

Dynamics of a Heroin Epidemic Model with Very Population

Xiaoyan Wang¹, Junyuan Yang¹, Xuezhi Li²

¹Department of Applied Mathematics, Yuncheng University, Yuncheng, China

²Department of Mathematics, Xinyang Normal University, Xinyang, China

E-mail: yangjunyuan00@126.com

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Abstract

Based on the model provided by the Mulone and Straughan [1], we relax the population which are constant and obtain the drug-free equilibrium which is global asymptotically stable under some conditions. The system has only uniqueness positive endemic equilibrium which is globally asymptotically stable by using the second compound matrix.

Keywords: Heroin Epidemic Model, Equilibrium, Global Stability

1. Introduction

Heroin is an opiate drug that is synthesized from morphine, a naturally occurring substance extracted from the seed pod of the Asian opium poppy plant. Heroin usually appears as a white or brown powder or as a black sticky substance, known as “black tar heroin” [2]. Heroin users are at high risk for addiction—it is estimated that about 23 percent of individuals who use heroin become dependent on it. The spread of heroin habituation and addiction presents many of the well-known phenomena of epidemics, including rapid diffusion and clear geographic boundaries. It is unrealistic to repeat the experiment on the human body for obtaining the statistic data. Mathematical models play very important role in dealing with these problems. Mathematical modellings are very useful tools to predict how classes of drug takers behave, and provide a good suggestion for the treatment strategies. It is interest to explore the problem mathematically, reducing the factors to the essential transmission mechanism, in real life, is much more complex because of imponderable and often intervening biological, psychological, and social conditions.

During recent years, many mathematical models have been developed to describe the Heroin epidemic model (see [1,3]). One of the recent model divided the mathematical problem into tree class, namely susceptibles, heroin users or alcoholics, and heroin users or alcoholics undergoing treatment which denote by $S(t)$, $U_1(t)$ and $U_2(t)$, respectively. The model is

$$\begin{aligned} \frac{dS}{dt} &= \Lambda - \frac{\beta_1 S U_1}{N} - \mu S, \\ \frac{dU_1}{dt} &= \frac{\beta_1 S U_1}{N} + \frac{\beta_3 U_1 U_2}{N} - (p + \mu + \delta_1) U_1, \\ \frac{dU_2}{dt} &= p U_1 - \frac{\beta_3 U_1 U_2}{N} - (\mu + \delta_2) U_2 \end{aligned} \quad (1.1)$$

where the number of individuals enter the susceptible population at a rate, Λ , die at rate μ . Infection of a drug user occurs through a simple mass action process $\beta_1 S U_1 / N$, where β_1 is probability of becoming a drug user, per unit time. The probability of a drug user in treatment relapsing to untreated use, per unit time is p . Infected heroin users die at enhanced rate δ_1 , treatment infected heroin users die at enhanced rate δ_2 . In this model, the authors considered the global stability of the drug equilibrium under some conditions. However, they treated the total population is a constant for simple. In fact, we sum the tree equations of (1.1) and get

$$\frac{dN}{dt} = \Lambda - \mu N - \delta_1 U_1 - \delta_2 U_2.$$

It is easy to see that the total population is not a constant. This assumption is not reasonable and maybe simply the problem. Since (1.1) can be changed a plane system under the constant population. In our paper we will relax this assumption and assume the total population is very according to the time. In our model we use the bilinear law incidence function instead of standard incidence. The model is changed into

$$\begin{aligned} \frac{dS}{dt} &= \Lambda - \beta_1 S U_1 - \mu S, \\ \frac{dU_1}{dt} &= \beta_1 S U_1 + \beta_3 U_1 U_2 - (p + \mu + \delta_1) U_1, \\ \frac{dU_2}{dt} &= p U_1 - \beta_3 U_1 U_2 - (\mu + \delta_2) U_2 \end{aligned} \quad (1.2)$$

