

## A MATHEMATICAL MODEL OF HIV-1 INFECTION INCLUDING THE SATURATION EFFECT OF HEALTHY CELL PROLIFERATION

MAHIÉDDINE KOUICHE \*, BEDR'EDDINE AINSEBA \*\*

\* Laboratory of Applied Mathematics  
Badji-Mokhtar University, BP 12, Annaba 23000, Algeria  
e-mail: kouchebone@yahoo.fr

\*\* Institute of Mathematics of Bordeaux, UMR CNRS 52 51, Case 36  
Université Victor Segalen Bordeaux 2, 3 ter place de la Victoire, F 33076 Bordeaux Cedex, France  
e-mail: bea@u-bordeaux2.fr

In this paper we derive a model describing the dynamics of HIV-1 infection in tissue culture where the infection spreads directly from infected cells to healthy cells through cell-to-cell contact. We assume that the infection rate between healthy and infected cells is a saturating function of cell concentration. Our analysis shows that if the basic reproduction number does not exceed unity then infected cells are cleared and the disease dies out. Otherwise, the infection is persistent with the existence of an infected equilibrium. Numerical simulations indicate that, depending on the fraction of cells surviving the incubation period, the solutions approach either an infected steady state or a periodic orbit.

**Keywords:** HIV, periodic oscillations, persistence, stability, tissue culture.

### 1. Introduction

Since the early 1980s there has been a tremendous effort made in mathematical modeling of the Human Immunodeficiency Virus type 1 (HIV-1), which causes AIDS (Acquired Immune Deficiency Syndrome). The major target of HIV-1 infection is a class of lymphocytes or white blood cells known as  $CD4^+$  T-cells, which are the most abundant white blood cells of the immune system. It is thought that HIV-1, although attacking many different cells, wreaks the most havoc on the  $CD4^+$  T-cells by causing their destruction and decline, and decreasing the body's ability to fight the infection. The process of infection is as follows: First, HIV-1 enters its target T-cell, inside this cell it makes, with the help of the reverse transcriptase enzyme RT, a DNA copy of its viral RNA. The viral DNA is inserted into the DNA of the host cell (T-cell), which will itself produce viral particles that can bud off the cell to infect other healthy cells.

Many mathematical models have been derived in order to describe the dynamics of HIV-1 infection in the bloodstream where cell-free-viral spread is the predominant route of viral spread (De Leenheer and Smith, 2003; Kirschner, 1996; Nowak and Bangham, 1996; Perelson, 1989; Perelson *et al.*, 1993; Perelson and Nelson, 1999;

Nowak and May, 2000). Upon infection with HIV-1, there is a short "intracellular eclipse phase", during which the cell is infected but has not yet begun producing the virus. This intracellular eclipse phase is modeled by introducing an explicit class of latently infected cells. The models are then defined by considering four compartments: cells that are uninfected or healthy cells, cells that are latently infected, cells that are actively infected, and the free virus. The dynamics of these populations are described by a system of four ordinary differential equations. To take into account the "immune responsiveness" of the system, the class of  $CD8^+$  T-cells is introduced in the proliferation of  $CD4^+$  T-cells (Kirschner and Webb, 1996). These cells cannot become infected with the virus, but do destroy infected cells. It was shown that these models can mimic the symptom of AIDS observed clinically such as the long latency period, low levels of the free virus in the body, and the depletion of  $CD4^+$  T-cells.

Herz *et al.* (1996) proposed to model the latency period by introducing a delay into the class of infected cells by assuming that cells become infected  $\tau$  times units after initial infection and obtained a new model of three delay differential equations. Tam (1999) investigated the delay effect in a model which describes the interaction between

a replicating virus and host cells. Culshaw and Ruan (2000) showed that such an intracellular delay can cause the cell and virus populations to fluctuate in the early stage of infection (called infective oscillations in the literature), and in a long term they will converge to the infected steady state values. Mittler *et al.* (1998) proposed to model the eclipse phase by a distributed delay assuming that the variation among cells with respect to this intracellular delay can be approximated by a gamma distribution which can mimic a variety of biologically plausible delays. Another type of discrete or distributed delay models has been considered recently in the literature to include antiviral therapy and the effects of drugs (Nelson *et al.*, 2000; Nelson and Perelson, 2002; Wang and Li, 2006).

It was reported that HIV-1 infection is achieved in tissue culture as the lymph nodes and the brain where cell-to-cell spread would be a much more important mode of infection than cell-free viral spread (Culshaw *et al.*, 2003; Philips, 1994; Dimitrov *et al.*, 1993). To that end, Spouge *et al.* (1996) derived a model of HIV-1 cell-to-cell infection kinetics in tissue culture in terms of mathematical models and showed that its asymptotic behavior is similar to that of a model representing cell-free viral spread. The model by Spouge *et al.* (1996) does not include the latent period after cells have been infected. Culshaw *et al.* (2003) generalized the model by Spouge *et al.* (1996) by introducing a distributed delay to model the intracellular eclipse phase. They proved that the intracellular eclipse phase does change the dynamics of the model. More precisely, their results show that, when a fraction of infected cells survives the incubation period and surpasses a critical values, then a Hopf bifurcation occurs inducing periodic fluctuations in cell concentrations.

In the models by Spouge *et al.* (1996) and Culshaw *et al.* (2003), it is assumed that HIV-1 infection spreads directly from infected cells to healthy cells by neglecting the free virus and assuming that the infection rate is bilinear, i.e., dependent only on the product of the concentrations of the two cell populations. This last notion, the so-called “mass action principle”, is valid when the system is well mixed, i.e., there are no significant spatial concentration heterogeneities, and there are significant quantities of each reactant. However, interactions between two populations are not always so simple, and the form of such an expression can change, for example, to take into account at least three parameters such as the contact rate between infected and healthy cells, the fraction of healthy cells which are activated and hence susceptible to infection, and the fraction of interaction between activated cells and the virus which result in productive infected cells. In a non-well-mixed tissue culture these parameters are not constant and vary, among others, the total size of the populations. It is well known that models for T-cell proliferation are most reasonable when they are based upon a saturation function such that individual T-cells have a well-defined maximal

rate of proliferation (De Boer and Perelson, 1995; Callaway and Perelson, 1998).

In this paper we propose a new model of cell-to-cell spread of HIV-1 infection in tissue culture which takes into account the saturation effect of the proliferation of healthy cells. We assume that the infection rate between infected and healthy cells in the model by Culshaw *et al.* (2003) is a saturating function of cell concentration, i.e.,

$$\begin{aligned} \frac{dx(t)}{dt} &= rx(t) \left( 1 - \frac{x(t) + y(t)}{K} \right) - \alpha \frac{x(t)y(t)}{1 + qx(t)}, \\ \frac{dy(t)}{dt} &= \beta \int_{-\infty}^t \frac{f(t-s)x(s)y(s)}{1 + qx(s)} ds - \delta y(t), \end{aligned} \quad (1)$$

where  $x(t)$ , and  $y(t)$  denote the concentrations of healthy and infected cells at time  $t$ , respectively,  $r$  is the effective reproductive rate of healthy cells (the term is the total reproductive rate for healthy cells  $r_1$  minus the death rate for healthy cells  $\gamma$ ),  $K$  is the effective carrying capacity of the system,  $\delta$  is the death rate of infected cells,  $\alpha$  is the maximum rate of infection,  $\beta$  is such that  $\beta/\alpha$  represents the fraction of cells surviving the incubation period, and  $1/q$  denotes the half saturation constant of the proliferation process. The values of these parameters are given in Table 1.

