1 + \delta_1 s_{h1}^* & -(\lambda_h + v_1 + a_1 b m i_v^* - \delta_1 i_{h1}^*) & -a_1 b m s_{h1}^* \\ a_1 c (1 - i_{v1}^*) & 0 & -a_1 c i_{h1}^* - \lambda_v \end{bmatrix}$$

Then the Jacobian matrix of above matrix at EEP $(i_{h1}^*, s_{h1}^*, i_{v1}^*)$ is given by

$$J(i_{h1}^*, s_{h1}^*, i_{v1}^*) = \begin{bmatrix} -(\lambda_h + r_1 + \delta_1) + 2\delta_1 i_{h1}^* & -a_1 b m i_v^* & a_1 b m s_{h1}^* \\ -v_1 + \delta_1 s_{h1}^* & -(\lambda_h + v_1 + a_1 b m i_v^* - \delta_1 i_{h1}^*) & -a_1 b m s_{h1}^* \\ a_1 c (1 - i_{v1}^*) & 0 & -a_1 c i_{h1}^* - \lambda_v \end{bmatrix}$$

Obtaining the eigenvalues as usual from of the formula $|J(H^* - \lambda I)| = 0$

$$\Rightarrow \begin{vmatrix} -(\lambda_h + r_1 + \delta_1) + 2\delta_1 i_{h1}^* - \lambda & -a_1 b m i_v^* & a_1 b m s_{h1}^* \\ -v_1 + \delta_1 s_{h1}^* & -(\lambda_h + v_1 + a_1 b m i_v^* - \delta_1 i_{h1}^*) - \lambda & -a_1 b m s_{h1}^* \\ a_1 c (1 - i_{v1}^*) & 0 & -a_1 c i_{h1}^* - \lambda_v - \lambda \end{vmatrix} = 0$$

Since the model is not expressed explicitly then we use Matlab or Mathematica and the results discussed in chapter four:

3.3.6. Calculating Reproductive Number R_{01} for the second age group

To calculate the reproductive number of age group two we consider equation (3.5.15)-(3.5.17).

$$\begin{aligned} a_2 b m s_{h2} i_v - (r_2 + \delta_2 - \varepsilon_1) i_{h2} - \varepsilon_1 i_{h1} + \delta_2 i_{h2}^2 &= 0 \text{ (i)} \\ -a_2 b m s_{h2} i_{v2} - \varepsilon_1 s_{h1} + \delta_2 s_{h2} i_{h2} - \varepsilon_2 s_{h2} + v_2 (1 - s_{h2} - i_{h2}) &= 0 \text{ (ii)} \\ c a_2 (1 - i_{v2}) i_{h2} - \lambda_v i_{v2} &= 0 \text{ (iii)} \end{aligned}$$

In the absence of infection ($i_{h2}=i_{v2}=0$) equation (i)-(iii) becomes

$$\begin{aligned} \mathcal{E}_1 S_{h1} - \mathcal{E}_1 S_{h2} + \nu_2 (1 - S_{h2}) &= 0 \\ \Rightarrow \mathcal{E}_1 S_{h1} - S_{h2} (\mathcal{E}_1 + \nu_2) + \nu_2 &= 0 \\ \Rightarrow S_{h2} &= \frac{\mathcal{E}_1 S_{h1} + \nu_2}{\nu_2 + \mathcal{E}_1} \\ &= \frac{\mathcal{E}_1 + \nu_2}{\nu_2 + \mathcal{E}_1} = 1 \end{aligned}$$

To find reproduction number R_0 for the second age group will refer to the ODEs (3.5.15)-(3.5.17) it is important to distinguish new infections from all other changes in population.

Gains terms

$$F_1 = a_2 b m i_v \quad F_2 = 0 \quad F_3 = c a_2 (1 - i_v) i_{h2}$$

Loss terms

$$\begin{aligned} \nu_1 &= (r_2 + \delta_2 - \mathcal{E}_1) i_{h2} + \delta_1 i_{h2}^2 - \mathcal{E}_1 i_{h1} \\ \nu_2 &= a_2 b m s_{h2} i_v + \nu_2 (1 - s_{h2} - i_{h2}) + \delta_2 s_{h2} i_{h2} + \mathcal{E}_1 s_{h1} + \mathcal{E}_1 s_{h2} \\ \nu_3 &= \lambda_v i_v \end{aligned}$$

Then the Jacobian matrix of F (gain terms) is given by

$$F(i_{h2}, s_{h2}, i_v) = \begin{bmatrix} 0 & a_2 b m i_v & a_2 b m s_{h2} \\ 0 & 0 & 0 \\ a_2 c (1 - i_v) & 0 & -a_2 c i_{h2} \end{bmatrix}$$

Then the Jacobian of F at equilibrium point $E_0 = (i_{h2}^*, s_{h2}^*, i_{v2}^*) = (0, 1, 0)$ is given by

$$F(0, 1, 0) = \begin{bmatrix} 0 & 0 & a_2 b m \\ 0 & 0 & 0 \\ c a_2 & 0 & 0 \end{bmatrix}$$

And the Jacobian matrix of v (loss terms) is given by

$$V(i_{h2}, s_{h2}, i_v) = \begin{bmatrix} w_1 & 0 & 0 \\ w_2 & w_3 & w_4 \\ 0 & 0 & \lambda_v \end{bmatrix}$$

$$\text{where } w_1 = \delta_2 + r_2 - \mathcal{E}_1 + 2\delta_2 i_{h2}$$

$$w_2 = \nu_2 + \delta_2 s_{h2}$$

$$w_3 = a_2 b m i_v + \nu_2 + \delta_2 i_{h2}$$

$$w_4 = a_2 b m s_{h2}$$

Then the Jacobian matrix of loss terms v at DFE (0, 1, 0) is given by

$$V(0, 1, 0) = \begin{bmatrix} w_{11} & 0 & 0 \\ w_{22} & w_{33} & w_{44} \\ 0 & 0 & \lambda_v \end{bmatrix}$$

$$\text{Where: } w_{11} = \delta_2 + r_2 - \mathcal{E}_1 \ \& \ w_{22} = \nu_2 + \delta_2, \ w_{33} = \nu_2 + \mathcal{E}_1, \ w_{44} = a_2 b m$$

The matrix FV^{-1} is called the next generation matrix of the epidemiological model.

$$\begin{aligned} &\text{To find } v^{-1} \text{ let } A = \begin{pmatrix} a & b & c \\ d & e & f \\ g & h & i \end{pmatrix}. \text{ Then } v \times v^{-1} = I \\ &\Rightarrow \begin{pmatrix} a & b & c \\ d & e & f \\ g & h & i \end{pmatrix} \times \begin{bmatrix} w_{11} & 0 & 0 \\ w_{22} & w_{33} & w_{44} \\ 0 & 0 & \lambda_v \end{bmatrix} = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{pmatrix} \\ &= \begin{pmatrix} aw_{11} + bw_{22} & bw_{33} & bw_{44} + chv \\ dw_{11} + ew_{22} & ew_{33} & ew_{44} + fhv \\ gw_{11} + hw_{22} & hw_{33} & hw_{44} + ihv \end{pmatrix} = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{pmatrix} \\ &\Rightarrow \begin{cases} aw_{11} + bw_{22} = 1 \\ bw_{33} = 0 \\ bw_{44} + chv = 0 \end{cases} \Rightarrow b = c = 0 \text{ and } a = \frac{1}{w_{11}} \\ &\begin{cases} dw_{11} + ew_{22} = 0 \\ ew_{33} = 1 \\ ew_{44} + fhv = 0 \end{cases} \Rightarrow d = \frac{-w_{22}}{w_{11}w_{33}}, f = \frac{-w_{44}}{w_{33}hv} \text{ and } e = \frac{1}{w_{33}} \\ &\begin{cases} gw_{11} + hw_{22} = 0 \\ hw_{33} = 0 \\ hw_{44} + ihv = 1 \end{cases} \Rightarrow h = g = 0 \text{ and } i = \frac{1}{hv} \\ &\Rightarrow v^{-1} = \begin{pmatrix} \frac{1}{w_{11}} & 0 & 0 \\ -\frac{w_{22}}{w_{11}w_{33}} & \frac{1}{w_{33}} & \frac{-w_{44}}{w_{33}hv} \\ 0 & 0 & \frac{1}{hv} \end{pmatrix} \\ &\text{Hence } Fv^{-1} = \begin{bmatrix} 0 & 0 & a_2 b m S_{h2} \\ 0 & 0 & 0 \\ c a_2 & 0 & 0 \end{bmatrix} \times \begin{pmatrix} \frac{1}{w_{11}} & 0 & 0 \\ -\frac{w_{22}}{w_{11}w_{33}} & \frac{1}{w_{33}} & \frac{-w_{44}}{w_{33}hv} \\ 0 & 0 & \frac{1}{hv} \end{pmatrix} \\ &= \begin{pmatrix} 0 & 0 & \frac{a_2 b m}{hv} \\ 0 & 0 & 0 \\ \frac{c a_2}{w_{11}} & 0 & 0 \end{pmatrix} \end{aligned}$$

Then the spectral radius (the dominant Eigen value) Fv^{-1} is equal to R_0 .

$$\text{That is } |J - \lambda I| = 0$$

$$\Rightarrow \begin{vmatrix} -\lambda & 0 & \frac{a_2bm}{\lambda_v} \\ 0 & -\lambda & 0 \\ \frac{ca_2}{w_{11}} & 0 & -\lambda \end{vmatrix} = 0$$

$$\Rightarrow -\lambda^3 + \frac{a_2^2bcm}{w_{11}\lambda_v} \lambda = 0$$

$$= -\lambda \left(\lambda^2 - \frac{a_2^2bcm}{w_{11}\lambda_v} \right) = 0$$

$$\Rightarrow \lambda = 0 \text{ or } \lambda^2 = \frac{a_2^2bcm}{w_{11}\lambda_v} = \frac{a_2^2bcm}{(\delta_2 + r_2 - \varepsilon_1)\lambda_v}$$

Therefore the dominant Eigen values of Fv^{-1} is $\lambda^2 = \frac{a_2^2bcm}{(\delta_2 + r_2 - \varepsilon_1)\lambda_v}$

$$\Rightarrow R_{01}^2 = \frac{a_2^2bcm}{w_{11}\lambda_v}, \text{ then } R_{01} = \sqrt{\frac{a_2^2bcm}{w_{11}\lambda_v}} (*)$$

3.3.7. Local Stability analysis (ih2*,sh2*,iv2*) = (0,1,0)

To determine the stability of the system, we will consider linearizing the systems of equations (3.5.14)– (3.5.17) about the equilibrium points by taking the Jacobian of them.

