Stability Analysis and Dynamics Preserving Non-Standard Finite Difference Schemes for a Malaria Model

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10⁻⁷ 18 0.02

0.08333

0.003704

 3.454×10

 2.279×10

 $0.6 \\ 0.8333$

0.08333

0.1

0.1

0.0035

9×10

0.13

0.033 2×10 0.5 0.48

0.048

0.091

(11)

(13)

(14)

(15)

(16)

(18)

(19)

0.1

1.8×10

 $0.33 \\ 0.24$

0.024

0.083

Table 2: Description of parameters and three sets of values used in numerical simulation

 $\mathcal{G} = \left\{ \left(\begin{array}{ccc} S_h & E_h & I_h & R_h & S_v & E_v & I_v \end{array} \right) \in \mathcal{D}; \, N_h^\# \leq N_h \leq N_h^*, \, N_v = N_v^* \right\}$

Theorem 1 The set \mathcal{G} in (10) is GAS for the dynamical system (1)–(7) defined on \mathcal{D} . (Thus, the study of the system (1)–(7) can be reduced from \mathcal{D} to \mathcal{G} .)

 $R_0 \le \xi$

 $R_0 = c(N_h^*, N_v^*) \sqrt{\frac{\beta_{hv} \nu_h \nu_v \left(\beta_{vh} + \frac{\gamma_h + \tilde{\beta}_{vh}}{\rho_h + f_h(N_h^*)}\right) N_h^* N_v^*}{f_v(N_v^*) (\nu_h + f_h(N_h^*)) (\nu_v + f_v(N_v^*)) (\gamma_h + f_h(N_h^*) + \delta_h)}}.$

 $\sqrt{\frac{\sigma_{h}N_{h}^{\#}+\sigma_{v}N_{v}^{*}}{\sigma_{h}N_{h}^{*}+\sigma_{v}N_{v}^{*}}}\times\frac{\nu_{h}+\mu_{1h}+\mu_{2h}N_{h}^{\#}}{\nu_{h}+\mu_{1h}+\mu_{2h}N_{h}^{\#}}\times\frac{\gamma_{h}+\delta_{h}+\mu_{1h}+\mu_{2h}N_{h}^{\#}}{\gamma_{h}+\delta_{h}+\mu_{1h}+\mu_{2h}N_{h}^{\#}}\times\frac{\beta_{vh}+\tilde{\beta}_{vh}\frac{\gamma_{h}}{\rho_{h}+\mu_{1h}+\mu_{2h}N_{h}^{*}}}{\beta_{vh}+\tilde{\beta}_{vh}\frac{\gamma_{h}}{\rho_{h}+\mu_{1h}+\mu_{2h}N_{h}^{*}}}\times\frac{\beta_{vh}+\tilde{\beta}_{vh}\frac{\gamma_{h}}{\rho_{h}+\mu_{1h}+\mu_{2h}N_{h}^{*}}}{\beta_{vh}+\tilde{\beta}_{vh}\frac{\gamma_{h}}{\rho_{h}+\mu_{1h}+\mu_{2h}N_{h}^{*}}}$

Remark 3 Since $\xi \le 1$, Theorem 2 is consistent with the bifurcation analysis in [2]: at $R_0 = 1$ there is forward bifurcation if $\delta_h = 0$ ($\xi = 1$) and possible backward bifurcation if $\delta_h > 0$ ($\xi < 1$).

 $\begin{array}{ll} {n+1 - S_h^n \over \phi(\Delta t)} & = & \Lambda_h + \psi_h N_h^n + \rho_h R_h^{n+1} - c(N_h^n, N_v^n) \beta_{hv} I_v^n S_h^{n+1} - f_h(N_h^n) S_h^{n+1}, \end{array}$

 $\begin{array}{ll} \frac{1}{s+1} - S_v^n \\ \frac{1}{\phi(\Delta t)} & = & \psi_v N_v^n - c(N_h^n, N_v^n) (\beta_{vh} I_h^n + \tilde{\beta}_{vh} R_h^n) S_v^{n+1} - f_v(N_v^n) S_v^{n+1}, \end{array}$

 $\frac{E_v^{n+1} - E_v^n}{\phi(\Delta t)} \ = \ c(N_h^n, N_v^n)(\beta_{vh}I_h^n + \tilde{\beta}_{vh}R_h^n)S_v^{n+1} - \nu_v E_v^{n+1} - f_v(N_v^n)E_v^{n+1},$

 $\phi \equiv \phi(\Delta t) = \Delta t + O[(\Delta t)^2].$ $N_v^{n+1} = F_v(N_v^n)$ $\underline{F}_h(N_h^{n-1}) =: \underline{N}_h^n \leq N_h^n \leq \overline{N}_h^n := \overline{F}_h(N_h^{n-1})$

where F_v , \underline{F}_h and \overline{F}_h are suitable maps with the same fixed-points N_{ν}^* , $N_h^{\#}$ and N_h^* , respectively, as for the continuous model, and $\phi(\Delta t) = (\Lambda_h \mu_{2h})^{-\frac{1}{2}} [1 - e^{-\Delta t (\Lambda_h \mu_{2h})^{\frac{1}{2}}}].$ **Theorem 4** The set G in (10) is GAS for the discrete system (13)–(19) defined on D under the Theorem 5 The DFE is a GAS for the discrete dynamical system (13)-(19), with (22), on D.