Table 1. Assumed parameter values.

Parameters	Values	Ref.
$r$ healthy cell reproductive rate	0.68/day	[3]
$K$ carrying capacity of the system	$2 \times 10^6$ /mL	[3]
$\delta$ death rate of infected cells	0.3/day	[3]
$k$ empirical initial growth of virus	1/day	[26]
$x_0$ initial concentration of healthy cells	$5 \times 10^5$ /mL	[26]
$y_0$ initial concentration of infected cells	$5 \times 10^2$ /mL	[26]
$\frac{1}{q}$ half saturation constant	$0.45 \times 10^7$ /mL	sec. 5
$\alpha$ maximum rate of infection	$2.84 \times 10^{-6}$ /day	sec. 5

The term in the denominator of the infection rate serves to “saturate” the infection rate when  $x$  gets large. For example, if  $qx \gg 1$ , then  $\alpha xy/(1 + qx) \approx (\alpha/q)y$ . In this case the infection rate depends almost exclusively on the number of infected cells and the amount of infected cells present limits the infection rate. A similar saturation term has been used to describe cell growth in models of T cell proliferation and HIV-1 immune control (Callaway and Perelson, 1998; De Boer and Perelson, 1995; Kirschner and Webb, 1996). We point out that

as  $q \rightarrow 0$ , the system (1) is reduced to that described by Culshaw *et al.* (2003). The model of Culshaw *et al.* (2003) with the simple mass-action infection term is then an extreme case of the HIV-1 model (1) in this paper. The initial values of the system are

$$x(s) = \phi(s), \quad y(s) = \psi(s), \quad s \in (-\infty, 0],$$

with  $\phi, \psi \in X$ , where  $X = C((-\infty, 0]; \mathbb{R}_+)$  is the Banach space of non-negative, continuous and bounded functions on  $(-\infty, 0]$ . The distributed delay in the second equation of the system (1) models the incubation period by assuming that cells which are productively infectious at time  $t$  were infected  $\tau$  time units ago, where  $\tau$  is the average delay defined by (MacDonald, 1978)

$$\tau = \int_0^\infty sf(s) ds.$$

In the rest of this paper we take as a delay kernel  $f(s)$  the gamma distribution function of order 0 called a weak kernel,

$$f(s) = \mu e^{-\mu s}, \quad s \geq 0,$$

for which the average delay is  $\tau = 1/\mu$ . Such kernels were used in mathematical models of HIV-1 infections by Mittler *et al.* (1998).

Setting

$$z(t) = \int_{-\infty}^t \mu e^{-\mu(t-s)} \frac{x(s)y(s)}{1+qx(s)} ds,$$

we obtain the following system of three ordinary differential equations:

$$\begin{aligned} \frac{dx(t)}{dt} &= rx(t) \left( 1 - \frac{x(t)+y(t)}{K} \right) - \alpha \frac{x(t)y(t)}{1+qx(t)}, \\ \frac{dy(t)}{dt} &= \mu z'(t) - \delta y(t), \\ \frac{dz'(t)}{dt} &= \beta \frac{x(t)y(t)}{1+qx(t)} - \mu z'(t), \end{aligned} \tag{2}$$

with the initial values

$$\begin{aligned} x_0 &= \phi(0) \geq 0, \quad y_0 = \psi(0) \geq 0, \\ z_0 &= \chi(0) = \int_{-\infty}^0 \mu e^{\mu s} \frac{\phi(s)\psi(s)}{1+q\phi(s)} ds \geq 0, \end{aligned} \tag{3}$$

where

$$\chi(s) = \int_{-\infty}^s \mu e^{\mu s} \frac{\phi(s)\psi(s)}{1+q\phi(s)} ds,$$

$s \leq 0$  and  $\phi, \psi \in X$ .

To find equilibria of system (2) we set

$$\begin{aligned} rx \left( 1 - \frac{x+y}{K} \right) - \alpha \frac{xy}{1+qx} &= 0, \\ \beta z - \delta y &= 0, \\ \mu \frac{xy}{1+qx} - \mu z &= 0. \end{aligned} \tag{4}$$

It is easy to see that the algebraic system (4) has three equilibria: the trivial equilibrium  $E_0 = (0, 0, 0)$ , the healthy equilibrium  $E_1 = (K, 0, 0)$ , and, when

$$\frac{\beta K}{\delta(1+qK)} > 1,$$

there is a third positive equilibrium  $E^* = (x^*, y^*, z^*)$ , called the infected equilibrium, given by

$$\begin{aligned} x^* &= \frac{\delta}{\beta - q\delta}, \\ y^* &= \frac{r\beta(K\beta - qK\delta - \delta)}{(\beta - q\delta)(\beta r + K\alpha\beta - qK\alpha\delta)}, \\ z^* &= \frac{r\delta(K\beta - qK\delta - \delta)}{(\beta - q\delta)(\beta r + K\alpha\beta - qK\alpha\delta)}. \end{aligned}$$

In the following section we derive the basic reproduction number  $R_0$  of the model (2). In Section 3 we prove that, when  $R_0 < 1$ , the healthy equilibrium  $E_1$  is locally asymptotically stable, while  $E_0$  is a saddle point. When  $R_0 > 1$ , we prove that the infected equilibrium  $E^*$  exists and is locally asymptotically stable under some hypotheses on the parameters. In Section 4 we are interested in the global dynamics of the model (2). We prove that, when  $R_0 \leq 1$ , the healthy equilibrium  $E_1$  is in fact globally asymptotically stable, and if  $R_0 > 1$ , we prove that the infection is persistent. By using the average delay as a bifurcation parameter, we prove in this case that the system (2) exhibits two Hopf bifurcations  $\mu_1$  and  $\mu_2$ , such that the infected equilibrium  $E^*$  is locally asymptotically stable for  $\mu \in (0, \mu_1) \cup (\mu_2, \infty)$ , and periodic oscillations occur when  $\mu \in (\mu_1, \mu_2)$ . In Section 5, we report some numerical simulations to illustrate the stability and bifurcation results. As in the work of Culshaw *et al.* (2003), our model exhibits infective periodic oscillations in realistic parameter regimes, but in our case there are two Hopf bifurcations and the infected steady state is locally asymptotically stable either if the average delay is small enough or sufficiently large.

## 2. Basic reproduction number $R_0$

To see whether the virus establishes an infection, a crucial quantity is the basic reproduction number  $R_0$ , which is defined as the number of newly infected cells that arise from any infected cell, over its entire infectious period  $1/\delta$ , when most cells are uninfected (Nowak and May, 2000). In epidemiological models,  $R_0$  is defined as the threshold parameter, such that the disease dies out irrespective of the initial number cases when  $R_0 \leq 1$ , and is persistent with the existence of an infected equilibrium when  $R_0 > 1$ . In this paper, following Diekmann *et al.* (1990), we derive the basic reproduction number  $R_0$  for the model (2) by using the method of the next generation matrix, which was modified later by Van den Driessche and Watmough

(2002), as well as Esteva-Peralta and Velasco-Hernandez (2002).