The Jacobian matrix at any point (ih2*,sh2*,iv2*) is given by

$$J(i_{h2}^*, S_{h2}^*, i_v^*) = \begin{bmatrix} -(r_2 + \delta_2 - \varepsilon_1) + 2\delta_2 i_{h2} & a_2 b m i_v & a_2 b m S_{h2} \\ -v_2 + \delta_2 S_{h2} & -a_2 b m i_v + \delta_2 i_{h2} - \varepsilon_1 - v_2 & -a_2 b m S_{h2} \\ a_2 c (1 - i_v) & 0 & -a_2 c i_{h2} - \lambda_v \end{bmatrix}$$

Then the Jacobian matrix of the above matrix at DFE point (0,1,0) is given by

$$J(0,1,0) = \begin{bmatrix} -(r_2 + \delta_2 - \varepsilon_1) & 0 & a_2 b m \\ -v_2 + \delta_2 & -\varepsilon_1 - v_2 & -a_2 b m \\ a_2 c & 0 & -\lambda_v \end{bmatrix} \text{ (ii)}$$

Now applying $|J - \lambda I_3| = 0$

$$\Rightarrow \begin{bmatrix} -(r_2 + \delta_2 - \varepsilon_1) - \lambda & 0 & a_2 b m \\ -v_2 + \delta_2 & -\varepsilon_1 - v_2 - \lambda & -a_2 b m \\ a_2 c & 0 & -\lambda_v - \lambda \end{bmatrix} \text{ (iii)}$$

The corresponding characteristic equation of (iii) is

$$\lambda^3 + b_1 \lambda^2 + b_2 \lambda + b_3 = 0$$

Where b_1 , b_2 , and b_3 are the coefficients given by the expressions:

$$b_3 = \delta_2 \varepsilon_1 \lambda_v + r_2 v_2 \lambda_v + \varepsilon_1^2 \lambda_v + a_2^2 b c m (v_2 + \varepsilon_2) + r_2 \varepsilon_1 \lambda_v + \delta_2 v_2 \lambda_v + \varepsilon_1 v_2 \lambda_v$$

$$b_2 = \left(\frac{v_2}{R_{01}^2 \lambda_v} - 1 \right) + (\delta_2 - \varepsilon_1 + r_2) \varepsilon_1 + (r_2 + v_2) \lambda_v$$

$$b_1 = v_2 + r_2 + \lambda_v + \delta_2$$

Using Routh-Hurwitz criteria (Edelstein-Keshet 1988), the equilibrium is locally stable if the following conditions are satisfied. If these conditions are true, then all roots of the characteristic polynomial equation have negative real part, which concludes that there is a stable equilibrium.

The conditions are:

- (i) $b_1 > 0$
- (ii) $b_3 > 0$ and
- (iii) $b_1 b_2 - b_3 > 0$ must hold.

To show (i), (ii), and (iii) we use

- (i) $b_1 > 0$

Clearly $b_1 > 0$ since all parameters are positive.

- (ii) $b_3 > 0$

$$\begin{aligned}
 b_3 &= \delta_2 \varepsilon_1 \lambda_v + r_2 v_2 \lambda_v - \varepsilon_1^2 \lambda_v - a_2^2 b c m (v_2 + \varepsilon_1) + r_2 \varepsilon_1 \lambda_v + \delta_2 \varepsilon_1 \lambda_v - \varepsilon_1 v_2 \lambda_v \\
 &= (\delta_2 + r_2 - \varepsilon_1) v_2 \lambda_v - a_2^2 b c m v_2 - a_2^2 b c m \varepsilon_1 + r_2 \varepsilon_1 \lambda_v + \delta_2 \varepsilon_1 \lambda_v - \varepsilon_1^2 \lambda_v \\
 &= ((\delta_2 + r_2 - \varepsilon_1) \lambda_v - a_2^2 b c m) v_2 - a_2^2 b c m \varepsilon_1 + r_2 \varepsilon_1 \lambda_v + \delta_2 \varepsilon_1 \lambda_v - \varepsilon_1^2 \lambda_v \\
 &= (r_2 + \delta_2 - \varepsilon_1) \lambda_v v_2 - a_2^2 b c m v_2 + (\delta_2 + r_2 - \varepsilon_1) \lambda_v \varepsilon_1 - a_2^2 b c m \varepsilon_1 \\
 &= \left(\frac{1}{R_{01}^2} - 1 \right) v_2 + (\delta_2 + r_2 - \varepsilon_1) \lambda_v \varepsilon_1 - a_2^2 b c m \varepsilon_1 \\
 &= \left(\frac{1}{R_{01}^2} - 1 \right) v_2 + \left(\frac{1}{R_{01}^2} - 1 \right) \varepsilon_1 \\
 &= \left(\frac{1}{R_{01}^2} - 1 \right) (\varepsilon_1 + v_2)
 \end{aligned}$$

Clearly we see that the disease free equilibrium point is locally asymptotically stable if $R_{01}^2 < 1$. If $R_{01}^2 > 1$ then $b_3 < 0$. Hence, there is only one sign change in the sequence, b_1, b_2, b_3 and so by Descartes rule of sign, there is one eigenvalue with positive real part. This implies that the disease-free equilibrium point $(0, I, 0)$ is unstable. (i.e Since the second condition failed then the DFE point unstable). Finally, we show that $b_1 b_2 - b_3 > 0$. The expression for b_2 is

$$b_2 = \left(\frac{v_2}{R_{01}^2 \lambda_v} - 1 \right) + (\delta_2 - \varepsilon_1 + r_2) \varepsilon_1 + (r_2 + v_2) \lambda_v$$

- (iii) $b_1 b_2 - b_3 > 0$

$$\left(\frac{v_2}{R_{01}^2 \lambda_v} - 1 \right) + (\delta_2 - \varepsilon_1 + r_2) \varepsilon_1 + (r_2 + v_2) \lambda_v (r_2 + v_2 + \lambda_v + \delta_2) - \left(\frac{1}{R_{01}^2} - 1 \right) (\varepsilon_1 + v_2)$$

3.3.8. Calculating Reproductive Number R_{02} for the third age group

$$\frac{di_{h3}}{dt} = a_3 b m S_{h3} i_{v2} - (r_3 + \delta_3 + \varepsilon_2) i_{h3} + \varepsilon_2 i_{h2} + \delta_3 i_{h3}^2$$

$$\frac{ds_{h3}}{dt} = -a_3 b m S_{h3} i_{v3} - \varepsilon_2 S_{h3} + \varepsilon_2 S_{h2} + \varepsilon_2 S_{h2} + v_3 (1 - S_{h3} - i_{h3}) + \delta_3 S_{h3} i_{h3}$$

$$\frac{di_{v1}}{dt} = c a_3 (1 - i_{v3}) i_{h3} - \lambda_v i_{v3}$$

To determine the reproduction number it is important to distinguish new infections from all other changes in population. Forming the matrices for F and V at the disease free equilibrium of the form $E_0 = (i_{h3}^*, s_{h3}^*, i_{v3}^*) = (0, 1, 0)$ such that there is no infection in the population,

Gains terms

$$F_1 = a_3 b m s_{h3} i_{v3} \quad F_2 = 0 \quad F_3 = c a_3 (1 - i_{v3}) i_{h3}$$

Loss terms

$$\begin{aligned} v_1 &= (r_3 + \delta_3 + \varepsilon_2) i_{h3} + \delta_1 i_{h2}^2 + \varepsilon_2 i_{h2} + \delta_3 i_{h3}^2 \\ v_2 &= a_3 b m s_{h3} i_{v3} + v_3 (1 - s_{h3} - i_{h3}) + \delta_3 s_{h3} i_{h3} + \varepsilon_2 s_{h3} + \varepsilon_2 s_{h2} \\ v_3 &= \lambda_v i_{v3} \end{aligned}$$

Then the Jacobian matrix of F (gain terms) is given by

$$F(i_{h3}, s_{h3}, i_{v3}) = \begin{bmatrix} 0 & a_3 b m i_{v3} & a_3 b m s_{h3} \\ 0 & 0 & 0 \\ c a_3 (1 - i_{v3}) & 0 & -a_3 c i_{h3} \end{bmatrix}$$