 $\frac{E_h^{n+1} - E_h^n}{\phi(\Delta t)} \ = \ c(N_h^n, N_v^n) \beta_{hv} I_v^n S_h^{n+1} - \nu_h E_h^{n+1} - f_h(N_h^n) E_h^{n+1},$

 $\frac{n+1}{\phi(\Delta t)} - \frac{I_h^n}{I_h^n} = \nu_h E_h^{n+1} - (\gamma_h + f_h(N_h^n) + \delta_h) I_h^{n+1},$

 $\frac{R_h^{n+1} - R_h^n}{\phi(\Delta t)} = \gamma_h I_h^{n+1} - \rho_h R_h^{n+1} - f_h(N_h^n) R_h^{n+1},$

 $\frac{n+1}{\phi(\Delta t)} - I_v^n = \nu_v(t)E_v^{n+1} - f_v(N_v^n)I_v^{n+1}$

is the basic reproduction number and the additional threshold number ξ is given by

3 A nonstandard finite difference scheme Consider the following NSFD scheme in the sense of [1, 4]:

Description

recovery rate
loss of immunity rate
disease-induced death rate

relative birth rate density-independent death rate density-dependent death rate bites required by a mosquito per unit time probability of transmission of infection from infective human probability of transmission of infection from recovered human transfer rate to infective transfer rate to infective in the probability of the probability of transmission of infection from recovered human transfer rate to infective in the probability of the property of the probability of the probability

relative birth rate

 $\tilde{\boldsymbol{\beta}}_{vh}$

2 GAS results

Compact biologically feasible region:

Following [3] for the model on G, we have: Theorem 2 The DFE is GAS on D when



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We extend the results in [2] by proving the GAS of the DFE and specifying the region of possible backward bifurcation. Furthermore, we design a nonstandard finite difference (NSFD) scheme, which is dynamically consistent with the continuous model

Fig. 1, Table 1 and Table 2 correspond to the model

$\frac{dS_h}{dt}$	=	$\Lambda_h + \psi_h N_h + \rho_h R_h - c(N_h, N_v) \beta_{hv} I_v S_h - f_h(N_h) S_h,$	(1)
$\frac{dE}{dt}$	=	$c(N_h,N_v)\beta_{hv}I_vS_h - \nu_hE_h - f_h(N_h)E_h,$	(2)
$\frac{dI_h}{dt}$	=	$\nu_h E_h - [\gamma_h + f_h(N_h) + \delta] I_h,$	(3)
$\frac{dR_h}{dt}$	=	$\gamma_h I_h - \rho_h R_h - f_h(N_h) R_h,$	(4)
$\frac{dS_v}{dt}$	=	$\psi_v N_v - c(N_h, N_v) (\beta_{vh} I_h + \tilde{\beta}_{vh} R_h) S_v - f_v(N_v) S_v,$	(5)
$\frac{dE_v}{dt}$	=	$c(N_h,N_v)[\beta_{vh}I_h+\tilde{\beta}_{vh}R_h]S_v-\nu_vE_v-f_v(N_v)E_v,$	(6)
$\frac{dI_v}{dt}$	=	$\nu_v(t)E_v - f_v(N_v)I_v$	(7)

$$\begin{array}{lll} f_h & = & \mu_{1h} + \mu_{2h} N_h, & f_v = \mu_{1v} + \mu_{2c} N_v, \\ N_h & = & S_h + E_h + I_h + R_h, & N_v = S_v + E_v + I_v, \\ c(N_h, N_v) & = & \frac{\sigma_v \sigma_h}{\sigma_h N_h + \sigma_v N_v}. \end{array}$$

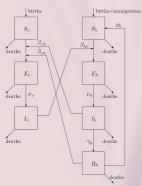


Figure 1: Compartmental Flow Diagram

Unbounded biologically feasible region:

 $D = \{(S_h \ E_h \ I_h \ R_h \ S_v \ E_v \ I_v) \in \mathbb{R}^7_+\}$

Conservation laws for vector and host:

$$\begin{split} \frac{dN_v}{dt} &= (\psi_v - \mu_{1v} - \mu_{2v} N_v) N_v \text{ with GAS equilibrium } N_v^\star = \frac{\psi_v - \mu_{1v}}{\mu_{2v}}. \\ &\qquad \underline{N}_h(t) \leq N_h(t) \leq \overline{N}_h(t) \end{split}$$

with, $\overline{N}_h(t)$ and $\underline{N}_h(t)$ being suitable "upper" and "lower" solutions, equilibrium

