# The dynamic of a SIV epidemic disease model with vertical infection and pulse Vaccination

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#### Abstract

In this paper, we conside a SIV epidemic disease model with two delays and vertical infection, and the dynamic behaviors of the model under pulse vaccination are analyzed. Using a new modeling method, we obtain sufficient condition for the permanence of the epidemic model with pulse vaccination.

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## **1** Introduction

Many infectious diseases in nature transmit through both horizontal and vertical modes [2, 3]. These diseases include such human diseases as rubella, herpes simplex Hepatitis B, AIDS, and so on. For human and animal diseases, horizontal transmission typically occur through direct or indirect physical contact with infectious host, or through disease vector such as mosquitoes, tick, or other biting insects. Vertical transmission can be accomplished through transplacental transfer of disease agents. Busenberg and Cooke [1] discussed a variety of diseases that transmit both vertically and horizontally and gave a comprehensive survey of the formulation and the mathematical analysis of compartment models that incorporate vertical transmission. In our paper, we assume a fraction of the offspring of infected hosts is infected at birth, and hence the infected birth flux will enter class I. Besides a susceptible individual goes through an infectious period. Since time delay has important biologic meaning in epidemic models.

Therefore, in our paper, we consider two time delays, that is, the number of the susceptible individuals during an infectious period and temporary immunity period should be considered, denoted by  $\tau$ ,  $\omega$ , respectively. Now we create a new delay SIV epidemic model with vertical infection, pulse vaccination into epidemic model and to obtain some important qualitative properties with delays and valid pulse vaccination strategy.

## 2 Main Results

In the following, we consider the SIV epidemic model with vaccination.

$$\begin{cases} \frac{dS}{dt} = (1-\alpha)(bN - pbI) + \lambda I + \theta V - \frac{\beta S(t-\tau)I(t-\tau)}{N} - bS, \\ \frac{dI}{dt} = pbI + \frac{\beta S(t-\tau)I(t-\tau)}{N} + \frac{\zeta\beta V(t-\omega)I(t)}{N} - (b+\lambda)I, \\ \frac{dV}{dt} = \alpha(bN - pbI) - \frac{\zeta\beta V(t-\omega)I(t)}{N} - (b+\theta)V, \end{cases} \quad t \neq nT,$$

$$\begin{cases} 1 \\ S(t^{+}) = (1-\delta)S(t), \\ I(t^{+}) = I(t), \\ V(t^{+}) = V(t) + \delta S(t), \end{cases} t = nT. \end{cases}$$

Since the natural birth rate and death rate are the same and the disease is assumed not to inflict death on the infected host, so the total population is constant, without loss of generality, let N(t) = 1, thus, the system (1) can be reduced as following:

$$\begin{cases} \frac{dS}{dt} = (1-\alpha)(b-pbI) + \lambda I + \theta V - \beta S(t-\tau)I(t-\tau) - bS, \\ \frac{dI}{dt} = pbI + \beta S(t-\tau)I(t-\tau) + \zeta \beta V(t-\omega)I(t) - (b+\lambda)I, \\ \frac{dV}{dt} = \alpha(b-pbI) - \zeta \beta V(t-\omega)I(t) - (b+\theta)V, \end{cases} t \neq nT,$$

$$\begin{cases} 20 \\ S(t^{+}) = (1-\delta)S(t), \\ I(t^{+}) = I(t), \\ V(t^{+}) = V(t) + \delta S(t), \end{cases} t = nT. \end{cases}$$

Since V(t) = 1 - S(t) - I(t), therefore we may just discuss S(t), I(t). Let

$$C_h^+ = \{ \phi = (\phi_1(s) \phi_2(s)) \in C_h : \phi_i(0) > 0 \},\$$

where  $\phi_i$  is positive, bounded and continuous function for  $s \ s \in [-\theta, 0]$ , where  $\theta = \max{\{\tau, \omega\}}$ .

Denote

$$R_{2} = \frac{\hbar((1-\alpha)b+\theta)(1-\delta)(1-e^{-(b+\theta)T})}{(b+\theta)[1-(1-\delta)e^{-(b+\theta)T}]}, \qquad m_{2}^{*} = \frac{((1-\alpha)b+\theta)(R_{2}-1)}{((1-\alpha)pb+\theta+\beta)R_{2}},$$

where

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$$\hbar = \frac{\beta}{(1-p)b + \lambda}$$

**Theorem 2.1** Suppose  $\lambda + (1-p)b - \zeta\beta > 0$ . If  $R_2 > 1$ , then there exists a positive constant  $m_2$  such that  $I(t) \ge m_2$  for t large enough.