In order to compute  $R_0$ , we regard (2) as an epidemiological model with three compartments:  $x$ ,  $y$  and  $z$ . We first write the system (2) the following form:

$$\begin{aligned} \frac{dX}{dt} &= F(X) - V(X), \\ V(X) &= V^-(X) - V^+(X), \\ X &= (x, y, z)^T, \end{aligned}$$

where  $F, V^+, V^- : \mathbb{R}^3 \rightarrow \mathbb{R}^3$  are defined by

$$\begin{aligned} F(X) &= \left( 0, 0, \mu \frac{xy}{1+qx} \right), \\ V^+(X) &= \left( r_1 x \left\{ 1 - \frac{x+y}{K} \right\}, \beta z, 0 \right) \\ V^-(X) &= \left( \alpha \frac{xy}{1+qx} + \gamma x \left( 1 - \frac{x+y}{K} \right), \delta y, \mu z \right). \end{aligned}$$

$F(X)$  is the rate of the appearance of new infected cells in each compartment,  $V^+$  is the rate of the transfer of cells into each compartment by all other means, and  $V^-$  is the rate of the transfer of cells out of the compartments. Denote by  $\partial F/\partial x$ ,  $\partial V/\partial x$  the Jacobian matrices of  $F$  and  $V$  at  $E_1 = (K, 0, 0)$ , respectively. We have

$$\begin{aligned} \frac{\partial F}{\partial x} &= \begin{pmatrix} 0 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & \frac{\mu K}{1+qK} & 0 \end{pmatrix}, \\ \frac{\partial V}{\partial x} &= \begin{pmatrix} r_1 + \gamma & r_1 + \frac{\alpha K}{1+qK} & 0 \\ 0 & \delta & -\beta \\ 0 & 0 & \mu \end{pmatrix}. \end{aligned}$$

According to (Diekmann *et al.*, 1990) we call  $\frac{\partial F}{\partial x} \left( \frac{\partial V}{\partial x} \right)^{-1}$  the next generation matrix of the model (2) and its spectral radius is the corresponding basic reproduction number  $R_0$ . We have

$$\frac{\partial F}{\partial x} \left( \frac{\partial V}{\partial x} \right)^{-1} = \begin{pmatrix} 0 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & \frac{\mu K}{\delta(1+qK)} & \frac{\beta K}{\delta(1+qK)} \end{pmatrix},$$

and its characteristic polynomial is given by

$$Q(\lambda) = \lambda^2 \left( \lambda - \frac{\beta K}{\delta(1+qK)} \right).$$

The corresponding eigenvalues are

$$\lambda_{1,2} = 0, \quad \lambda_3 = \frac{\beta K}{\delta(1+qK)}.$$

The spectral radius of the next generation matrix  $\frac{\partial F}{\partial x} \left( \frac{\partial V}{\partial x} \right)^{-1}$  is then  $\beta K/\delta(1+qK)$ . According to Diekmann *et al.* (1990), we define the basic reproduction number  $R_0$  for model (2) by

$$R_0 = \frac{1}{\delta} \left[ \frac{\beta K}{1+qK} \right].$$

The term in brackets describes the newly infected cells that arise from any infected cell when almost all cells are uninfected. We will prove in Section 4 that the dynamics of the system (2) are largely determined by the basic reproduction number  $R_0$ .

### 3. Local stability of equilibria

In this section we analyze the local stability of the equilibria of the system (2). Denote by  $J(E)$  the Jacobian matrix of the system (2) at  $E = (x, y, z)$ . We have

$$J(E) = \begin{pmatrix} r - 2r \frac{x}{K} - r \frac{y}{K} - \frac{\alpha y}{(1+qx)^2} & & & & & \\ & 0 & & & & \\ & \frac{\mu y}{(1+qx)^2} & & & & \\ & & -r \frac{x}{K} - \alpha \frac{x}{1+qx} & & 0 & \\ & & -\delta & & & \beta \\ & & \mu \frac{x}{1+qx} & & & -\mu \end{pmatrix}.$$

**Theorem 1.**  $E_0$  is a saddle point and  $E_1$  is locally asymptotically stable provided that  $R_0 < 1$ .

*Proof.* Denote by  $P(\lambda)$  the characteristic polynomial of the system (2) at  $(x, y, z)$ . We have, after some calculations, that

$$\begin{aligned} P(\lambda) &= \lambda^3 + (\delta + \mu - A)\lambda^2 + \left( \delta\mu - A\delta - A\mu \right. \\ &\quad \left. - \beta \frac{\mu x}{1+qx} \right)\lambda + \frac{A\beta\mu x}{1+qx} \\ &\quad + B \frac{\beta\mu y}{(1+qx)^2} - A\mu\delta, \end{aligned} \tag{5}$$

where

$$\begin{aligned} A &= r - \frac{r}{K}y - 2\frac{r}{K}x - \frac{\alpha y}{(1+qx)^2}, \\ B &= \frac{r}{K}x + \frac{\alpha x}{1+qx}. \end{aligned}$$

Case 1. Trivial equilibrium  $E_0 = (0, 0, 0)$ :

Set  $x = y = z = 0$ . The characteristic polynomial (5) takes the form

$$\begin{aligned} P(\lambda) &= \lambda^3 + (\delta + \mu - r)\lambda^2 + (\delta\mu - r\delta - r\mu)\lambda \\ &\quad - r\mu\delta \\ &= (r - \lambda)(\lambda + \mu)(\lambda + \delta), \end{aligned}$$

and therefore  $P$  has two negative real roots  $\lambda_1 = -\delta$ ,  $\lambda_2 = -\mu$  and one positive real root  $\lambda_3 = r > 0$ . The trivial equilibrium is a saddle point.

Case 2. Healthy equilibrium  $E_1 = (K, 0, 0)$ :

In this case the characteristic polynomial has the form

$$\begin{aligned}
 P(\lambda) &= \lambda^3 + (r + \mu + \delta)\lambda^2 + (r\delta + r\mu \\
 &\quad + \mu \left( \delta - \frac{\beta K}{1 + qK} \right)) \lambda \\
 &\quad + r\mu \left( \delta - \frac{\beta K}{1 + qK} \right) \\
 &= \lambda^3 + (r + \mu + \delta)\lambda^2 \\
 &\quad + \left( r\delta + r\mu + \frac{\mu}{\delta} (1 - R_0) \right) \lambda \\
 &\quad + r\frac{\mu}{\delta} (1 - R_0).
 \end{aligned}$$

Since  $R_0 < 1$ , all the coefficients of  $P(\lambda)$  are positive, and the inequality

$$(r + \mu + \delta) \left[ r\delta + r\mu + \frac{\mu}{\delta} (1 - R_0) \right] > r\frac{\mu}{\delta} (1 - R_0)$$

holds. So by the Routh–Hurwitz theorem all the roots of  $P(\lambda)$  have negative real parts. The healthy equilibrium  $E_1$  is locally asymptotically stable. ■

We now turn to proving the local stability of the infected equilibrium  $E^*$ . We showed in Section 2 that, if  $R_0 > 1$ , then the infected equilibrium exists. In the following theorem we give some hypotheses on the parameters under which  $E^*$  is locally asymptotically stable.