And the Jacobian matrix of F (loss terms) is given by

$$v(i_{h3}^*, s_{h3}^*, i_{v3}^*) = \begin{bmatrix} m_1 & 0 & 0 \\ m_2 & m_3 & m_4 \\ 0 & 0 & \lambda_v \end{bmatrix}$$

Where

$$\begin{aligned} m_1 &= \delta_3 + r_3 + \varepsilon_2 + 2\delta_3 i_{h3} \\ m_{12} &= v_3 + \delta_3 s_{h3} \\ m_3 &= \varepsilon_2 + v_3 + a_3 b m i_{v3} + \delta_3 i_{h3} \\ m_4 &= a_3 b m s_{h3} \end{aligned}$$

Then the Jacobian matrix at DFE point of (0,1,0) is given by

$$V(0,1,0) = \begin{bmatrix} m_{11} & 0 & 0 \\ m_{12} & m_{13} & m_{14} \\ 0 & 0 & \lambda_v \end{bmatrix}$$

Where

$$\begin{aligned} m_{11} &= \delta_3 + r_3 + \varepsilon_2 \\ m_{12} &= v_3 + \delta_3 \\ m_{13} &= \varepsilon_2 + v_3 \\ m_{14} &= a_3 b m \end{aligned}$$

Now to find v^{-1} let $v^{-1} = \begin{pmatrix} a & b & c \\ d & e & f \\ g & h & i \end{pmatrix}$. Then $v \times v^{-1} = I_3$

$$\Rightarrow \begin{pmatrix} a & b & c \\ d & e & f \\ g & h & i \end{pmatrix} \times \begin{bmatrix} m_{11} & 0 & 0 \\ m_{12} & m_{13} & m_{14} \\ 0 & 0 & \lambda_v \end{bmatrix} = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{pmatrix}$$

$$\Rightarrow \begin{pmatrix} a m_{11} + b m_{22} & b m_{33} & b m_{44} + c \lambda_v \\ d m_{11} + e m_{22} & e m_{33} & e m_{44} + f \lambda_v \\ g m_{11} + h m_{22} & h m_{33} & h m_{44} + i \lambda_v \end{pmatrix} = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{pmatrix}$$

$$\begin{cases} am_{11} + bm_{22} = 1 \\ bm_{33} = 0 \\ bm_{44} + c\lambda_v = 0 \end{cases} \Rightarrow b = c = 0 \quad \text{and} \quad a = \frac{1}{m_{11}}$$

$$\begin{cases} dm_{11} + em_{22} = 0 \\ em_{33} = 1 \\ em_{44} + f\lambda_v = 0 \end{cases} \Rightarrow d = \frac{-m_{22}}{m_{11}m_{22}} \quad f = \frac{-m_{44}}{m_{33}\lambda_v} \quad \text{and} \quad e = \frac{1}{m_{33}}$$

$$\begin{cases} gm_{11} + hm_{22} = 0 \\ hm_{33} = 0 \\ hm_{44} + i\lambda_v = 1 \end{cases} \Rightarrow h = g = 0 \quad \text{and} \quad i = \frac{1}{\lambda_v}$$

$$\Rightarrow v^{-1} = \begin{pmatrix} \frac{1}{m_{11}} & 0 & 0 \\ -\frac{w_{22}}{m_{11}m_{33}} & \frac{1}{m_{33}} & \frac{-m_{44}}{m_{33}\lambda_v} \\ 0 & 0 & \frac{1}{\lambda_v} \end{pmatrix}$$

$$\text{Hence } Fv^{-1} = \begin{bmatrix} 0 & 0 & a_3 b m_{h3} \\ 0 & 0 & 0 \\ ca_3 & 0 & 0 \end{bmatrix} \times \begin{pmatrix} \frac{1}{m_{11}} & 0 & 0 \\ -\frac{w_{22}}{m_{11}m_{33}} & \frac{1}{m_{33}} & \frac{-m_{44}}{m_{33}\lambda_v} \\ 0 & 0 & \frac{1}{\lambda_v} \end{pmatrix}$$

$$= \begin{pmatrix} 0 & 0 & \frac{a_3 b m}{\lambda_v} \\ 0 & 0 & 0 \\ \frac{ca_3}{m_{11}} & 0 & 0 \end{pmatrix}$$

Then the spectral radius (the dominant eigen value) Fv^{-1} is equal to R_0 .

Obtaining the eigenvalues as usual form of the formula $|J - \lambda I| = 0$

$$\Rightarrow \begin{vmatrix} -\lambda & 0 & \frac{a_3 b m}{\lambda_v} \\ 0 & -\lambda & 0 \\ \frac{ca_3}{m_{11}} & 0 & -\lambda \end{vmatrix} = 0$$

$$\Rightarrow -\lambda^3 + \frac{a_3^2 b c m}{m_{11} \lambda_v} \lambda = 0$$

$$= -\lambda \left(\lambda^2 - \frac{a_3^2 b c m}{m_{11} \lambda_v} \right) = 0$$

$$\Rightarrow \lambda = 0 \text{ or } \lambda^2 = \frac{a_3^2bcm}{m_{11}\lambda_v}$$

Therefore the dominant Eigen values of Fv^{-1} is $\lambda^2 = \frac{a_3^2bcm}{m_{11}\lambda_v}$

$$\Rightarrow R_{02}^2 = \frac{a_3^2bcm}{m_{11}\lambda_v}, \text{ then } R_{02} = \sqrt{\frac{a_3^2bcm}{m_{11}\lambda_v}}$$

Clearly if $a_3^2bcm > m_{11}\lambda_v$, then the reproduction number is $R_{02}^2 > 1$ and the disease is highly affect the community. If $a_3^2bcm < m_{11}\lambda_v$, then the reproduction number is $R_{02}^2 < 1$ and the malaria disease is die out. In the context of epidemiology modeling, it is generally known that if $R_0 < 1$, then the disease free equilibrium is locally asymptotically stable and the disease will be eradicated from the community.

3.4. Local Stability analysis of DFE point $(ih_3^*, sh_3^*, iv^*) = (0, 1, 0)$

To determine the stability of the system, we will consider linearizing the systems of equations (3.5.18)– (3.5.20) about the equilibrium points by taking the Jacobian of them.

The Jacobian matrix at any point (ih_3^*, sh_3^*, iv^*) is given by

$$J(i_{h_3}^*, s_{h_3}^*, i_v^*) = \begin{bmatrix} -(r_3 + \delta_3 + \varepsilon_2) + 2\delta_3 i_{h_3} & a_3 b m i_v & a_3 b m s_{h_3} \\ -v_3 + \delta_3 S_{h_3} & -a_3 b m i_v + \delta_3 i_{h_3} - v_3 - \varepsilon_2 & -a_3 b m s_{h_3} \\ c a_3 (1 - i_v) & 0 & -c a_3 i_{h_3} - \lambda_v \end{bmatrix}$$

Then the Jacobian matrix of the above matrix at DFE point $(0, 1, 0)$ is given by

$$J(0, 1, 0) = \begin{bmatrix} -(r_3 + \delta_3 + \varepsilon_2) & 0 & a_3 b m \\ -v_3 + \delta_3 & -(v_3 + \varepsilon_2) & -a_3 b m \\ c a_3 & 0 & -\lambda_v \end{bmatrix} \text{ (ii)}$$

The corresponding characteristic polynomial is

$$\begin{vmatrix} -(r_3 + \delta_3 + \varepsilon_2) - \lambda & 0 & a_3 b m \\ -v_3 + \delta_3 & -(v_3 + \varepsilon_2) - \lambda & -a_3 b m \\ c a_3 & 0 & -\lambda_v - \lambda \end{vmatrix} = 0 \text{ (iii)}$$

The corresponding characteristic equation (iii) is given by

$$\lambda^3 + d_1 \lambda^2 + d_2 \lambda + d_3 = 0$$

Where d_1, d_2 and d_3 are the coefficients given by the expressions:

$$d_3 = -a_3^2bcm\varepsilon_2 - a_3^2bcmv_3 + \delta_3v_3\lambda_v + \varepsilon_2^2\lambda_v + r_3v_3\lambda_v + r_3\varepsilon_2\lambda_v + \delta_3\varepsilon_2\lambda_v + v_3\varepsilon_2\lambda_v$$

$$d_2 = \left(\frac{1}{R_{02}^2} - 1 \right) + \varepsilon_2v_3 + \varepsilon_2^2 + r_3\varepsilon_2 + r_3v_3 + \lambda_vv_3 + \delta_3\varepsilon_2 + \lambda_v\varepsilon_2 + \delta_3v_3$$

$$d_1 = v_3 + \lambda_v + \delta_3 + r_3 + 2\varepsilon_2$$

Using Routh-Hurwitz criteria (Edelstein-Keshet 1988), the equilibrium is locally stable if the following conditions are satisfied. If these conditions are true, then all roots of the characteristic polynomial equation have negative real part, which concludes that there is a stable equilibrium.