**Proof:** Suppose that x(t) = (S(t), I(t)) is any positive solution of system (2). The second equation of system (2) may be rewritten as follow:

$$\frac{dI}{dt} = \beta S(t-\tau)I(t-\tau) - (b+\lambda - pb - \zeta\beta V(t-\omega))I(t)$$

Define

$$R(t) = I(t) + \beta \int_{t-\tau}^{t} S(\theta) I(\theta) d\theta$$
(3)

calculating the derivative of R(t) along the solution (2), it follows from (3)

$$\frac{dR(t)}{dt} = \frac{dI(t)}{dt} + \beta S(t)I(t) - \beta S(t-\tau)I(t-\tau)$$

$$\geq (\lambda + (1-p)b - \zeta\beta)(\hbar S(t) - 1)I(t).$$
(4)

since  $R_2 > 1$ , then  $m_2^* > 0$  and there exists a positive constant  $\varepsilon_1$  small enough such that  $\hbar \sigma > 1$ . Where

$$\varpi = \frac{(1-\alpha)(b-pbm_{2}^{*}) + (1-m_{2}^{*})\theta - m_{2}^{*}\beta}{b+\theta} \cdot [1-\frac{\delta}{1-(1-\delta)\exp(-(b+\theta)T)}] - \varepsilon_{1} > 0,$$

for any positive constant  $t_0$ , we claim that the inequality  $I(t) < m_2^*$  can not hold for  $t \ge t_0$  otherwise, there is a positive constant  $t_0$ , such that  $I(t) < m_2^*$  for all  $t \ge t_0$ . From the first and the fourth equations of system (2), we have

$$\begin{cases} \frac{dS}{dt} \ge (1-\alpha)(b-pbm_{2}^{*}) + (1-m_{2}^{*})\theta - \beta m_{2}^{*} - (b+\theta)S, & t \neq nT, \ n \in N. \\ S(t^{+}) = (1-\delta)S(t), & t = nT, \ n \in N. \end{cases}$$
(5)

we know that there exists such  $T_1 \ge t_0 + \tau$  for  $t \ge T_1$ , that

$$\frac{S(t) >}{\frac{(1-\alpha)(b-pbm_2^*) + (1-m_2^*)\theta - \beta m_2^*}{b+\theta}} \cdot \left[1 - \frac{\delta}{1 - (1-\delta)\exp(-(b+\theta)T)}\right] - \varepsilon_1 = \varpi$$
<sup>(6)</sup>

We have that  $S(t) > \varpi$  for  $t \ge T_1$ .

By (4) and (6), we see that

$$\frac{dR(t)}{dt} > (\lambda + (1-p)b - \zeta\beta)(\hbar \varpi - 1)I(t), \quad t \ge T_1.$$
(7)

Let

$$I^{l} = \min_{t \in [T_{1}, T_{1} + \tau]} I(t)$$

We show that  $I(t) \ge I^{l}$  for all  $t \ge T_{1}$ , otherwise, there exists a nonnegative constant  $T_{2}$  such that  $I(t) \ge I^{l}$  for  $t \in [T_{1}, T_{1} + \tau + T_{2}], I(T_{1} + \tau + T_{2}) = I^{l}$  and  $\frac{dI(T_{1} + \tau + T_{2})}{dt} \le 0$ . Thus from the second equation of (2), (4) and (7), we easily see

that

$$\frac{dI(T_1+\tau+T_2)}{dt} \ge (\lambda+b-pb-\zeta\beta)(\hbar\varpi-1)I^l > 0.$$

This contradiction. Hence, we get that  $I(t) \ge I^{l} > 0$ , for all  $t \ge T_{1}$ . From (7), we have

$$\frac{dR(t)}{dt} > (\lambda + b - pb - \zeta\beta)(\hbar \, \varpi - 1)I^{l} > 0,$$

This implies  $R(t) \to +\infty$  as  $t \to +\infty$ . This is a contradiction to  $R(t) \le 1 + \beta \tau$  for t large enough. Therefore, for any positive constant  $t_0$ , the inequality  $I(t) < m_2^*$ can not hold for all  $t \ge t_0$ .