The Jacobian matrix  $J(E^*)$  of (2) at point  $E^* = (x^*, y^*, z^*)$  is

$$J(E^*) = \begin{pmatrix} -a & -b & 0 \\ 0 & -\delta & \beta \\ \frac{\mu y^*}{(1 + qx^*)^2} & \mu \frac{\delta}{\beta} & -\mu \end{pmatrix},$$

where

$$\begin{aligned}
 a &= -r + y^* \frac{r}{K} + 2 \frac{r}{K} x^* + \frac{\alpha y^*}{(1 + qx^*)^2}, \\
 b &= r \frac{x^*}{K} + \alpha \frac{\delta}{\beta}.
 \end{aligned}$$

To proceed further, we make the following assumption on the parameters of the system (2):

$$r - 2r \frac{x^*}{K} - r \frac{y^*}{K} - \frac{\alpha y^*}{(1 + qx^*)^2} < 0. \tag{6}$$

Write

$$\begin{aligned}
 \Delta &= \left[ a - \frac{b}{a} \frac{\beta y^*}{(1 + qx^*)^2} \right]^2 - 4\delta \frac{b}{a} \frac{\beta y^*}{(1 + qx^*)^2}, \\
 D &= 2\delta + a - \frac{b}{a} \frac{\beta y^*}{(1 + qx^*)^2}.
 \end{aligned}$$

**Theorem 2.** Assume that  $R_0 > 1$ , and (6) holds.

(i) If either  $\Delta < 0$  or  $\Delta = 0$  and  $D > 0$ , then  $E^*$  is

locally asymptotically stable for all  $\mu > 0$ .

(ii) If  $\Delta = 0$  and  $D < 0$ , there is a  $\mu_0 > 0$ , such that  $E^*$  is locally asymptotically stable for  $\mu > 0$  and  $\mu \neq \mu_0$  and unstable for  $\mu = \mu_0$ .

(iii) If  $\Delta > 0$ , we have the following two cases:

1.  $D > 0$ , then  $E^*$  is locally asymptotically stable for all  $\mu > 0$ .
2.  $D < 0$ , then there are  $0 < \mu_1 < \mu_2$ , such that  $E^*$  is locally asymptotically stable for  $\mu < \mu_1$  or  $\mu > \mu_2$ , and is unstable for  $\mu \in (\mu_1, \mu_2)$ . Further, the system (2) exhibits Hopf bifurcations at  $\mu = \mu_1$  and  $\mu = \mu_2$ .

*Proof.* Let  $P(\lambda)$  be the characteristic polynomial of the Jacobian matrix  $J(E^*)$ . We have

$$\begin{aligned}
 P(\lambda) &= \lambda^3 + (a + \mu + \delta)\lambda^2 + (a\mu + a\delta)\lambda \\
 &\quad + b \frac{\mu \beta y^*}{(1 + qx^*)^2}.
 \end{aligned} \tag{7}$$

From (6) it follows that  $a > 0$  and all the coefficients of  $P(\lambda)$  are positive. Applying the Routh–Hurwitz theorem to (7), we find that  $E^*$  is locally asymptotically stable if and only if

$$\begin{aligned}
 A(\mu) &= (a + \delta + \mu)(a\delta + a\mu) \\
 &\quad - b \frac{\mu \beta y^*}{(1 + qx^*)^2} > 0.
 \end{aligned} \tag{8}$$

Taking  $\mu > 0$  as a bifurcation parameter, we rewrite (8) into the following form:

$$\begin{aligned}
 A(\mu) &= \mu^2 + \left[ 2\delta + a - \frac{b}{a} \frac{\beta y^*}{(1 + qx^*)^2} \right] \mu + \delta^2 + a\delta \\
 &= \mu^2 + D\mu + \delta^2 + a\delta,
 \end{aligned}$$

for which the discriminant is  $\Delta$ . We then have the following cases:

- If  $\Delta < 0$ , or  $\Delta = 0$  and  $D > 0$ , then  $A(\mu)$  has no real roots or one negative real root, and in this case  $A(\mu) > 0$  while  $E^*$  is locally asymptotically stable for all  $\mu > 0$ .
- If  $\Delta = 0$  and  $D < 0$ ,  $A(\mu)$  has one positive real root  $\mu_0$  and  $A(\mu) > 0$  for all  $\mu > 0$  and  $\mu \neq \mu_0$ .  $E^*$  is then locally asymptotically stable for all  $\mu > 0$ , and  $\mu \neq \mu_0$  and unstable for  $\mu = \mu_0$ .
- Assume now that  $\Delta > 0$ . Then, if  $D > 0$ ,  $A(\mu)$  has two negative real roots, so  $A(\mu) > 0$ . In this case the infected equilibrium  $E^*$  is locally asymptotically stable for all  $\mu > 0$ .
- When  $D < 0$ ,  $A(\mu)$  has two positive real roots  $0 < \mu_1 < \mu_2$  such that  $A(\mu) > 0$ , for  $\mu \in (0, \mu_1) \cup$

$(\mu_2, +\infty)$  and  $A(\mu) < 0$ , for  $\mu \in (\mu_1, \mu_2)$ .  $E^*$  is then locally asymptotically stable for  $\mu \in (0, \mu_1) \cup (\mu_2, +\infty)$  and unstable for  $\mu \in (\mu_1, \mu_2)$ . Further,

$$\begin{aligned} \frac{dA(\mu_1)}{d\mu} &= 2\mu_1 + D = \sqrt{\Delta} > 0, \\ \frac{dA(\mu_2)}{d\mu} &= 2\mu_2 + D = -\sqrt{\Delta} < 0. \end{aligned} \tag{9}$$

Let  $\lambda(\mu) = \eta(\mu) + i\omega(\mu)$  be the eigenvalue of  $P(\lambda)$ , where  $\eta(\mu)$  and  $\omega(\mu)$  depend on  $\mu$ . We first show that for  $\mu = \mu_{1,2}$ ,  $P(\lambda)$  has one negative real root and two conjugate purely imaginary roots. Since

$$(a + \delta + \mu_{1,2})(a\delta + a\mu_{1,2}) = b \frac{\mu_{1,2}\beta y^*}{(1 + qx^*)^2},$$

we have

$$P(\lambda) = (\lambda + a + \delta + \mu_{1,2})(\lambda^2 + a\delta + a\mu_{1,2}).$$

Hence  $P(\lambda)$  has one negative real root  $\lambda_1 = -(a + \delta + \mu_{1,2}) < 0$  and two conjugate purely imaginary roots  $\lambda_{2,3} = \pm i\sqrt{a\delta + a\mu_{1,2}}$ . Substituting  $\lambda(\mu) = \eta(\mu) + i\omega(\mu)$  into (7), we find that

$$\begin{aligned} \eta^3 - 3\eta\omega^2 + 3i\eta^2\omega - i\omega^3 + (a + \delta + \mu)(\eta^2 - \omega^2) \\ + 2i(a + \delta + \mu)\eta\omega + \eta(a\delta + a\mu) + i\omega(a\delta + a\mu) \\ + b \frac{\mu\beta y^*}{(1 + qx^*)^2} = 0. \end{aligned}$$