The three conditions are:

$$(i) \quad d_1 > 0$$

$$(ii) \quad d_3 > 0$$

$$(iii) \quad d_1d_2 - d_3 > 0 \text{ must hold.}$$

To show (i), (ii), and (iii) we use

$$(i) \quad d_1 = v_3 + \lambda_v + \delta_3 + r_3 + 2\varepsilon_2$$

Clearly $d_1 > 0$ since all parameters have positive values. Now to check the second condition we check for

$$d_3 > 0$$

$$\begin{aligned} d_3 &= -a_3^2bcm\varepsilon_2 - a_3^2bcmv_3 + \delta_3v_3\lambda_v + \varepsilon_2^2\lambda_v + r_3v_3\lambda_v + r_3\varepsilon_2\lambda_v + \delta_3\varepsilon_2\lambda_v + v_3\varepsilon_2\lambda_v \\ &= \left((\varepsilon_2 + r_3 + \delta_3)\lambda_v - a_3^2bcm \right) v_3 + \varepsilon_2^2\lambda_v + r_3\varepsilon_2\lambda_v + \delta_3\varepsilon_2\lambda_v - a_3^2bcm\varepsilon_2 \\ &= \left(\frac{1}{R_{02}^2} - 1 \right) v_3 + \varepsilon_2^2\lambda_v + r_3\varepsilon_2\lambda_v + \delta_3\varepsilon_2\lambda_v - a_3^2bcm\varepsilon_2 \\ &= \left(\frac{1}{R_{02}^2} - 1 \right) v_3 + (\varepsilon_2 + r_3 + \delta_3)\lambda_v\varepsilon_2 - a_3^2bcm\varepsilon_2 \\ &= \left(\frac{1}{R_{02}^2} - 1 \right) v_3 + ((\varepsilon_2 + r_3 + \delta_3)\lambda_v - a_3^2bcm)\varepsilon_2 \\ &= \left(\frac{1}{R_{02}^2} - 1 \right) v_3 + \left(\frac{1}{R_{02}^2} - 1 \right) \varepsilon_2 \\ &= \left(\frac{1}{R_{02}^2} - 1 \right) (v_3 + \varepsilon_2) \end{aligned}$$

So we have seen that the disease free equilibrium point is locally asymptotically stable if $R_{02} < 1$. If $R_{02} > 1$, we can see that $d_3 < 0$. Hence, there is only one sign change in the sequence, d_1, d_2, d_3 and so by Descartes rule of sign, there is one eigenvalue with positive real part. This implies that the disease-free equilibrium point $(0, l, 0)$ is unstable. (i.e Since the second condition failed then the DFE point unstable)

Finally, we show that $d_1d_2 - d_3 > 0$. The expression for d_2 is

$$d_2 = \left(\frac{1}{R_{02}^2} - 1 \right) + \varepsilon_2v_3 + \varepsilon_2^2 + r_3\varepsilon_2 + r_3v_3 + \lambda_vv_3 + \delta_3\varepsilon_2 + \lambda_v\varepsilon_2 + \delta_3v_3$$

Hence we get

$$d_1d_2 - d_3 =$$

$$\begin{aligned} & \left[v_3 + \lambda_v + \delta_3 + r_3 + 2\varepsilon_2 \right] \left[\left(\frac{1}{R_{02}^2} - 1 \right) + \varepsilon_2v_3 + \varepsilon_2^2 + r_3\varepsilon_2 + r_3v_3 + \lambda_vv_3 + \delta_3\varepsilon_2 + \lambda_v\varepsilon_2 + \delta_3v_3 \right] \\ & - \left(\frac{1}{R_{02}^2} - 1 \right) (v_3 + \varepsilon_2) \end{aligned}$$

Thus the third condition $d_1d_2 - d_3 > 0$ also satisfied if and only if $R_{02} < 1$. If $R_{02} > 1$, then $d_1d_2 - d_3 < 0$.

3.4.1. Local Stability analysis of endemic equilibrium points for the age group three

The endemic equilibrium state is the state where the disease cannot be totally eradicated but remains in the population.

For the disease to persist in the population, the susceptible class, the Infectious class and the Recovered Class must not be zero at equilibrium state (i.e $i_{h1}, i_v \neq 0$).

In other words, if E_1 is the endemic equilibrium state, then in (i.e. equilibrium where the infected components of the age-structured model (3.5.18) – (3.5.20) are non-zero or where the disease persist in the population (all state variables are positive) will be derived. When the disease free equilibrium point is unstable, then there exists an endemic equilibrium point

The endemic equilibrium solution of (3.5.18)-(3.5.20) can be found by setting the right hand side to zero and solving the nonlinear system to obtain

$$\frac{di_{h3}}{dt} = a_3 b m s_{h3} i_{v2} - (r_3 + \delta_3 + \epsilon_2) i_{h3} + \epsilon_2 i_{h2} + \delta_3 i_{h3}^2 (*)$$

$$\frac{ds_{h3}}{dt} = -a_3 b m s_{h3} i_{v3} - \epsilon_2 s_{h3} + \epsilon_2 s_{h2} + \epsilon_2 s_{h2} + v_3 (1 - s_{h3} - i_{h3}) + \delta_3 s_{h3} i_{h3} (**)$$

$$\frac{di_{v1}}{dt} = c a_3 (1 - i_{v3}) i_{h3} - \lambda_v i_{v3} (***)$$

Now from (***) we have that $i_v = \frac{a_3 c i_{h3}}{a_3 c i_{h3} + \lambda_v}$ (i)

Substitute (i) in (*) and (**) solve for i_{h3} .

Then the Jacobian matrix of age group one model at endemic steady state $(i_{h3}^*, s_{h3}^*, i_v^*)$ given by:

$$J(i_{h3}^*, s_{h3}^*, i_v^*) = \begin{bmatrix} -(r_3 + \delta_3 + \epsilon_2) + 2\delta_3 i_{h3}^* & a_3 b m i_v^* & a_3 b m s_{h3}^* \\ -v_3 + \delta_3 s_{h3}^* & -a_3 b m i_v^* + \delta_3 i_{h3}^* - v_3 - \epsilon_2 & -a_3 b m s_{h3}^* \\ c a_3 (1 - i_v^*) & 0 & -c a_3 i_{h3}^* - \lambda_v \end{bmatrix}$$

Then see the EEP of stability for this model in chapter four.

NUMERICAL SIMULATION

We present numerical simulations using Matlab to illustrate the transmission of malaria disease for the three age group described above. The parameter values are given in Table 2.

4.1. Numerical simulation of the first age group model

Consider the model given by equation (3.5.12)-(3.5.14)

$$\frac{di_{h1}}{dt} = a b m s_{h1} i_v - (r_1 + \delta_1 + \lambda_h) i_{h1} + \delta_1 i_{h1}^2 (a)$$

$$\frac{ds_{h1}}{dt} = \lambda_h (1 - s_{h1}) - a b m s_{h1} i_v + v_1 (1 - s_{h1} - i_{h1}) + \delta_1 s_{h1} i_{h1} (b)$$

$$\frac{di_v}{dt} = c a_1 (1 - i_v) i_{h1} - \lambda_v i_v (c)$$

4.2. Reproduction number of age group one

The reproduction number measures the average number of new infected population by a single infected population. Since we have shown an explicit expression for R_0 , we can analytically evaluate its sensitivity to the different parameter values.

We can also numerically evaluate the sensitivity of the endemic equilibrium to the parameter values, as this will allow us to determine the relative importance to the age-structured malaria transmission and prevalence.

$$R_0^2 = \frac{a_1^2 b c m}{k_1 l_1 h v} \text{ or } R_0 = \sqrt{\frac{a_1^2 b m c}{k_1 l_1 h v}}$$

$$\Rightarrow R_0 = \sqrt{\frac{(0.46)^2 (0.5) (0.48) (0.2)}{(0.071) (0.00012 + 0.0014 + 0.00000214)}} = 9.6944 > 1$$

Since $R_0 > 1$, the prevalence of malaria disease will result in an epidemic. This is due to the fact that the rate of transmission is greater than the recovery rate.