2. Basic Reproduction Number and Stability of Drug-Free Equilibrium

The drug-free equilibrium is given by

$$E_0 = (S^0, 0, 0) = \left(\frac{\Lambda}{\mu}, 0, 0 \right).$$

The basic reproduction number R_0 is defined to be the expected number of secondary cases produced, in a completely susceptible population, by a typical infected individual during its entire period of infection. We follow the recipe of [4] to calculate the basic reproduction number. O. Diekmann *et al.* in [5] has shown R_0 as the spectral radius of the next generation matrix. Firstly we need to separate the new infections from other factors. We define $F(X)$ to be the vector which represents the rate of new infections that appear in the population where X is a vector given by $X = (U_1, U_2, S)$. In this model, we get $F = (\beta_1 S U_1, 0, 0)$,

$$\begin{aligned} V = & (-\beta_3 U_1 U_2 + (p + \mu + \delta_1) U_1, \beta_3 U_1 U_2 \\ & + (\mu + \delta_2) U_2 - p U_1, \beta_1 U_1 S + \mu S - \Lambda) \end{aligned}$$

F, V have the same definition as explained in [4]. According to the recipe presented in [4] we define the derivatives $DF(E_0)$ and $DV(E_0)$ in the following way

$$F = \begin{pmatrix} \frac{\beta_1 \Lambda}{\mu} & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix}$$

and

$$V = \begin{pmatrix} p + \mu + \delta_1 & 0 & 0 \\ -p & \mu + \delta_2 & 0 \\ \frac{\beta_1 \Lambda}{\mu} & 0 & \mu \end{pmatrix}.$$

Finally the basic reproduction number is obtained by calculating the spectral radius of FV^{-1} . Then the basic reproduction number for the model presented in system (1.2) is given by the following expression

$$R_0 = \rho(FV^{-1}) = \frac{\beta_1 \Lambda}{\mu(p + \mu + \delta_1)}.$$

This completes the calculation of basic reproduction number. The Jacobian matrix of (1.2) at drug-free equilibrium E_0 is

$$J_{E_0} = \begin{pmatrix} -\mu & -\frac{\beta_1 \Lambda}{\mu} & 0 \\ 0 & \frac{\beta_1 \Lambda}{\mu} - (p + \mu + \delta_1) & 0 \\ 0 & p & -(\mu + \delta_2) \end{pmatrix}$$

The characteristic roots of the matrix are

$$\lambda_1 = -\mu < 0, \lambda_2 = -(\mu + \delta_2) < 0, \lambda_3 = \frac{\beta_1 \Lambda}{\mu} - (p + \mu + \delta_1).$$

Hence the characteristic equation just has negative real part roots under some condition. Then we can obtain the following theorem.

Theorem 2.1. If $R_0 < 1$, the drug-free equilibrium is locally asymptotically stable otherwise it is unstable.

Theorem 2.2. If $\frac{\beta_1 \Lambda}{\mu(\mu + \delta_1)} < 1$ drug-free equilibrium

is globally stable.

Proof. We set a Lyapunov function

$$V(t, U_1, U_2) = U_1 + U_2.$$

Then

$$\begin{aligned} \left. \frac{dV}{dt} \right|_{(2.1)} &= U_1 + U_2 \\ &= \beta_1 S U_1 - (\mu + \delta_1) U_1 - (\mu + \delta_2) U_2 \\ &\leq (\mu + \delta_1) \left(\frac{\beta_1 \Lambda}{\mu(\mu + \delta_1)} - 1 \right) U_1 - (\mu + \delta_2) U_2 \end{aligned}$$

When $\frac{\beta_1 \Lambda}{\mu(\mu + \delta_1)} < 1$, it is easy to get $\frac{dV}{dt} \leq 0$. Also

$\dot{V} = 0$ if and only if U_1, U_2 are all zero. Note that the

largest invariant set is $M = \left\{ \left(\frac{\Lambda}{\mu}, 0, 0 \right) \mid \frac{dV}{dt} = 0 \right\}$, the

drug-free equilibrium E_0 is globally asymptotically stable by Lassale's invariance principle.