Separating the real and imaginary parts, we obtain

$$\begin{aligned} \eta^3 - 3\eta\omega^2 + (a + \delta + \mu)(\eta^2 - \omega^2) + \eta(a\delta + a\mu) \\ + b \frac{\mu\beta y^*}{(1 + qx^*)^2} = 0. \end{aligned} \tag{10}$$

Differentiating the expression in (10) with respect to  $\mu$  and using the fact that  $\eta(\mu_{1,2}) = 0$  and  $\omega^2(\mu_{1,2}) = a\delta + a\mu_{1,2}$ , we get

$$\begin{aligned} -3(a\delta + a\mu_{1,2}) \frac{d\eta}{d\mu} \Big|_{\mu=\mu_{1,2}} \\ - \frac{d}{d\mu} (a\delta + a\mu)(a + \delta + \mu) \Big|_{\mu=\mu_{1,2}} \\ + \frac{d}{d\mu} \left( b \frac{\mu\beta y^*}{(1 + qx^*)^2} \right) \Big|_{\mu=\mu_{1,2}} = 0. \end{aligned}$$

Hence we obtain, with the help of (8) and (9),

$$\begin{aligned} -3(a\delta + a\mu_{1,2}) \frac{d\eta}{d\mu} \Big|_{\mu=\mu_{1,2}} &= \frac{dA(\mu)}{d\mu} \Big|_{\mu=\mu_{1,2}} \\ &= \pm\sqrt{\Delta} \neq 0, \end{aligned}$$

thus a family of periodic solutions bifurcates from  $E^*$  when  $\mu$  passes through one of the critical values  $\mu_1$  or  $\mu_2$  (Hassard *et al.*, 1981). ■

#### 4. Global dynamics of the model

In this section we analyze the global asymptotic behavior of the system (2). We begin with the following lemma on the dissipativity of (2).

**Lemma 1.** *The positive orthant is positively invariant by the system (2) and there exists a constant  $M > 0$  independent of the solutions such that*

$$\begin{aligned} \limsup_{t \rightarrow +\infty} x(t) &\leq K, & \limsup_{t \rightarrow +\infty} y(t) &\leq M, \\ \limsup_{t \rightarrow +\infty} z(t) &\leq M. \end{aligned}$$

*Proof.* The invariance of the positive orthant is trivial. We have to prove the ultimate boundedness of the solutions. We have, from the first equation of the system (2),

$$\frac{dx(t)}{dt} \leq rx(t) \left( 1 - \frac{x(t)}{K} \right),$$

which gives, with the help of the comparison principle,

$$\limsup_{t \rightarrow +\infty} x(t) \leq K. \tag{11}$$

Define the function

$$W(t) = x(t) + \frac{\alpha}{\mu} z(t).$$

We have

$$\begin{aligned} \frac{dW}{dt} &\leq rx - \alpha z \\ &\leq (r + \mu)x - \mu x - \alpha z \\ &\leq (r + \mu)x - \mu W(t). \end{aligned} \tag{12}$$

Let  $\epsilon > 0$  be fixed. By (11), there is  $T^\epsilon > 0$  such that  $x(t) \leq K + \epsilon$  for  $t > T^\epsilon$ . Combining this last relation with (12), for  $t > T^\epsilon$  we have

$$\frac{dW}{dt}(t) \leq (r + \mu)(K + \epsilon) - \mu W(t),$$

which leads to

$$\limsup_{t \rightarrow +\infty} W(t) \leq \frac{K(r + \mu)}{\mu}.$$

Finally, from the second equation in (2) and the fact that  $z$  is uniformly bounded it follows that there is an

$$M > \frac{K(r + \mu)}{\mu}$$

such that

$$\limsup_{t \rightarrow +\infty} y(t) \leq M.$$

This completes the proof of the lemma. ■

In Section 3 we proved that, if  $R_0 < 1$ , the healthy equilibrium  $E_1$  is locally asymptotically stable. In the following theorem we prove that  $E_1$  is globally asymptotically stable when  $R_0 \leq 1$ .

**Theorem 3.** *Assume that  $R_0 \leq 1$ . Then the healthy equilibrium  $E_1$  is globally stable.*

*Proof.* We analyze separately two cases:  $R_0 < 1$  and  $R_0 = 1$ .

Case 1.  $R_0 < 1$

Define the set  $G = \{(\phi, \psi, \chi) \in X : 0 \leq \phi \leq K, \psi, \chi \geq 0\}$ . By Lemma 1,  $G$  attracts all solutions of the system (2). Let  $(\phi, \psi, \chi) \in G$ , and denote by  $(x(t), y(t), z(t))$  the corresponding solution of (2). We claim that  $x(t) \leq K$  for all  $t > 0$ . Otherwise, there is  $t_1 > 0$  such that  $x(t_1) = K$  and  $dx(t_1)/dt > 0$ . We have

$$\begin{aligned} \frac{dx}{dt}(t_1) &\leq rx(t_1) \left(1 - \frac{x(t_1) + y(t_1)}{K}\right) \\ &\leq -ry(t_1) \\ &\leq 0, \end{aligned}$$

which contradicts the fact that  $dx(t_1)/dt > 0$ . Thus  $G$  is positively invariant with respect to (2). For any  $(\phi, \psi, \chi) \in G$ , we define the functional  $V(\phi, \psi, \chi) = \mu y + \beta z$ , where  $(x(t), y(t), z(t))$  is the corresponding solution to (2). From the system (2) we have that

$$\frac{dV}{dt} = \left(\beta \frac{\mu x}{1 + qx} - \delta \mu\right) y. \quad (13)$$

Since  $G$  is positively invariant and  $R_0 < 1$ , we have  $x(t) \leq K$  for all  $t > 0$ . Hence (13) leads to

$$\begin{aligned} \frac{dV}{dt} &\leq \mu \left\{ \frac{\beta K}{1 + qK} - \delta \right\} y \\ &= \frac{\mu}{\delta} (R_0 - 1) y < 0, \quad t > 0. \end{aligned} \quad (14)$$

$V$  is then a Lyapunov functional on  $G$ . Define now the set

$$E = \{(\phi, \psi, \chi) \in G : \frac{dV}{dt}(\phi, \psi, \chi) = 0\},$$

and let  $M$  be the largest set in  $E$  which is invariant with respect to (2). Clearly,  $M$  is not empty since  $E_1 = (K, 0, 0) \in M$ . Let  $(\phi, \psi, \chi) \in M$  and  $(x(t), y(t), z(t))$  be the solution of (2) with the initial data  $(\phi, \psi, \chi)$ . From the invariance of  $M$ , we have  $(x_t, y_t, z_t) \in M$  for all  $t \in R$ . Thus  $dV/dt = 0$ , and, from (13),  $y(t) = 0$  for all  $t \in R$ . We obtain from the first and the third equations in (2) that  $x(t) \rightarrow K$  and  $z(t) \rightarrow 0$  as  $t \rightarrow +\infty$ . The invariance of  $M$  implies that  $x(t) = K$  and  $z(t) = 0$  for all  $t \in R$ . Therefore,  $M = \{(K, 0, 0)\} = E_1$ . Since it was shown in Theorem 1 that  $E_1$  is locally asymptotically stable when  $R_0 < 1$ , from the LaSalle invariance principle it follows that  $E_1$  is globally asymptotically stable.