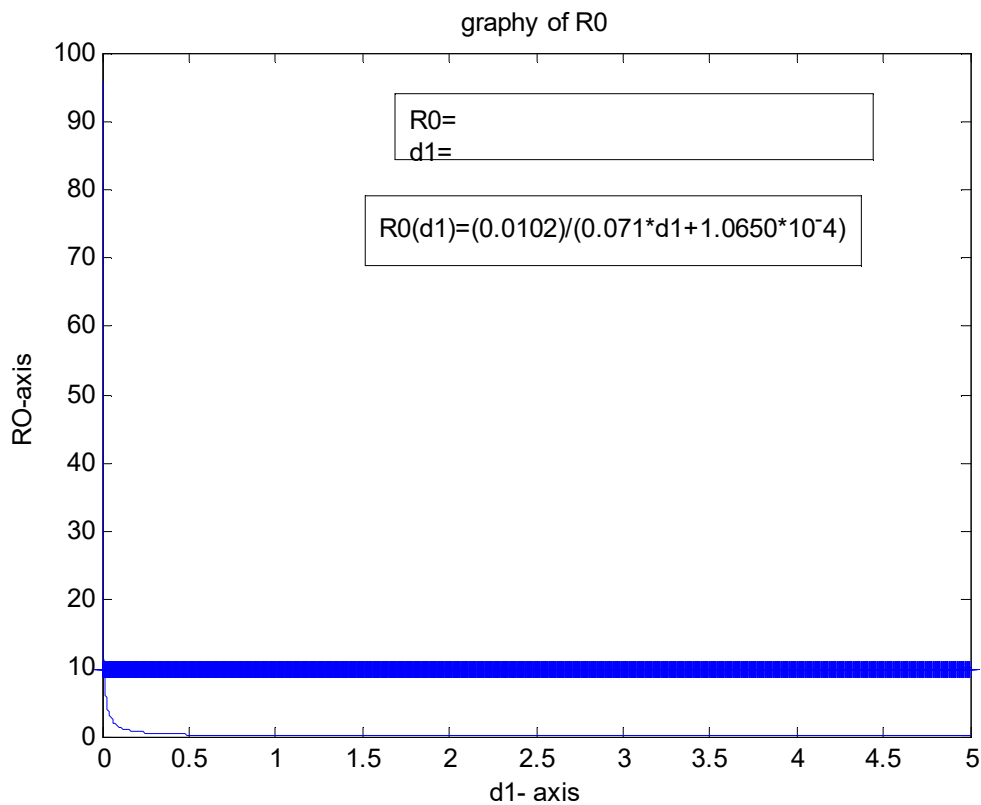


Figure 5.1 graph of the basic reproduction number R_0

From this graph we can observe that if the Per capita death rate δ_1 increases and λ_h, r_1 is constant then the reproduction number R_0 decreases. This means that when the Per capita death rate δ_1 decrease then the reproduction number is increasing and the susceptible population increasing. And when the Per capita death rate δ_1 increases R_0 decreases that means when the Per capita death rate increase then the number of infected population decrease since individual in the infected class dies out. Similarly if λ_h, δ_1 are constant and r_1 increases, then the reproduction number R_0 decreases and as r_1 decreases R_0 increases.

4.3. Stability analysis of DFE point of first age group

The Jacobian matrix of the given model at DFE point $(0,sh1,0)$ is given as follows

$$J(0,sh1,0) = \begin{bmatrix} -(\delta_1 + r_1 + \lambda_h) & 0 & a_1 b m s_{h1} \\ -v_1 + \delta_1 s_{h1} & 0 & -a_1 b m s_{h1} \\ c a_1 & 0 & -\lambda_v \end{bmatrix}$$

And the Jacobian matrix at DFE point $(0,1,0)$ is

$$J(0,1,0) = \begin{bmatrix} -0.0015 & 0 & 0.0460 \\ -0.0083 & -0.0084 & -0.0460 \\ -0.2208 & 0 & -0.071 \end{bmatrix}$$

The corresponding characteristic polynomial is $|J - \lambda I_3| = 0$

$$\Rightarrow \begin{vmatrix} -0.0015 - \lambda & 0 & 0.0460 \\ -0.0083 & -0.0084 - \lambda & -0.0460 \\ -0.2208 & 0 & -0.071 - \lambda \end{vmatrix} = 0 (*)$$

The characteristic equation of (*) after calculation is given below

$$\Rightarrow \lambda^3 + 0.0809\lambda^2 + 0.0103\lambda + 0.0013 = 0$$

λ^3	1	0.0103	0
λ^2	0.0809	0.0013	0
λ^1	b_1	b_2	0
λ^0	k_1	k_2	0

And we can calculate out all the coefficients:

$$b_1 = \frac{-1}{0.0809} \begin{vmatrix} 1 & 0.0103 \\ 0.0809 & 0.0013 \end{vmatrix} = -0.0058$$

$$b_2 = \frac{-1}{0.0809} \begin{vmatrix} 1 & 0 \\ 0.0809 & 0 \end{vmatrix} = 0$$

$$k_1 = \frac{-1}{-0.0058} \begin{vmatrix} 0.0809 & 0.0013 \\ -0.0058 & 0 \end{vmatrix} = 0.0013$$

$$k_2 = \frac{-1}{-0.0058} \begin{vmatrix} 0.0809 & 0 \\ -0.0058 & 0 \end{vmatrix} = 0$$

And filling these values into our Routh Array, we can determine whether the system is stable or not:

λ^3	1	0.0103	0
λ^2	0.0809	0.0013	0
λ^1	-0.0058	0	0
λ^0	0.0013	0	0

Since the first columns do not contain the same sign, or the change of sign in the first column then by Routh-Hurwitz stability criteria the DFE point (0,1,0) is unstable. Using Matlab we obtain the values of the Eigen value as follows.

The roots of the equation is

$$\lambda_1 = 0.0111 + 0.1117i$$

$$\lambda_2 = 0.0111 - 0.1117i$$

$$\lambda_3 = -0.1032$$

4.4. Local Stability analysis of EE point of first age group

The endemic equilibrium solution of (3.5.12)-(3.5.14) can be found by setting the right hand side to zero and solving the nonlinear system to obtain

$$abms_{h1}i_v - (r_1 + \delta_1 + \lambda_h)i_{h1} + \delta_1 i_{h1}^2 = 0 \text{ (a)}$$

$$\lambda_h(1 - s_{h1}) - abms_{h1}i_v + v_1(1 - s_{h1} - i_{h1}) + \delta_1 s_{h1}i_{h1} = 0 \text{ (b)}$$

$$ca_1(1 - i_v)i_{h1} - \lambda_v i_v = 0 \text{ (c)}$$

Now from (c) we have that $i_v = \frac{a_1 c i_{h1}}{a_1 c i_{h1} + \lambda_v}$ (d) Substitute (d) in (a) and (b) solve for i_{h1}^* using Matlab we obtain

$$i_{h1}^* = 0.3433$$

Therefore the endemic equilibrium points for the respective states are:

$$(i_{h1}^*, s_{h1}^*, i_v^*) = (0.3433, 0.0458, 0.5164)$$

The Jacobian matrix of (3.5.2)-(3.5.14) at any endemic steady state $(i_{h1}^*, s_{h1}^*, i_v^*)$ is given by:

$$J(i_{h1}^*, s_{h1}^*, i_v^*) = \begin{bmatrix} -(r_1 + \delta_1 + \lambda_h) + 2\delta_1 i_{h1} & -a_1 b m i_v & a_1 b m s_{h1} \\ -v_1 + \delta_1 s_{h1} & -(\lambda_h + v_1) - a_1 b m i_v + \delta_1 i_{h1} & -a_1 b m s_{h1} \\ c a_1 (1 - i_v) & 0 & -c a_1 i_{h1} - \lambda_v \end{bmatrix}$$

The Jacobian matrix of this equilibrium point is evaluated at (0.9698, 0.0107, 0.7510) to obtain:

$$J(0.9698, 0.0107, 0.7510) = \begin{bmatrix} -0.0015 & -0.0238 & 0.0021 \\ -0.0083 & -0.0322 & -0.0021 \\ 0.1068 & 0 & -0.1468 \end{bmatrix}$$

The characteristic polynomial is given by

$$\begin{vmatrix} -0.0015 - \lambda & -0.0238 & 0.0021 \\ -0.0083 & -0.0322 - \lambda & -0.0021 \\ 0.1068 & 0 & -0.1468 - \lambda \end{vmatrix} = 0$$

Solving for the roots of the polynomial in the Jacobian matrix leads to the characteristic equation:

$$\lambda^3 + 0.1805\lambda^2 + 0.0050\lambda - 1.6439 \times 10^{-5} = 0$$

Now constructing Routh-Hurwitz array to determine stability in the following ways

λ^3	1	0.0050	0
λ^2	0.1805	-1.6439×10^{-5}	0
λ^1	b_1	b_2	0
λ^0	k_1	k_2	0

And we can calculate out all the coefficients:

$$b_1 = \frac{-1}{0.1805} \begin{vmatrix} 1 & 0.0050 \\ 0.1805 & -1.6439 \times 10^{-5} \end{vmatrix} = 0.0051$$

$$b_2 = \frac{-1}{0.1805} \begin{vmatrix} 1 & 0 \\ 0.1805 & 0 \end{vmatrix} = 0$$

$$k_1 = \frac{-1}{0.0051} \begin{vmatrix} 0.1805 & -1.6439 \times 10^{-5} \\ 0.0051 & 0 \end{vmatrix} = -1.6439 \times 10^{-5}$$

$$k_2 = \frac{-1}{0.0051} \begin{vmatrix} 0.1805 & 0 \\ 0.0051 & 0 \end{vmatrix} = 0$$

And filling these values into our Routh Array, we can determine whether the system is stable or not:

λ^3	1	0.0050	0
λ^2	0.1805	-1.6439×10^{-5}	0
λ^1	$b_1=0.0051$	$b_2=0$	0
λ^0	$k_1=$ -1.6439×10^{-5}	$k_2= 0$	0

Since all the first columns element contain the same sign, then by Routh-Hurwitz stability criteria the endemic equilibrium point is stable. Using Matlab we obtain the values of the Eigen value as follows.

The roots of the equation is

>>a=[10.1805 0.0050 - 1.6439 × 10⁻⁵];
 >> f=roots (a) λ₁ =-0.1453, λ₂ =-0.0382, λ₃ = 0.0030

All the roots dos not contain the same sign. Thus the EEP (0.3433, 0.0458, 0.5164) is unstable by Routh-Hurwitz stability criteria.