3. Existence and Stability of the Endemic Equilibrium

In this section we discuss the existence of the endemic equilibrium. We just consider the special situation. We find the positive endemic equilibrium for (1.2) which is obtained by solving the following sets of equations

$$\begin{cases} \Lambda - \beta_1 S^* U_1^* - \mu S^* = 0, \\ \beta_1 S^* U_1^* + \beta_3 U_2^* U_3^* - (p + \mu + \delta_1) U_1^* = 0, \\ p U_1^* - \beta_3 U_1^* U_2^* - (\mu + \delta_2) U_2^* = 0. \end{cases} \quad (3.1)$$

Solving the first and second equations of (3.1), we obtain

$$U_1^* = \frac{\Lambda - \mu S^*}{\beta_1 S^*}, U_2^* = \frac{p + \mu + \delta_1}{\beta_3}.$$

Substituting U_1^* and U_2^* into the third equation of (3.1), we obtain S^* should satisfies the following equation

$$P(E^*) = Ax^2 + Bx + C = 0.$$

where

$$\begin{aligned} A &= \beta_1^2 (\mu + \delta_2) - \beta_1 \beta_3 \mu, \\ B &= \beta_1 \beta_3 \Lambda + \beta_3 \mu (p + \mu + \delta_1) - \beta_1 (\mu + \delta_2) (\mu + \delta_1), \\ C &= -\beta_3 \Lambda (\mu + \delta_1). \end{aligned} \quad (3.2)$$

In our model, we just consider $\beta_1 > \beta_3$. It is easy to see $A > 0$ and $C < 0$. Due to Descartes' rule of sings, the number of real positive roots of $P(E^*)$ is determined by the following table

B	Number of Real Positive Roots
>0	1
<0	1

$$J_{E^*} = \begin{pmatrix} -\beta_1 U_1^* - \mu & -\beta_1 S^* & 0 \\ \beta_1 U_1^* & \beta_1 S^* + \beta_3 U_2^* - (p + \mu + \delta_1) & \beta_3 U_1^* \\ 0 & p - \beta_3 U_2^* & -\beta_3 U_1^* - (\mu + \delta_2) \end{pmatrix}.$$

Then the characteristic of the endemic equilibrium is

$$\lambda^3 + a_1 \lambda^2 + a_2 \lambda + a_3 = 0, \quad (3.3)$$

where

$$\begin{aligned} a_1 &= \beta_3 U_1^* + 2\mu + \delta_1 + \beta_1 U_1^* > 0 \\ a_2 &= (\beta_1 U_1^* + \mu)(\beta_3 U_1^* + \mu + \delta_2) + \beta_3 U_1^* (\mu + \delta_1) + \beta_1 (\beta_1 - \beta_3) S^* U_1^* > 0, \\ a_3 &= \beta_3 U_1^* (\beta_1 U_1^* + \mu)(\mu + \delta_1) + \beta_1 (\beta_1 - \beta_3) \mu S^* U_1^* + \beta_1^2 S^* U_1^* (\mu + \delta_2) > 0. \end{aligned}$$

Define $H(E^*) = a_1 a_2 - a_3$, and straightly calculate and then we obtain

$$H(E^*) = (\beta_3 U_1^* + \mu + \delta_2) \left[(\beta_1 U_1^* + \mu)^2 + (\beta_3 U_1^* + \mu + \delta_2) (\beta_1 U_1^* + \mu) - (\mu + \delta_2) \beta_3 U_2^* \right] + \beta_1^2 S^* U_1^* (\beta_1 U_1^* + \mu).$$

Define $y = (\beta_1 U_1^* + \mu)$, and $f(y) = y^2 + (\beta_3 U_1^* + \mu + \delta_2) - (\mu + \delta_2) \beta_3 U_2^*$. Note that $\Delta = (\beta_3 U_1^* + \mu + \delta_2)^2 + 4(\mu + \delta_2) \beta_3 U_2^* > 0$, it is easy to get $H(E^*) > 0$. Due to the Routh Hurtwiz [6], we get the following theorem.