Case 2.  $R_0 = 1$

By the first equation of (2),  $x(t)$  is always decreasing when above  $K$ . We can prove that if there exists some  $t_0 > 0$  such that  $x(t_0) < K$ , then  $x(t) < K$  for all  $t > t_0$ . Otherwise, there must exist some  $t_1 > t_0$  such that  $x(t_1) = K$  and  $dx(t_1)/dt \geq 0$ . This is impossible. Hence, there are two possible cases:

- (i)  $x(t) > K$ , for all  $t > 0$  and  $x(t) \rightarrow K$  as  $t \rightarrow +\infty$ .
- (ii) There exists  $t_0 > 0$  such that  $x(t_0) < K$ .

In Case (ii) we have that  $x(t) < K$  for all  $t > t_0$ . The global stability of  $E_1$  can be proved by using the same Lyapunov functional  $V$  as in the first part of the proof.

In Case (i) we rewrite the first equation in (2) as follows:

$$r \frac{xy}{K} \leq rx \left(1 - \frac{x}{K}\right) - \frac{dx}{dt}.$$

Now, since  $x(t) > K$ , from the preceding equation we obtain

$$ry(t) \leq -\frac{dx}{dt}, \quad t > 0.$$

Integrating the last relation over  $(0, t)$ , we find that

$$r \int_0^t y(s) ds \leq x(0) - x(t) \leq x(0), \quad t > 0,$$

which implies, with the boundedness of  $y(t)$ , that  $y(t) \rightarrow 0$ , and thus  $z(t) \rightarrow 0$  as  $t \rightarrow +\infty$ .  $E_1$  is then globally stable. ■

Theorem 3 completely determines the global dynamics of the system when  $R_0 \leq 1$  and can be interpreted as follows: If  $R_0 \leq 1$ , then the infection will not spread since every infected cell will on the average produce less than one new infected cell. The chain reaction is sub-critical. In the following theorem, we will prove that, if  $R_0 > 1$ , then a chain reaction takes place and leads to a chronic infection.

Before giving the second main result of this section, we recall some results about the theory of infinite dimensional systems (Thieme, 1993). Let  $X$  be a metric space with metric  $d$ . Assume that  $X$  is the union of two disjoint subsets  $X_1$  and  $X_2$ . A continuous semiflow on  $X_1$  is a continuous mapping  $\Phi : [0, \infty[ \times X_1 \rightarrow X_1$  such that

$$\Phi_t \circ \Phi_s = \Phi_{t+s}, \quad t, s \geq 0, \quad \Phi_0(x) = x, \quad \forall x \in X_1,$$

where  $\Phi_t$  denotes the mapping from  $X_1$  to  $X_1$  given by

$$\Phi_t(x) = \Phi(t, x).$$

We say that a subset  $K$  of  $X$  is a weak repeller of  $X_1$  if there is an  $\epsilon > 0$  such that

$$\limsup_{t \rightarrow \infty} d(\Phi_t(x), K) > \epsilon, \quad \forall x \in X_1.$$

$K$  is called a uniform strong repeller for  $X_1$  if there is an  $\epsilon > 0$  such that

$$\liminf_{t \rightarrow \infty} d(\Phi_t(x), K) > \epsilon, \quad \forall x \in X_1.$$

**Lemma 2.** (Thieme, 1993) Let  $X_1$  be open in  $X$  and the semiflow  $\Phi$  be dissipative, i.e, there is a bounded set  $B \subset X$  such that  $d(\Phi_t(x), B) \rightarrow 0$  for all  $x \in X_1$ . Further, assume that  $A_\partial$ ,

$$A_\partial = \bigcup_{y \in Y} \omega(y),$$

$$Y = \{x \in X_2 : \Phi_t(x) \in X_2, \forall t > 0\},$$

has an acyclic isolated covering  $M = \bigcup_{k=1}^m M_k$  such that each  $M_k$  is a weak repeller for  $X_1$ . Then  $X_2$  is a uniform strong repeller for  $X_1$ .

**Theorem 4.** Assume that  $R_0 > 1$ . Then the infection is persistent.

*Proof.* Recall that solutions of the system (2) are said to be persistent (or uniformly persistent) if there is an  $\epsilon > 0$  such that, for all solutions  $(x(t), y(t), z(t))$  of (2) for which  $x(0), y(0), z(0) > 0$ , we have

$$\liminf_{t \rightarrow +\infty} x(t) \geq \epsilon, \quad \liminf_{t \rightarrow +\infty} y(t) \geq \epsilon, \quad \liminf_{t \rightarrow +\infty} z(t) \geq \epsilon.$$

To this end we use the persistence theory for infinite dimensional systems (Thieme, 1993). In order to apply Theorem 4 of (Thieme, 1993), we define the sets

$$X = \{(x, y, z) \in \mathbb{R}^3 : x, y, z \geq 0\}, \quad X_1 = \text{int}(X),$$

$$X_2 = \{(x, y, z) \in \mathbb{R}^3 : x = 0 \text{ or } z = 0\}.$$

We have  $X = X_1 \cup X_2$  and  $X_1 \cap X_2 = \emptyset$ . Let  $U = \{(x, y, z) \in X : x \leq K, y \leq M, z \leq M\}$ . From Lemma 1, it is easy to see that  $U$  is a compact absorbing set. The semiflow associated with the system (2) is then dissipative. Denote by  $\omega(x_0, y_0, z_0)$  the omega limit set of the solution to the system (2) starting at  $(x_0, y_0, z_0) \in X$  (which exists by Lemma 1). We have to determine the following set:

$$A_\partial = \bigcup_{(x_0, y_0, z_0) \in Y} \omega(x_0, y_0, z_0),$$

where

$$Y = \{(x_0, y_0, z_0) \in X_2 : \Phi_t(x_0, y_0, z_0) \in X_2, \forall t > 0\},$$

in which  $\Phi_t(x_0, y_0, z_0)$  denotes the solution semiflow of the system (2) starting at  $(x_0, y_0, z_0)$ . It follows from Lemma 1 that  $X_1$  is an invariant set with respect to the semiflow of the system (2).

Let  $(x_0, y_0, z_0) \in \partial X_2$ . There are two distinct cases:

1.  $x_0 = 0$ . In this case we have, from the first equation of

the system (2),

$$\frac{dx}{dt} \leq rx \left(1 - \frac{x}{K}\right), \quad x(0) = 0,$$

which implies that  $x(t) = 0$ , for all  $t > 0$ . The third equation of the system (2) gives  $z(t) \rightarrow 0$  as  $t \rightarrow +\infty$ , and hence  $y(t) \rightarrow 0$  as  $t \rightarrow +\infty$ . So the omega limit set becomes  $\omega(x_0, y_0, z_0) = \{E_0\}$ .