4.2 Numerical simulation of the second age group model

Consider the model given by equation (3.5.15)-(3.5.17)

$$\frac{di_{h2}}{dt} = a_2 b m s_{h2} i_{v2} - (r_2 + \delta_2 - \epsilon_1) i_{h2} - \epsilon_1 i h_1 + \delta_2 i_{h2}^2 \tag{i}$$

$$\frac{ds_{h2}}{dt} = -a_2 b m s_{h2} i_{v2} - \epsilon_1 s_{h1} + \delta_2 s_{h2} i_{h2} - \epsilon_2 s_{h2} + v_2 (1 - s_{h2} - i_{h2}) \tag{ii}$$

$$\frac{di_{v2}}{dt} = c a_2 (1 - i_{v2}) i_{h2} - \lambda_v i_{v2} \tag{iii}$$

4.5. The reproduction number of age group two(5-25 years old)

We have that $R_{01}^2 = \frac{a_2^2 b c m}{w_1 l \lambda v}$, or $R_{01} = \sqrt{\frac{a_2^2 b c m}{w_1 l \lambda v}}$

$$\Rightarrow R_{01} = \sqrt{\frac{(0.4)^2 (0.5) (0.48) (0.2)}{(0.071) (0.0035 + 0.0000000978 - 0.0000502)}} = 5.5995 > 1$$

Still the malaria transmission is present highly at this age level but less transmission than the first age group.

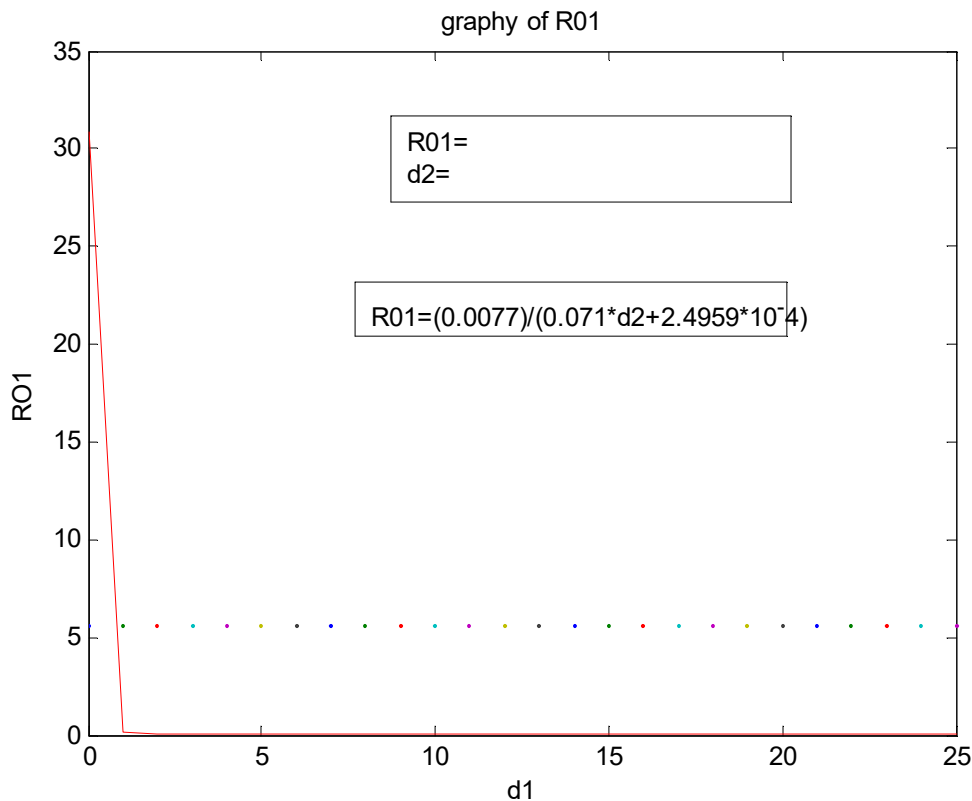


Figure 5.2 graph of the basic reproduction number R_{01}

From the above graph we can observe that if the Per capita death rate δ_2 increases then the reproduction number R_{01} decreases. This means that when the Per capita death rate δ_2 decrease then the reproduction number is

increasing and the susceptible population increasing. And when the Per capita death rate δ_2 increases R_{01} decreases that means when the when the Per capita death rate increase then the number of infected population decrease since individual in the infected class dies out.

4.6. Local Stability analysis of DFE point of second age group

Then the Jacobian matrix of the above matrix at DFE point $(0, 1, 0)$ is given by

$$J(0,1,0) = \begin{bmatrix} -(r_2 + \delta_2 - \varepsilon_1) & 0 & a_2bm \\ -v_2 + \delta_2 & -(v_2 + \varepsilon_1) & -a_2bm \\ ca_2 & 0 & -\lambda_v \end{bmatrix} \quad (\text{ii})$$

The characteristic polynomial is given by $|J - \lambda I_3| = 0$

$$\Rightarrow \begin{vmatrix} -(r_2 + \delta_2 - \varepsilon_1) - \lambda & 0 & a_2bm \\ -v_2 + \delta_2 & -(v_2 + \varepsilon_1) - \lambda & -a_2bm \\ ca_2 & 0 & -\lambda_v - \lambda \end{vmatrix} = 0$$

$$= \begin{vmatrix} -0.0034 - \lambda & 0 & 0.0400 \\ -0.0014 & -0.0015 - \lambda & -0.0400 \\ 0.1920 & 0 & -(0.071 + \lambda) \end{vmatrix} = 0$$

The corresponding characteristic equation is given by

$$\lambda^3 + a_1\lambda^2 + a_2\lambda + a_3 = 0$$

Where

$$a_1 = 0.0759, \quad a_2 = -0.0073, \quad a_3 = -5.4634 \times 10^{-4}$$

$$\Rightarrow \lambda^3 + 0.0759\lambda^2 - 0.0073\lambda - 5.4634 \times 10^{-4} = 0$$

Now we apply the array of Routh-Hurwitz stability criteria. If all the sign of the first column have the same sign, thus all the roots have the negative real parts. This shows that the DFE point $(0, 1, 0)$ is stable. Now to check stability we draw the following array:

λ^3	1	-0.0073	0
λ^2	0.0759	-5.4634×10^{-5}	0
λ^1	b_1	b_2	0
λ^0	k_1	k_3	0

And we can calculate out all the coefficients:

$$b_1 = \frac{-1}{0.0759} \begin{vmatrix} 1 & -0.0073 \\ 0.0759 & -5.4634 \times 10^{-5} \end{vmatrix} = -9.7101 \times 10^{-5}$$

$$b_2 = \frac{-1}{0.0759} \begin{vmatrix} 1 & 0 \\ 0.0759 & 0 \end{vmatrix} = 0$$

$$k_1 = \frac{-1}{-9.7101 \times 10^{-5}} \begin{vmatrix} 0.0759 & -5.4634 \times 10^{-4} \\ -9.7101 \times 10^{-5} & 0 \end{vmatrix} = -5.4634 \times 10^{-4}$$

$$k_3 = \frac{-1}{-9.7101 \times 10^{-5}} \begin{vmatrix} 0.07591 & 0 \\ -9.7101 \times 10^{-5} & 0 \end{vmatrix} = 0$$

And filling these values into our Routh Array, we can determine whether the system is stable or not:

λ^3	1	-0.0073	0
λ^2	0.00759	-5.4634×10^{-4}	0
λ^1	-9.7101×10^{-4}	0	0
λ^0	-5.4634×10^{-4}	0	0

Since all the coefficients in the first column of the R-H table all have no the same sign and there is sign change, then the system is unstable.

Since the first columns do not contain the same sign, then by Routh-Hurwitz stability criteria the DFE point is unstable. Using Matlab we obtain the values of the Eigen value as follows.

The roots of the equation is

$$\lambda_1 0.0852, \lambda_2 = -0.0889, \lambda_3 = -0.0722$$

4.7. Local Stability analysis of EE point of second age group

The endemic equilibrium solution of (3.5.15)-(3.5.17) can be found by setting the right hand side to zero and solving the nonlinear system to obtain:

$$a_2 b m s_{h2} i_v - (r_2 + \delta_2 - \epsilon_1) i_{h2} - \epsilon_1 i h_1 + \delta_2 i_{h2}^2 = 0 \tag{a}$$

$$-a_2 b m s_{h2} i_v - \epsilon_1 s_{h1} + \delta_2 s_{h2} i_{h2} - \epsilon_2 s_{h2} + v_2 (1 - s_{h2} - i_{h2}) = 0 \tag{b}$$

$$c a_2 (1 - i_{v2}) i_{h2} - \lambda_v i_{v2} = 0 \tag{c}$$

Now from (c) we have that $i_{v1} = \frac{a_2 c i_{h2}}{a_2 c i_{h2} + \lambda_v}$ (i)

Now substitute (i) in (a) and (b) solve for ih2*.

$$\Rightarrow ih2^* = 0.02068$$

Finally substitute the value of ih2* = 0.02068 in (a) we obtain

$$(ih2^{**}, sh2^{**}, iv2^{**}) = (0.2068, -0.0513, 0.3587). \text{ Then the Jacobian matrix of the model at any point}$$

$(ih2^{**}, sh2^{**}, iv2^{**})$ is given by

$$J(i_{h2}^*, s_{h2}^*, i_{v2}^*) =$$

$$\begin{bmatrix} -(r_2 + \delta_2 - \epsilon_1) + 2\delta_2 s_{h2} & a_2 b m i_{v2} & a_2 b m s_{h2} \\ -v_2 + \delta_2 s_{h2} & -a_2 b m i_{v2} + \delta_2 i_{h2} - \epsilon_1 - v_2 & -a_2 b m s_{h2} \\ c a_2 (1 - i_{v2}) & 0 & -c a_2 i_{h2} - \lambda_v \end{bmatrix}$$