From the expression of U_1^* and U_2^* , and positivity of the endemic equilibrium, we note that

$$S^* < \min \left\{ \frac{\Lambda}{\mu}, \frac{p + \mu + \delta_1}{\beta_1} \right\}$$

If $R_0 > 1$, $S^* < \frac{p + \mu + \delta_1}{\beta_1}$. We assume the two real positive roots of (1.2) are respectively x_- and x_+ , we check the positive real roots of (1.2)

x	Number of Real Positive Roots of (3.1)
$x_+ \leq \frac{p + \mu + \delta_1}{\beta_1}$	1
$x_+ > \frac{p + \mu + \delta_1}{\beta_1}$	0

Concluding the above discussion, then we can obtain the following theorem.

Theorem 3.1. 1) If $R_0 > 1$, and $x_+ \leq \frac{p + \mu + \delta_1}{\beta_1}$, (1.2)

has only unique real positive endemic equilibrium.

Next, we will discuss the locally stability of the endemic equilibrium. The Jacobian matrix at endemic equilibrium is

Theorem 3.2. Let E^* be the endemic equilibria of system (1.2) as defined before, and assume

$x_+ \leq \frac{p + \mu + \delta_1}{\beta_1}$ and $R_0 > 1$, then it is locally asymptotically stable.

Finally, we will consider the global stability of the endemic equilibrium is asymptotically orbitally stable with asymptotic phase.

Theorem 3.3. The endemic equilibrium of E^* to

$$\begin{pmatrix} \begin{pmatrix} -\beta_1 U_1 + \beta_1 S + \beta_3 U_2 \\ -(p + 2\mu + \delta_1) \end{pmatrix} & \beta_3 U_1 & 0 \\ p - \beta_3 U_2 & -\beta_1 U_1 - \beta_3 U_1 - (2\mu + \delta_2) & -\beta_1 S \\ 0 & \beta_1 U_1 & \begin{pmatrix} \beta_1 S + \beta_3 U_2 - \beta_3 U_1 \\ -(p + 2\mu + \delta_1 + \delta_2) \end{pmatrix} \end{pmatrix},$$

we can write the linear system (1.2) with respect to a solution $(S(t), U_1(t), U_2(t))$ of (1.2) as the following 3×3 system (3.4):

$$\begin{aligned} X' &= -(\beta_1 U_1 + p + 2\mu + \delta_1 - \beta_1 S - \beta_3 U_2)X + \beta_3 U_1 Y, \\ Y' &= (p - \beta_3 U_2)X - [(\beta_1 + \beta_3)U_1 + 2\mu + \delta_2]Y - \beta_1 S Z, \\ Z &= \beta_1 U_1 Y - [p + 2\mu + \delta_1 + \delta_2 - \beta_1 S - \beta_3 U_2 + \beta_3 U_1]Z. \end{aligned}$$

To show the asymptotic stability of the system (3.4) we consider the following function:

$$V(X, Y, Z; S, U_1, U_2) = |P(S, U_1, U_2) \cdot (X, Y, Z)^*|$$

where the matrix $P = \text{diag}(I, U_1/U_2, U_1/U_2)$ and $|\cdot|$ is the norm in R^3 defined by

$$D_+ |X(t)| \leq -(\beta_1 U_1 + p + 2\mu + \delta_1 - \beta_1 S - \beta_3 U_2)|X| + \beta_3 U_1 \left(\frac{U_1}{U_2} (|Y| + |Z|) \right),$$

and

$$\begin{aligned} D_+ |Y| &\leq (p - \beta_3 U_2)|X| - [(\beta_1 + \beta_3)U_1 + 2\mu + \delta_2]|Y| - \beta_1 S |Z|, \\ D_+ |Z| &\leq \beta_1 U_1 |Y| - [p + 2\mu + \delta_1 + \delta_2 - \beta_1 S - \beta_3 U_2 + \beta_3 U_1]|Z|. \end{aligned}$$

and thus

$$\begin{aligned} D_+ \frac{U_1}{U_2} (|Y| + |Z|) &= \left(\frac{U'_1}{U_1} - \frac{U'_2}{U_2} \right) \frac{U_1}{U_2} (|Y| + |Z|) + \frac{U_1}{U_2} D_+ (|Y| + |Z|), \\ &\leq \frac{U_1}{U_2} (p - \beta_3 U_2)|X| + \frac{U_1}{U_2} (|Y| + |Z|) \left(\frac{U'_1}{U_1} - \frac{U'_2}{U_2} - \beta_3 U_1 - 2\mu - \delta_2 + \beta_3 U_2 \right) (|Y| + |Z|). \end{aligned}$$