2.  $z_0 = 0$ . Then, by (3), either  $x(0) = 0$  or  $y(0) = 0$ . In the first case, we have  $x_0 = 0$ , which implies, according to the first part of the proof, that  $\omega(x_0, y_0, z_0) = \{E_0\}$ . If  $y(0) = 0$ , then by the second and third equations of the system (2) we obtain  $y(t), z(t) \rightarrow 0$  as  $t \rightarrow +\infty$ , and by the first one  $x(t) \rightarrow K$  as  $t \rightarrow +\infty$ . Thus  $\omega(x_0, y_0, z_0) = \{E_1\}$ . Hence  $A_\partial = \{E_0\} \cup \{E_1\}$  and  $\{E_0\}, \{E_1\}$  are an acyclic isolated covering for  $A_\partial$ .

It remains to prove that  $\{E_0\}, \{E_1\}$  are weak repellers for  $X_1$ , i.e.,  $W^s(E_i) \cap X_1 = \emptyset$  for  $i = 0, 1$ , where

$$W^s(E_i) = \{(x_0, y_0, z_0) \in X : \omega(x_0, y_0, z_0) = E_i\}$$

is the stable manifold of  $E_i$ . Assume that  $W^s(E_0) \cap X_1 \neq \emptyset$ . Then there is a solution  $(x(t), y(t), z(t))$  to the system (2) such that  $(x(t), y(t), z(t)) \rightarrow (0, 0, 0)$  as  $t \rightarrow +\infty$ . Hence from the first equation of the system (2), there is a  $T_0 > 0$  such that

$$\frac{d(\ln x)}{dt} \geq r \left\{1 - \frac{x}{K}\right\} - \alpha \frac{y}{1 + qx} \geq \frac{r}{2}, \quad t > T_0,$$

which implies that  $x(t) \rightarrow +\infty$  as  $t \rightarrow +\infty$ , which is a contradiction.

In the second case assume, on the contrary, that there holds  $W^s(E_1) \cap X_1 \neq \emptyset$ . There is then a solution  $(x(t), y(t), z(t))$  of the system (2) such that  $(x(t), y(t), z(t)) \rightarrow (K, 0, 0)$  as  $t \rightarrow +\infty$ . Since  $R_0 > 1$ , there are  $\epsilon > 0$  and  $T_1 > 0$  such that

$$x(t) > K - \epsilon, \quad t > T_1,$$

$$\frac{\beta(K - \epsilon)}{1 + q(K - \epsilon)} > \delta. \tag{15}$$

From the system (2) and the first equation in (15), we have

$$\frac{dy}{dt} = \beta z - \delta y,$$

$$\frac{dz}{dt} \geq \mu \frac{(K - \epsilon)}{1 + q(K - \epsilon)} y - \mu z.$$

The second inequality in (15) implies

$$\frac{d}{dt} (\mu y(t) + \beta z(t))$$

$$= \mu y(t) \left( \frac{\beta(K - \epsilon)}{1 + q(K - \epsilon)} - \delta \right) > 0,$$

for all  $t > T_1$ . We deduce that  $\mu y(t) + \beta z(t) > \mu y(T_1) + \beta z(T_1) > 0$  for all  $t > T_1$ , which contradicts the fact that  $y(t), z(t) \rightarrow 0$  as  $t \rightarrow +\infty$ . By Lemma 2,  $X_2$  is a uniform strong repeller for  $X_1$ . This means that solutions of system (2) are uniformly persistent. ■





which are the maximal infected rate  $\alpha$  and the half saturation constant of the proliferation process  $1/q$ . There are no empirical values of these parameters in the literature. However, we propose their estimates. Firstly, we use the fact that the basic reproduction number  $R_0$ , which represents the number of newly infected cells produced by one infected cell at the beginning of the infection, can be computed by using the parameter values given by Culshaw *et al.* (2003). If we assume that all cells survive the incubation period, we find that  $R_0$  is around 13. We assume that this number is approximately the same as in our model. Further, by Section 2 and the assumption that all cells survive the incubation period (i.e.,  $\alpha = \beta$ ), we know that  $R_0 = \frac{\alpha K}{\delta(1+qK)}$ , and then

$$\frac{\alpha K}{\delta(1+qK)} \simeq 13. \tag{17}$$

Secondly, it is well known that, for HIV-1 cell-to-cell spread, the initial exponential growth of infected cells  $k$  is a linear function of the initial concentration of infected cells (Spouge *et al.*, 1996; Dimitrov *et al.*, 1993) and is around 1 per day. Since the exponential growth of infected cells at time  $t = 0$  is given by  $y_0^{-1}dy(0)/dt$ , we obtain from the second equation in (2) that  $k \simeq \alpha z_0/y_0 - \delta$ . If we assume now that the initial concentrations of healthy and infected cells are constants around  $x_0 = 5 \times 10^5$  cells/mL and  $y_0 = 5 \times 10^2$  cells/mL, respectively, we obtain that

$$k \simeq \frac{\alpha x_0}{1+qx_0} - \delta. \tag{18}$$

Knowing  $K$ ,  $x_0$ ,  $\delta$  and  $k$  (see Table 1) we can compute  $q$  and  $\alpha$  from Eqns. (17) and (18), and we find that  $q = 2.2 \times 10^{-7}/\text{mL}$  and  $\alpha = 2.84 \times 10^{-6}/\text{day}$ .

Suppose that 50% of infected cells survive incubation, which corresponds to  $\beta = 1.42 \times 10^{-6}$ . Then  $R_0 = 6.25$ ,  $a = 0.05$ ,  $\Delta = 7.33$ ,  $D = -2.78$ . The second condition of Part (iii) in Theorem 2 is then satisfied. The critical parameters are  $\mu_1 = 0.038$ ,  $\mu_2 = 2.74$ . If we take as the average delay  $\mu = 1$ , which is a realistic value since the incubation period is around 1 day (Spouge *et al.*, 1996), then Corollary 1 predicts that the solution to the system (1) approaches a periodic solution. Figure 1 shows that in this case periodic oscillations of cell concentrations occur with a period around 25 days.

Assume now that 32% of infected cells survive incubation, which corresponds to  $\beta = 0.9 \times 10^{-6}$ . We obtain the following values:  $R_0 = 4.16$ ,  $a = 0.086$ ,  $\Delta = 0.77$  and  $D = -1.11$ . The corresponding values are  $\mu_1 = 0.11$ ,  $\mu_2 = 0.99$ , which is biologically realistic for  $\mu_2$ . If we take  $\mu = 0.9$ , we obtain Fig. 2, which shows periodic oscillations in cell concentrations. If we increase  $\mu_2$  to the value of 1.1, numerical simulations give the plots in Fig. 3, which shows damped oscillations of cell concentrations followed by convergence to the infected steady state.

Finally, assume that 25% of infected cells survive incubation, corresponding to  $\beta = 0.71 \times 10^{-6}$ . We have in this case  $R_0 = 3.25$ ,  $a = 0.11 > 0$ ,  $\Delta = -0.23 < 0$ . Condition (i) in Theorem 2 is satisfied and the theorem predicts that the infected equilibrium  $E^*$  is locally asymptotically stable. Numerical simulations (see Fig. 4) show that in this case the concentrations of healthy and infected cells converge to the infected equilibrium  $E^*$ .