Then the Jacobian matrix of the model at EE point $(ih2^{**}, sh2^{**}, iv2^{**}) = (0.2068, -0.0513, 0.3587)$ is given by

$$J(i_{h2}^*, s_{h2}^*, i_{v2}^*) =$$

$$\begin{bmatrix} -(r_2 + \delta_2 - \epsilon_1) + 2\delta_2(-0.0513) & a_2 b m(0.3587) & a_2 b m(-0.0513) \\ -v_2 + \delta_2(-0.0513) & -a_2 b m(0.3587) + \delta_2(0.2068) - \epsilon_1 - v_2 & -a_2 b m(-0.0513) \\ c a_2 (1 - 0.3587) & 0 & -c a_2(0.2068) - \lambda \end{bmatrix}$$

$$J(0.2068, -0.0513, 0.3587) = \begin{bmatrix} -0.0034 & 0.0143 & 0.0021 \\ -0.0014 & 0.0129 & -0.0021 \\ 0.1231 & 0 & -0.1107 \end{bmatrix}$$

The characteristic polynomial is given by $|J - \lambda I_3| = 0$

$$\Rightarrow \begin{vmatrix} -0.0034 - \lambda & 0.143 & 0.0021 \\ -0.0014 & 0.0129 - \lambda & -0.0021 \\ 0.1231 & 0 & -(0.1107 + \lambda) \end{vmatrix} = 0$$

The characteristic equation are given by $\lambda^3 + 0.1270\lambda^2 - 0.0013\lambda + 5.5762 \times 10^{-6}$

Now we apply the array of Routh-Hurwitz stability criteria. If all the sign of the first column have the same sign, thus all the roots have the negative real parts. This shows that the EE point (0.2068, -0.0513, 0.3587) is stable. Now to check stability we draw the following array:

λ^3	1	-0.0013	0
λ^2	0.1270	5.5762×10^{-6}	0
λ^1	b_1	b_2	0
λ^0	k_1	k_2	0

And we can calculate out all the coefficients:

$$b_1 = \frac{-1}{0.1270} \begin{vmatrix} 1 & -0.0013 \\ 0.1270 & -5.5762 \times 10^{-6} \end{vmatrix} = -0.0013$$

$$b_2 = \frac{-1}{0.1270} \begin{vmatrix} 1 & 0 \\ 0.1270 & 0 \end{vmatrix} = 0$$

$$k_1 = \frac{-1}{-0.0013} \begin{vmatrix} 0.0759 & -5.5762 \times 10^{-6} \\ -0.0013 & 0 \end{vmatrix} = -5.5762 \times 10^{-6}$$

$$k_2 = \frac{-1}{-0.0013} \begin{vmatrix} 0.1270 & 0 \\ -0.0013 & 0 \end{vmatrix} = 0$$

And filling these values into our Routh Array, we can determine whether the system is stable or not:

λ^3	1	-0.0013	0
λ^2	0.1270	-5.5762×10^{-6}	0
λ^1	-0.0013	0	0
λ^0	-5.5762×10^{-6}	0	0

Since all the coefficients in the first column of the R-H table all have no the same sign and there is sign change, then the system is unstable.

Since the first columns do not contain the same sign, then by Routh-Hurwitz stability criteria the DFE point is unstable. Using Matlab we obtain the values of the Eigen value as follows.

The roots of the equation is

$$\lambda_1 = -0.1368, \lambda_2 = 0.0049 + 0.0041i, \lambda_3 = 0.0049 - 0.0041i$$

4.8. Numerical simulation of the third age group model

To determine the reproduction number and stability of age group three we consider the following model

$$\frac{di_{h3}}{dt} = a_3 b m s_{h3} i_{v2} - (r_3 + \delta_3 + \epsilon_2) i_{h3} + \epsilon_2 i_{h2} + \delta_3 i_{h3}^2$$

$$\frac{ds_{h3}}{dt} = -a_3 b m s_{h3} i_{v3} - \epsilon_2 s_{h3} + \epsilon_2 s_{h2} + \epsilon_2 s_{h2} + v_3 (1 - s_{h3} - i_{h3}) + \delta_3 s_{h3} i_{h3}$$

$$\frac{di_{v1}}{dt} = ca_3(1-i_{v3})i_{h3} - \lambda_{v1}i_{v3}$$

4.9. The reproduction number of third age group

From the given model of age group model (3.5.18)-(3.5.20) we obtain the reproduction number:

$$R_{02}^2 = \frac{a_3^2 b c m}{m_1 l \lambda v}, \text{ or } R_{02} = \sqrt{\frac{a_3^2 b m c}{m_1 l \lambda v}}$$

$$\Rightarrow R_{02} = \sqrt{\frac{(0.2)^2(0.5)(0.48)(0.2)}{(0.071)(0.01 + 0.00000000667 + 0.0000502)}} = 1.6463 > 1$$

This indicate that the disease parasite in the community but less than the first age group.

graphy of R02

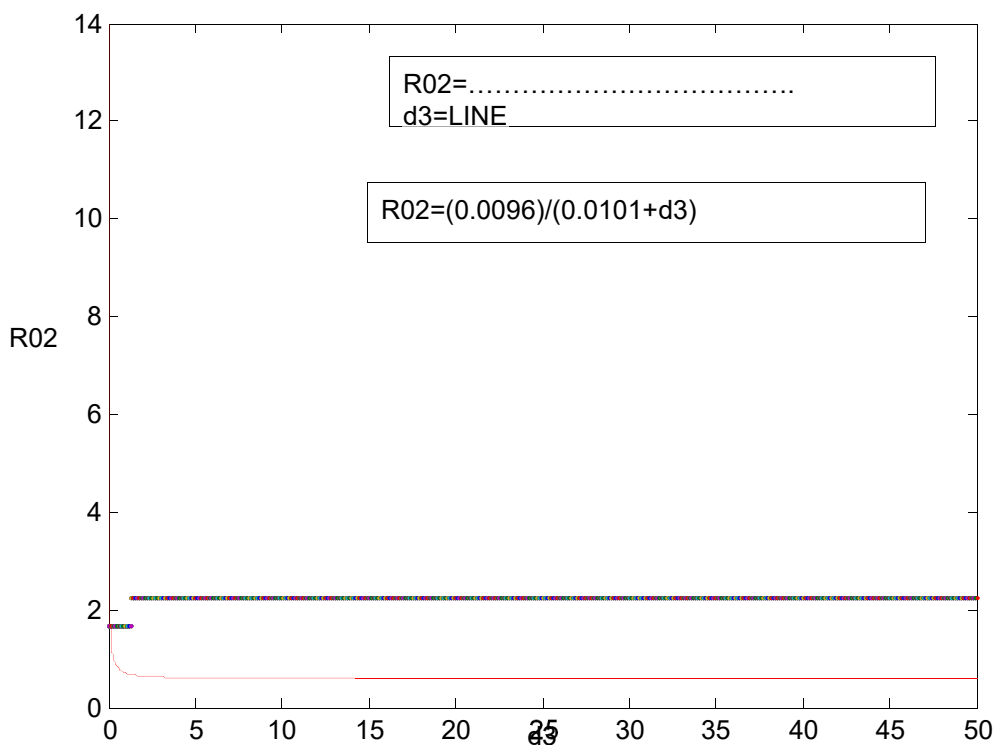


Figure 5.3 graph of the basic reproduction number R_{02}

From this graph we can observe that similar approach R_0 and phenomena observed. If the Per capita death rate δ_3 increases then the reproduction number R_0 decreases. This means that when the Per capita death rate δ_3 decrease then, the reproduction number is increasing and the susceptible population increasing. And when the Per capita death rate δ_3 increases R_{02} decreases that means when the when the Per capita death rate increase then the number of infected population decrease since individual in the infected class dies out.

4.10. Stability analysis of DFE point of third age group

The Jacobian matrix of the dynamical system (3.5.18)-(3.5.20) at any point $(ih2^*, sh2^*, iv2^*)$ is

$$J(i_{h_3}^*, s_{h_3}^*, i_v^*) = \begin{bmatrix} -(r_3 + \delta_3 + \varepsilon_2) + 2\delta_3 i_{h_3} & a_3 b m i_v & a_3 b m s_{h_3} \\ -v_3 + \delta_3 s_{h_3} & -a_3 b m i_v + \delta_3 i_{h_3} - \varepsilon_2 - v_3 & -a_3 b m s_{h_3} \\ c a_3 (1 - i_v) & 0 & -a_3 c i_{h_3} - \lambda_v \end{bmatrix}$$

Then the Jacobian matrix of the above matrix at DFE point (0,1,0) is given by

$$J(0,1,0) = \begin{bmatrix} -(r_3 + \delta_3 + \varepsilon_2) & 0 & a_3 b m \\ -v_3 + \delta_3 & -(v_3 + \varepsilon_2) & -a_3 b m \\ a_3 c & 0 & -\lambda_v \end{bmatrix} \text{ (ii)}$$

If we substitute the values of parameters in the given Jacobian matrix

$$J(0,1,0) = \begin{bmatrix} -0.0101 & 0 & 0.0200 \\ -7.9999 \times 10^{-4} & -8.5052 \times 10^{-4} & -0.0200 \\ 0.0960 & 0 & -0.071 \end{bmatrix}$$