 $\begin{array}{ll} N_h^* &=& (\psi_h - \mu_{1h} + \sqrt{(\psi_h - \mu_{1h})^2 + 4\mu_{2h}\Lambda_h})/2\mu_{2h} \; \text{when} \; \; \delta_h = 0 \\ N_h^\# &=& (\psi_h - \mu_{1h} - \delta_h + \sqrt{(\psi_h - \mu_{1h} - \delta_h)^2 + 4\mu_{2h}\Lambda_h})/2\mu_{2h} \; \text{when} \; \delta_h \neq 0 \end{array}$

Disease free equilibrium (DFE):

 $DFE = (N_h^*, 0, 0, 0, N_v^*, 0, 0)$

Humans			Mosquito		
S_h :	Number of susceptible humans	S_v :	Number of susceptible mosquito		
E_h :	Number of exposed humans	E_v :	Number of exposed mosquito		
I_h :	Number of infective humans	I_v :	Number of infective mosquito		
R_h :	Number of recovered (immune an	d asymp	tomatic, but slightly infectious) humans		

Table 1: The state variables of the model (1) (7)

	Set 1	Set 2	Set 3
R_0	0.9503	0.9898	4.4402
ξ	0.9583	0.4124	not relevant
Threshold condition	$R_0 \le \xi$	$\xi < R_0 < 1$	$R_0 > 1$
Stability of DFE	GAS	asymptotically stable (possibly co-exists with EE)	unstable

Table 3: Threshold numbers for the three sets of parameter values and the stability of DFE



Figures 2, 3 and 4 represent human population by compartment (left); total population, its lower bound X_h and its upper bound \overline{N}_h in terms of (9) (right). Figures 5, 6 and 7 provide phase diagrams of Infected or Disease Carriers $(E_h + I_h + R_h)$ versus susceptible (S_h) . The five pointed stars indicate the initial points of the trajectories. The shaded area is the projection of the set \mathcal{G} . An invariant manifold of one dimension less is clearly indicated on each figure. It is of interest to notice the coexistence of EE and DFE on figure 6. There are two asymptotically stable equilibria, denoted by circled stars and an unstable equilibrium denoted by a circle all within the region \mathcal{G} . This unstable equilibrium, which one can also see is a saddle point, actually accounts for the dip in the population size observed on Fig. 3.

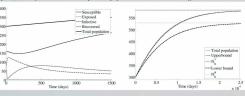


Figure 2: Parameter values from Table 2, Set 1: The rate of change of the compartments is even tually comparable with the rate of change of the total population (left). The solution approache DFE (right). The total population remains between N_{λ} and \overline{N}_{h}

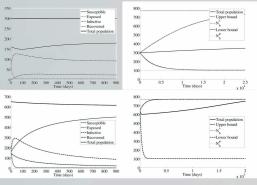


Figure 3: Parameter values from Table 2, Set 2: Two typical solutions are represented one conve ing to DFE (bottom) and one converging to an EE (top). In both cases the conservation law (9 preserved (right). It is of interest to observe also the initial dip in the total population when is solution approaches DFE (bottom, right) which cannot be assimilated through a logistic equation

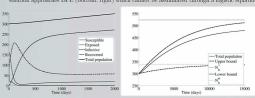
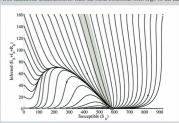


Figure 4: Parameter values from Table 2, Set 3: A typical solution initialed at a point outside the disease free manifold demonstrates that all such solutions converge to an EE.



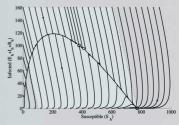


Figure 6: Backward bifurcation: Parameter values from Table 2, Set 2.

References

- [2] N. Chitnis, J.M. Cushin and J.M. Hyman, Bifurcation analysis of a mathematical model for Malaria transmission, SIAM J. Applied Math., 67 (2006), 24–45
- [3] J. C. Kamgang and G. Sallet, Computation of threshold conditions for epidemiological models and global stability of the disease free equilibrium, Math. Biosc. 213 (2008), 1–12.

4 Numerical simulations

immigration rate relative birth rate density-independent death/emmigration rate density-dependent death/emmigration rate bites tolerated by a human per unit time probability of transmission of infection from infective mosquito transfer rate to infective $\frac{1}{ah_0}$ = average duration of the latent period recovery rate 0.041 5.5×10^{-5} 8.8×10^{-6} 2×10^{-7} 4.3 0.0220.033 1.1×10^{-4} 1.6×10^{-5} 3×10^{-7} 19 0.022