We claim that

$$D_+ V(t) \leq \sup \{g_1(t), g_2(t)\} V(t)$$

where

$$\begin{aligned} g_1(t) &= -\beta_1 U_1 - p - 2\mu - \delta_1 + \beta_1 S, \\ g_2(t) &= \frac{U'_1}{U_1} - \frac{U'_2}{U_2} - \beta_3 U_1 - 2\mu - \delta_2 + \beta_3 U_2 \end{aligned}$$

Using (1.2), we find

(1.2), if it exists, is asymptotically orbitally stable with asymptotic phase.

Proof. The second compound matrices $J(S, U_1, U_2)$ of (1.2) is given by

$$|(X, Y, Z)| = \sup \{|X|, |Y| + |Z|\}. \tag{3.5}$$

Suppose that the solution $(S(t), U_1(t), U_2(t))$ is periodic of least period $w > 0$. The matrix P and its inverse are thus well defined and smooth along γ . There exists constant $c > 0$ such that

$$V(X, Y, Z; S, U_1, U_2) \geq c |(X, Y, Z)|. \tag{3.6}$$

for all $(X, Y, Z) \in R^3$ and $(S, E, I) \in \gamma$. Let $(X(t), Y(t), Z(t))$ be a solution to (3.4) and

$$V(t) = V(X, Y, Z; S, U_1, U_2) = \sup \left\{ |X|, \frac{U_1}{U_2} (|Y| + |Z|) \right\}.$$

The right-hand derivative of $V(t)$ exists and its calculation is described in [7]. In fact, direct calculation yields

$$g_1(t) = \frac{U'_1}{U_1} - \beta_1 U_1 - \beta_3 U_2 - \mu \leq \frac{U'_1}{U_1} - \mu,$$

and

$$g_2(t) = -\frac{p U_1}{U_2} < 0$$

and thus

$$\int_0^w \sup \{g_1(t), g_2(t)\} dt \leq \log U_1(t) \Big|_0^w - \mu w < 0$$

which, together with (3.9), implies that $V(t) \rightarrow 0$ as $t \rightarrow \infty$ and in turn that $(X(t), Y(t), Z(t)) \rightarrow 0$ as $t \rightarrow \infty$ by (3.6). As a result, the linear system (1.2) is asymptotically stable and the periodic solution $(S(t), U_1(t), U_2(t))$ is asymptotically orbitally stable with asymptotic phase by Theorem 3.1 [7].

4. Conclusions and Simulation

In this paper we have addressed the problem of investigation the existence and stability of the equilibria heroin epidemic model provided by Mulone and Straughan. We relax the condition the total population is a constant. Under some condition, the drug-free equilibrium is global asymptotically stable; the number of equilibria are determined by Theorem 3.1 and we show that the endemic equilibrium is also globally asymptotically stable under any conditions. The natures of the model are not different from the epidemic models [1,3]. We use numerical simulations to illustrate the theoretical results obtained in previous sections. As an example, we take the parameter values as follows: $\Lambda = 1, \beta_1 = 0.008, \beta_3 = 0.001, p = 0.04, \mu = 0.01, \delta_1 = 0.05, \delta_2 = 0.06$. By using the classical implicit format solving the ordinary differential equations and the method of steps for differential equations, we can solve the numerical solutions of (1.2) via the software package Matlab (see **Figure 1**) is globally asymptotically stable with asymptotic phase. The initial conditions are $S_0 = 10, U_{10} = 1, U_{20} = 1$.