The characteristic polynomial is given by

$$\begin{vmatrix} -0.0101 - \lambda & 0 & 0.0200 \\ -7.9999 \times 10^{-4} & -8.5020 \times 10^{-4} - \lambda & -0.0200 \\ 0.0960 & 0 & -(0.071 + \lambda) \end{vmatrix} = 0 \text{ (iii)}$$

The corresponding characteristic polynomial is

$$\lambda^3 + d_3 \lambda^2 + d_2 \lambda + d_1 = 0 \text{ where}$$

$$d_3 = 6.0968 \times 10^{-7} - 1.6154 \times 10^{-6} = -1.0257 \times 10^{-6}$$

$$d_2 = 7.1710 \times 10^{-4} + 8.5870 \times 10^{-6} + 6.0364 \times 10^{-5} = -0.0011$$

$$d_1 = 0.0101 + 0.071 + 8.5020 \times 10^{-4} = 0.0819$$

Now we apply the array of **Routh-Hurwitz stability criteria**. If all the sign of the first column have the same sign, thus all the roots have the negative real parts. This shows that the DFE point (0, 1, 0) is stable. Now to check whether the DFE point (0,1,0) is stability we draw the following array

λ^3	1	0.0819	0
λ^2	-0.0011	$a_4 = -1.0257 \times 10^{-6}$	0
λ^1	b_1	b_3	0
λ^0	k_1	k_3	0

Where

$$b_1 = \frac{-1}{-0.0011} \begin{vmatrix} 1 & 0.0819 \\ -0.0011 & -1.0257 \times 10^{-6} \end{vmatrix} = 0.0810$$

$$b_2 = \frac{-1}{-0.0011} \begin{vmatrix} 1 & 0 \\ -0.0011 & 0 \end{vmatrix} = 0$$

$$k_1 = \frac{-1}{0.0810} \begin{vmatrix} -0.0011 & -1.0257 \times 10^{-6} \\ 0.0810 & 0 \end{vmatrix} = -1.0257 \times 10^{-6}$$

$$k_2 = \frac{-1}{0.0810} \begin{vmatrix} -0.0661 & 0 \\ 0.0097 & 0 \end{vmatrix} = 0$$

And final we substitute

λ^3	1	0.0819	0
λ^2	-0.0011	$a_4 = -1.0257 \times 10^{-6}$	0
λ^1	0.0810	0	0
λ^0	-1.0257×10^{-6}	0	0

Where

From this array, we can clearly see that all of the signs of the first column do not contain positive value, there are sign changes, then by Routh-Hurwitz stability criteria the dynamical system of the given model is unstable.

Using Matlab we obtain

Roots :

$$\lambda_1 = -0.0371, \lambda_2 = 0.0298, \lambda_3 = -0.0009$$

4.11. Local Stability analysis of EE point of third age group

The endemic equilibrium solution of (3.5.18)-(3.5.20) can be found by setting the right hand side to zero and solving the nonlinear system to obtain:

$$\frac{di_{h3}}{dt} = a_3 b m s_{h3} i_{v2} - (r_3 + \delta_3 + \epsilon_2) i_{h3} + \epsilon_2 i_{h2} + \delta_3 i_{h3}^2 \quad (i)$$

$$\frac{ds_{h3}}{dt} = -a_3 b m s_{h3} i_{v3} - \epsilon_2 s_{h3} + \epsilon_2 s_{h2} + \epsilon_2 s_{h2} + v_3 (1 - s_{h3} - i_{h3}) + \delta_3 s_{h3} i_{h3} \quad (ii)$$

$$\frac{di_{v1}}{dt} = c a_3 (1 - i_{v3}) i_{h3} - \lambda_v i_{v3} \quad (iii)$$

Now from (c) we have that $i_v = \frac{a_3 c i_{h3}}{a_3 c i_{h3} + \lambda_v} \quad (iv)$

Now substitute (iv) in (a) and (b) solve for $i_{h3}^* \Rightarrow i_{h3}^* = -0.1274$

Finally substitute the value of $i_{h3}^* = -0.1274$ in (a) we obtain

$(i_{h3}^*, s_{h3}^*, i_v^*) = (-0.1274, 0.2281, 0.2853)$. Then the Jacobian matrix of the model at any point

$(i_{h3}^*, s_{h3}^*, i_v^*)$ is given by

$$J(i_{h3}^*, s_{h3}^*, i_v^*) =$$

$$\begin{bmatrix} -(r_3 + \delta_3 + \epsilon_2) + 2\delta_3 i_{h3}^* & a_3 b m i_v^* & a_3 b m s_{h3}^* \\ -v_3 + \delta_3 s_{h3}^* & -a_3 b m i_v^* + \delta_3 i_{h3}^* - \epsilon_2 - v_3 & -a_3 b m s_{h3}^* \\ c a_3 (1 - i_v^*) & 0 & -a_3 c i_{h3}^* - \lambda_v \end{bmatrix}$$

The Jacobian matrix of the third age group at any point $(i_{h3}^*, s_{h3}^*, i_v^*) = (-0.1274, 0.2281, 0.2853)$ is given as

$$J(-0.1274, 0.2281, 0.2853) = \begin{bmatrix} -0.0101 & -0.0057 & 0.0046 \\ 8.0000 \times 10^{-4} & -0.0049 & -0.0046 \\ 0.1234 & 0 & -0.0832 \end{bmatrix} \quad (*)$$

The characteristic polynomial (*) is given by

$$\begin{bmatrix} -0.0101 - \lambda & -0.0057 & 0.0046 \\ 8.0000 \times 10^{-4} & -0.0049 - \lambda & -0.0046 \\ 0.1234 & 0 & -0.0832 - \lambda \end{bmatrix}$$

The usual characteristic equation of $|J - \lambda I_3| = 0$ is given by

$$\begin{vmatrix} -0.0101 - \lambda & 0 & 0.0200 \\ -7.9999 \times 10^{-4} & -8.5020 \times 10^{-4} - \lambda & -0.0200 \\ 0.0960 & 0 & -(0.071 + \lambda) \end{vmatrix} = 0$$

$$= \lambda^3 + 0.0982\lambda^2 + 7.2985 \times 10^{-4}\lambda + 1.3362 \times 10^{-6} = 0$$

Now we apply the array of Routh-Hurwitz stability criteria. If all the sign of the first column have the same sign, thus all the roots have the negative real parts. This shows that the EE point (-0.1274,0.2281,-0.2853) is stable. Now to check whether the EEP(-0.1274,0.2281,-0.2853) is stability we draw the following array

λ^3	1	7.2985×10^{-4}	0
λ^2	0.0982	1.3362×10^{-6}	0
λ^1	b_1	b_2	0
λ^0	k_1	k_2	0

And we can calculate out all the coefficients:

$$b_1 = \frac{-1}{0.0982} \begin{vmatrix} 1 & 7.2985 \times 10^{-4} \\ 0.0982 & 1.3362 \times 10^{-6} \end{vmatrix} = 7.1624 \times 10^{-4}$$

$$b_2 = \frac{-1}{0.0982} \begin{vmatrix} 1 & 0 \\ 0.0982 & 0 \end{vmatrix} = 0$$

$$k_1 = \frac{-1}{7.1624 \times 10^{-4}} \begin{vmatrix} 0.0982 & 1.3362 \times 10^{-6} \\ 7.1624 \times 10^{-4} & 0 \end{vmatrix} = 1.3362 \times 10^{-6}$$

$$k_2 = \frac{-1}{7.1624 \times 10^{-4}} \begin{vmatrix} 0.0982 & 0 \\ 7.1624 \times 10^{-4} & 0 \end{vmatrix} = 0$$

And filling this values in the routh array we have

λ^3	1	7.2985×10^{-4}	0
λ^2	0.0982	1.3362×10^{-6}	0
λ^1	7.1624×10^{-4}	0	0
λ^0	1.3362×10^{-6}	0	0

Since all the coefficients in the first column of the R-H table all have the same sign and there is no sign change, then EEP (-0.1274,0.2281,-0.2853) is stable by Routh-Hurwitz stability criteria .Using Matlab we obtain the values of the Eigen value as follows.The roots of the equation is

$$\lambda_1 = -0.0903, \lambda_2 = -0.0049, \lambda_3 = -0.0030$$

5. DISCUSSION OF RESULT

In this paper a mathematical model of the prevalence and transmission of malaria was studied. The standard SIR differential equation model was used to predict the transmission and spread of malaria. By analyzing the model we found, the threshold parameter, R_0 found which the basic reproduction number. From our analysis, we observe that the disease free equilibrium is stable when the threshold parameter is less than unity. i.e if $R_0 < 1$, then the epidemic will not spread and when $R_0 > 1$ the disease was parasite in the population and hence become endemic.

The model has two equilibrium points namely the disease free equilibrium point and endemic equilibrium points. The existence and stability of disease free equilibrium point, endemic equilibrium points and sensitivity analysis of the model where preformed. We developed a model for the spread of malaria disease by human host vector population. The model divided three age classes children (age 0-5 years), youngest population (age 6-25), the

remaining age classes and three epidemiological classes susceptible, infected and recovered. The model incorporates death rate due to malaria infection, death rate due to natural, new born of population, infection contact rate depending on the local population densities. Births are assumed to be continuous throughout the year. The model is not considered the migration and emigration of the population in to compartment and from compartment.

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