On the other hand, if  $I(t) \ge m_2^*$  holds true for all t large enough, then our aim is obtained. On the other hand, I(t) is oscillatory about  $m_2^*$ . where, The dynamic of a SIV epidemic disease model ...

$$m_2 = \min\{\frac{m_2^*}{2}, m_2^*e^{-(\lambda+b-pb)\tau}\}.$$

Critical values of some parameters of systems (2) ( $R_2 > 1$  must be satisfied). The condition for the permanence of epidemic sease

$$\delta < \delta^*, \quad \delta^* = 1 - \frac{(b+\theta)e^{(b+\theta)T}}{((1-\alpha)b+\theta)\hbar(e^{(b+\theta)T}-1) + (b+\theta)}$$
$$T > T^*, \quad T^* = \frac{1}{b+\theta} In \left(1 + \frac{\delta(b+\theta)}{\hbar((1-\alpha)b+\theta)(1-\delta) - (b+\theta)}\right)$$

In the following, we shall show that  $I(t) \ge m_2$ . There exists two positive constant  $\bar{t}, \psi$  such that  $I(\bar{t}) = I(\bar{t}+\psi) = m_2^*$ , and  $I(t) < m_2^*$  for  $\bar{t} < t < \bar{t}+\psi$ . When  $\bar{t}$  is large enough, the inequality  $S(t) > \varpi$  holds true for  $\bar{t} < t < \bar{t}+\psi$ , since I(t) is continuous and ultimately bounded and is not effected by impulses. We conclude that I(t) is uniformly continuous.

Hence there exists a constant  $T_3$  (with  $0 < T_3 < \theta$  and  $T_3$  is independent of the choice of  $\bar{t}$ ) such that  $I(t) > \frac{m_2^*}{2}I(t) > \frac{m_2^*}{2}$  for all  $\bar{t} < t < \bar{t} + T_3$ . If  $\psi \le T_3$ , our aim is obtained.

If  $T_3 < \psi \le \theta$ , from the second equation of system (2) we have that

$$\frac{dI(t)}{dt} \ge -(\lambda + b - pb)I(t) \text{ for } \vec{t} < t < \vec{t} + \psi.$$

Then we have

$$I(t) \ge m_2^* e^{-(\lambda + b - pb)\theta}$$
, for  $t < t < t + \psi \le t + \theta$ .

Since  $I(t) = m_2^*$ , it is obvious that  $I(t) > m_2^*$  for  $t < t < t + \psi$ . If  $\psi \ge \theta$ , then we have that  $I(t) \ge m_2$  for  $t < t \le t + \theta$ . The same argument can be continued, we can obtain  $I(t) \ge m_2$ , for  $t + \theta \le t \le t + \psi$ . Since the interval  $[t, t + \psi]$  is

arbitrarily chosen, we get that  $I(t) \ge m_2$  for t large enough. In view of our arguments above, the choice of  $m_2$  is independent of the positive solution of system (2) which satisfies that  $I(t) \ge m_2$  for sufficiently large t. This completes the proof.

**Theorem 2.2** If  $R_2 > 1$ , then system (2) is uniformly permanent.

**Proof:** Suppose that X(t) = (S(t), I(t)) is any positive solution of system (2) with initial conditions (3). From the first and the fourth equations of (2), we have that

$$\begin{cases} \frac{dS}{dt} \ge b(1-\alpha)(1-p) - \beta - (b+\theta)S, & t \ne nT, \ n \in N. \\ S(t^+) = (1-\delta)S(t), & t = nT, \ n \in N. \end{cases}$$
(8)

we can get such t large enough and  $\varepsilon > 0$  small enough that

$$S(t) \ge \frac{\left((1-\alpha)(b-pb)-\beta\right)}{b+\theta} \frac{(1-\delta)(1-e^{-(b+\theta)T})}{1-(1-\delta)e^{-(b+\theta)T}} - \varepsilon = m_1, \quad \text{for} \quad t > T_4$$

Set  $D = \{(S, I) \in \mathbb{R}^2 : m_1 \leq S(t) \leq 1, m_2 \leq I(t) \leq 1\}$ . Then *D* is a bounded compact region in which has positive distance from coordinate hyperplanes. One obtains that every solution of system (2) eventually enters and remains in the region *D*. The proof is completed.

# References

- S. Busenberg and K. Cooke, Vertically Transmitted Disease, Model and Dynamics, Biomathematical, 23, Springer-Verlag, Berlin, 1993.
- [2] Z. Lu, X. Chi and L. Chen, The effect of constant and pulse vaccination on sir epidemic model with horizontal and vertical transmission, *Math. Comput. Model*, 36, (2002), 1039-1105.

[3] W. Wang, Global behavior of an SEIR epidemic model with two delays, *Appl.Math.Biol.*, **35**, (1996), 240-260.