In future, there are some problems that will be solved. We continue the problems of system (2.1) on substituting the bilinear into standard incidence. Whether there are some kinds of bifurcations for the model or not is still open. If we assume the input is impulsive input or sto-

chastic perturbations, what results will occur. We give some numerical results. For the impulsive perturbation, we give the following system (4.1).

$$\begin{cases} \frac{dS}{dt} = -\beta_1 S U_1 - \mu S, & t \neq nT, \\ S(t^+) = S(t) + \Lambda, & t = nT, \\ \frac{dU_1}{dt} = \beta_1 S U_1 + \beta_3 U_1 U_2 - (p + \mu + \delta_1) U_1, & t \neq nT, \\ U_1(t^+) = U_1(t), & t = nT, \\ \frac{dU_2}{dt} = p U_1 - \beta_3 U_1 U_2 - (\mu + \delta_2) U_2, & t \neq nT, \\ U_2(t^+) = U_2(t), & t = nT. \end{cases}$$

Theorem 4.1. The drug-free periodic solution of (4.1) is globally asymptotically stable, and the system is persistent.

As another example, we take the parameter values as follows: $\Lambda = 2, \beta_1 = 1, \beta_3 = 0.1, p = 0.4, \mu = 0.01, \delta_1 = 0.05, \delta_2 = 0.06, T = 1$. We can solve the numerical solutions of (4.1) via the software package Matlab (see **Figure 2**).

For the stochastic perturbation, we give the following system (4.2).

$$\begin{cases} \frac{dS}{dt} = \Lambda - \beta_1 S U_1 - \mu S + \sigma d\xi_t^1, \\ \frac{dU_1}{dt} = \beta_1 S U_1 + \beta_3 U_1 U_2 - (p + \mu + \delta_1) U_1 + \sigma d\xi_t^2, \\ \frac{dU_2}{dt} = p U_1 - \beta_3 U_1 U_2 - (\mu + \delta_2) U_2 + \sigma d\xi_t^3. \end{cases}$$