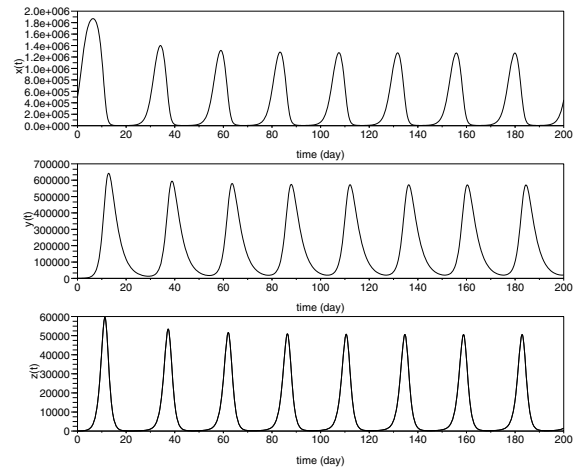


Fig. 1. Half infected cells survive incubation:  $\mu = 1$ .

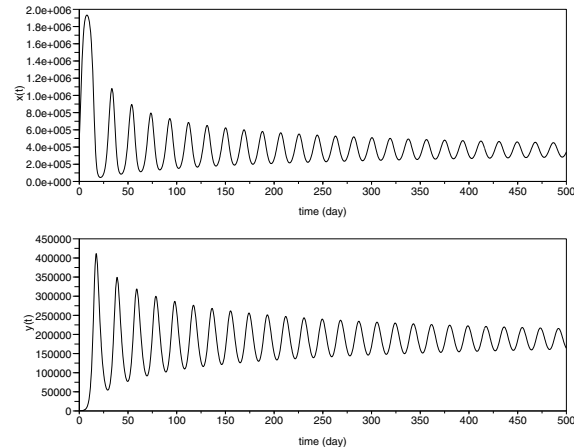


Fig. 2. 32% of infected cells survive incubation:  $\mu = 0.9$ .

## 6. Discussion

It is well known that HIV-1 is thought to be active in areas such as the brain and the lymph nodes, where 98% of lymphocytes reside and in which cell-to-cell spread would be a much more important mode of infection than cell-free viral spread since viral replication in a system with rapid cell turnover kinetics depends on cell-to-cell transfer of virus (Culshaw *et al.*, 2003; Dimitrov *et al.*, 1993; Spouge *et al.*, 1996).

In this paper we modified the infection rate between infected and healthy cells in the model proposed by Cul-

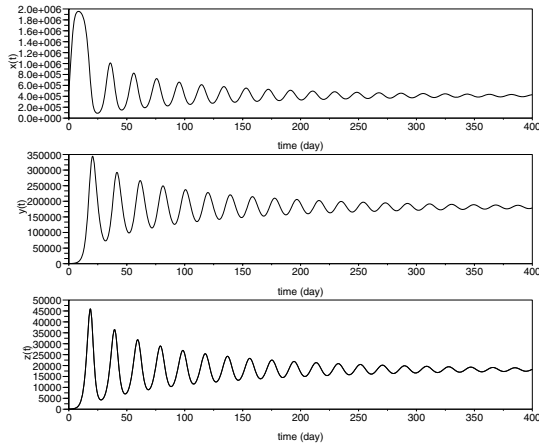


Fig. 3. 32% of infected cells survive incubation:  $\mu = 1.1$ .

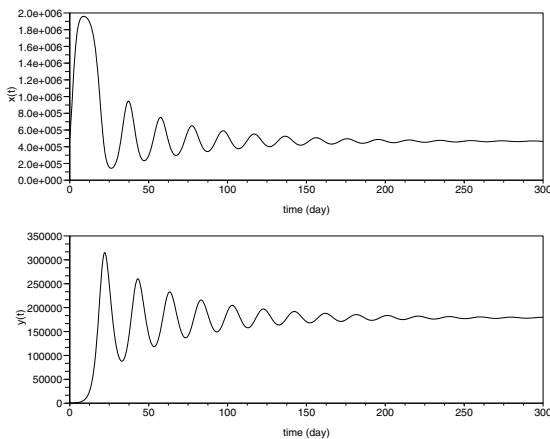


Fig. 4. 25% of infected cells survive incubation.

shaw *et al.* (2003) by assuming that the infection rate term is a saturating function of cell concentration. By using the method of the next generation matrix, the basic reproduction number  $R_0$  of the model is derived. We also proved that the model has at most three equilibria: trivial, healthy and, in case  $R_0 > 1$ , there is a third equilibrium called the infected equilibrium. Stability analysis shows that, when  $R_0 \leq 1$ , infected cells die out and the healthy equilibrium is globally stable. When  $R_0 > 1$ , we proved that the infection is persistent. When the infected equilibrium is unstable, infective oscillations may occur but, unlike the model of Culshaw *et al.* (2003), the system exhibits two Hopf bifurcations  $\mu_1$  and  $\mu_2$  such that the infected equilibrium is locally asymptotically stable either when the average delay is sufficiently large ( $\mu > \mu_2$ ) or small ( $\mu < \mu_1$ ).

Numerical simulations show that, if no more than 7.7% of infected cells survive the incubation period, infected cells die out and the system converges to its healthy equilibrium. If between 7.7% and 30% of infected cells survive the incubation, then the system stabilizes at its in-

fect equilibrium, and when more than 30% of infected cells survive the incubation, then we observe periodic oscillations in cell concentrations. Further, an increase in the fraction of cells surviving the incubation  $\beta/\alpha$ , would increase the periods and amplitudes of cell concentrations, and a reduction in such a fraction leads to damped oscillations and eventually steady states.

Qualitatively, our model differs from that of Culshaw *et al.* (2003) since under realistic parameter regimes it exhibits two Hopf bifurcations and the infected steady state is locally asymptotically stable either when the average delay is sufficiently large or small. For the other values of the delay, the model exhibits stable periodic solutions. The existence of stable periodic solutions is due to the third compartment, which is the compartment of latently infected cells, since the system with only two compartments (healthy and infected cells) does not generate a periodic solution. This result confirms the experimental work of Grossman *et al.* (1998), which showed that infected and latently infected cells may not contribute significantly to the viral load but may be instrumental in sustaining the infection (Spouge *et al.*, 1996).

Quantitatively, we observe in our model some overestimates in the periods of cell concentrations and the critical bifurcation value of cells surviving the incubation period. If 75% of infected cells survive the incubation, our model indicates a period of around 30 days of cell concentrations and a critical bifurcation value around 30, for equivalent values of around 25 days and 22.5%, respectively, in the model by Culshaw *et al.* (2003).

The model proposed in this paper is a first step in studying cell-to-cell HIV-1 spread in non-well-mixed tissue culture. Another compartmental models can be considered to take into account more significantly spatial heterogeneities in tissue culture. We leave this for a future work.

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**Mahieddine Kouche** was born in Annaba, Algeria, in 1970. He obtained his doctorate in applied mathematics from Badji-Mokhtar University in 2007. His research interests include delay differential equations, dynamical systems, and mathematical biology. Currently he is an assistant professor at the Department of Mathematics of Badji-Mokhtar University.

**Bedr'eddine Ainseba** was born in Tlemcen, Algeria, in 1968. He received his Ph.D. from Compiègne University in 1992. Presently he is professor at the University of Bordeaux. His current interests involve control theory, age structured systems, dynamical systems, and mathematical biology.

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