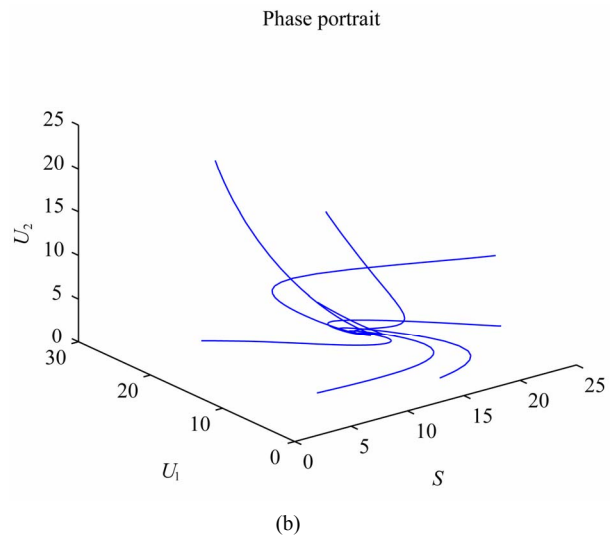
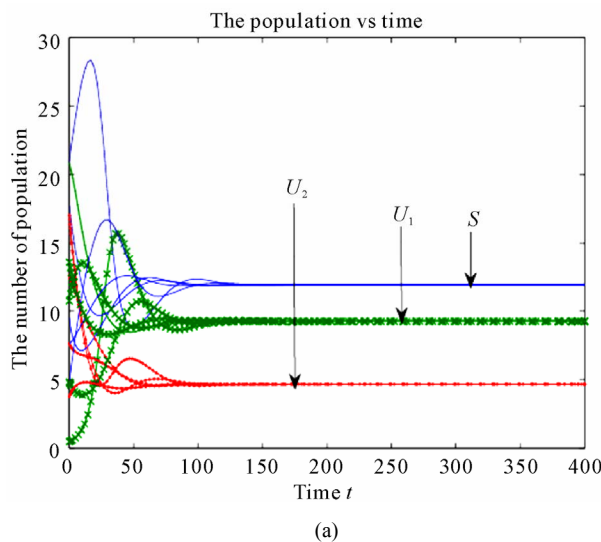
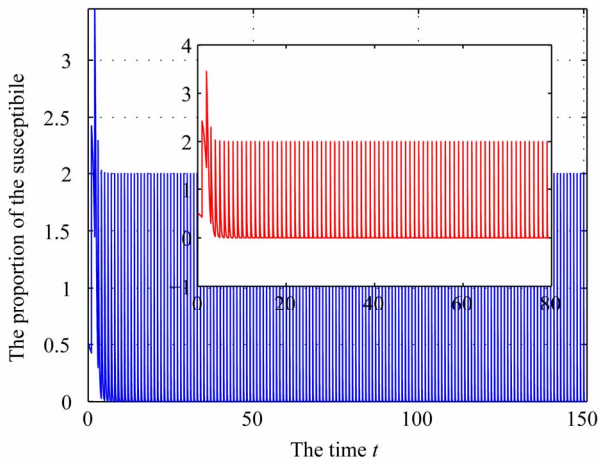
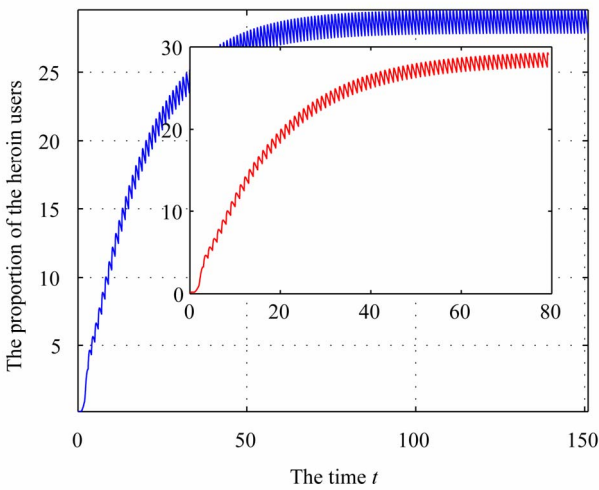


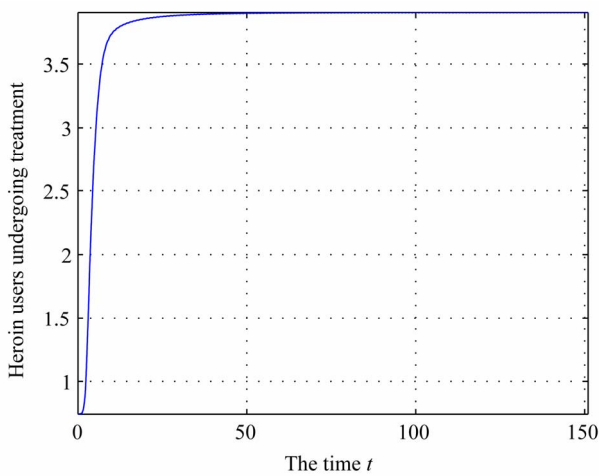
Figure 1. (a)-(b) show that endemic equilibrium to (1.2).



(a)

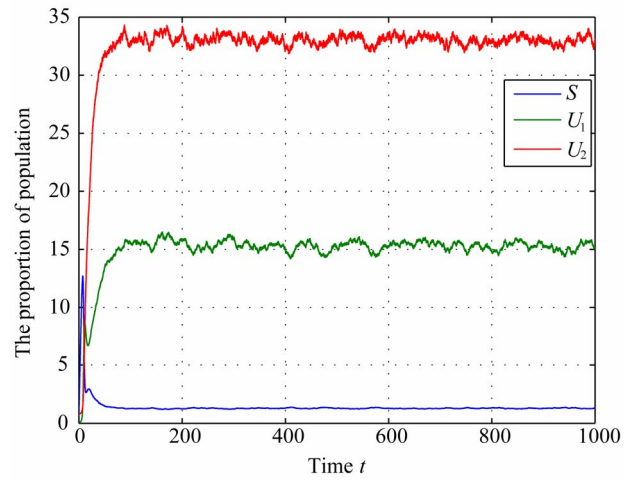


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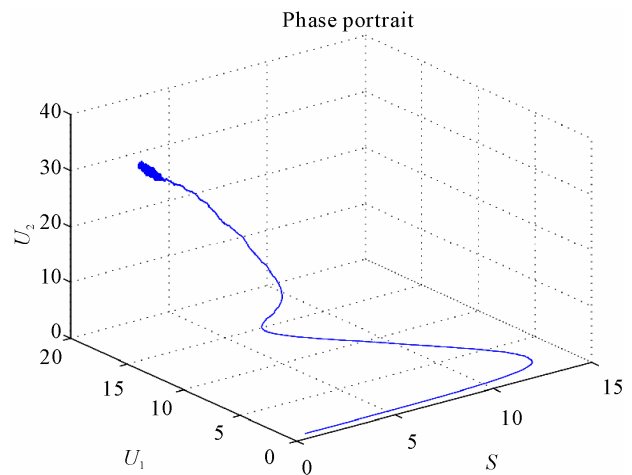


(c)

Figure 2. (a)-(c) show that the system (4.1) is persistent. The initial conditions are $S_0 = 0.5126$, $U_{10} = 0.1688$, $U_{20} = 0.7448$.



(a)



(b)

Figure 3. (a)-(b) show that the zero solution of system (4.2) is probably asymptotically stable. The initial conditions are $S_0 = 0.5126$, $U_{10} = 0.1688$, $U_{20} = 0.7448$.

We take the parameter values as follows: $\Lambda = 2$,

$$\beta_1 = 0.1, \beta_3 = 0.01, p = 0.4, \mu = 0.01,$$

$$\delta_1 = 0.05, \delta_2 = 0.06, \sigma = 0.002.$$

we can solve the numerical solutions of (4.1) via the software package Matlab (see **Figure 3**).

Furthermore, we will give the detail analysis proofs for the theorems in the future.

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6. References

- [1] G. Mulone and B. Straughan, "A Note on Heroin Epidemics," *Mathematical Biosciences*, Vol. 218, No. 2, 2009, pp. 138-141. [doi:10.1016/j.mbs.2009.01.006](https://doi.org/10.1016/j.mbs.2009.01.006)
- [2] NIDA InfoFacts: Heroin.
<http://www.nida.nih.gov/infofacts/heroin.html>
- [3] E. White and C. Comiskey, "Heroin Epidemics, Treatment and ODE Modelling," *Mathematical Biosciences*, Vol. 208, No. 1, 2007, pp. 312-324. [doi:10.1016/j.mbs.2006.10.008](https://doi.org/10.1016/j.mbs.2006.10.008)
- [4] P. Driessche and W. James, "Reproduction Numbers and Subthreshold Endemic Equilibria for Compartmental Models of Disease Transmission," *Mathematical Biosciences*, Vol. 180, No. 1-2, 2002, pp. 29-48. [doi:10.1016/S0025-5564\(02\)00108-6](https://doi.org/10.1016/S0025-5564(02)00108-6)
- [5] O. Diekmann, J. A. P. Heesterbeek and J. A. J. Metz, "On the Definition and the Computation of the Basic Reproduction Ratio R_0 in Models for Infectious Diseases in Heterogenous Populations," *Journal of Mathematical Biology*, Vol. 28, No. 4, 1990, pp. 365-382. [doi:10.1007/BF00178324](https://doi.org/10.1007/BF00178324)
- [6] M. Y. Li and J. S. Muldowney, "Global Stability for the SEIR Model in Epidemiology," *Mathematical Biosciences*, Vol. 125, No. 2, 1995, pp. 155-164. [doi:10.1016/0025-5564\(95\)92756-5](https://doi.org/10.1016/0025-5564(95)92756-5)
- [7] J. K. Hale, "Ordinary Differential Equations," John Wiley and Sons, New York, 1969.