

# Stochastic Stability of the non linear epidemic model with temporary immunity

Laid Chahrazed.

Department of Mathematics, Faculty of Exacte Sciences,  
University Constantine 1, Algeria.

**Abstract:** In this paper, addresses a time-delayed epidemiologic model by experiencing the disease; whenever the quarantine will return to the susceptible. First, the equilibrium and global stabilities of the endemic equilibrium. Second, Stochastic Stability. Finally, the equilibrium and stability of the epidemic model with age.

**Keywords:** Epidemic model, Global asymptotic stability, Lyaponov functional, Stochastic Stability.

## 1 Introduction

This paper considers the following epidemic model with temporary immunity:

$$\begin{cases} \dot{S}(t) = \rho - (\mu_1 + d)S(t) - \beta S(t)Q(t), \\ \dot{I}(t) = \beta S(t)Q(t) - (\mu_2 + d)I(t) - \gamma e^{-\mu_2\tau}S(t-\tau)Q(t-\tau), \\ \dot{Q}(t) = \gamma e^{-\mu_2\tau}S(t-\tau)Q(t-\tau) - (\mu_3 + d)Q(t), \end{cases} \quad (1)$$

Consider a population of size  $N(t)$  at time  $t$ , this population is divided into three subclasses, with  $N(t) = S(t) + I(t) + Q(t)$ ; where  $S(t)$ ,  $I(t)$ , and  $Q(t)$  denote the sizes of the population susceptible to disease, and infectious members, quarantine members with the possibility of infection through temporary immunity, respectively. The positive constants  $\mu_1$ ,  $\mu_2$ , and  $\mu_3$  represent the death rates of susceptible, infectious and quarantine. Biologically, it is natural to assume that  $\mu_1 \leq \min \{\mu_2, \mu_3\}$ . The positive constant  $d$  is natural mortality rate. The positive constant  $\gamma$  represent the recovery rate of infection. The positive constant  $\beta$  is the average numbers of contacts infective for  $S$  and  $I$ .  $\rho$  the positive constant is the parameter of immigration. The term  $\gamma e^{-\mu_2\tau}S(t-\tau)Q(t-\tau)$  reflects the fact that an individual has recovered from infection and still are alive after infectious period  $\tau$ , where  $\tau$  is the length of immunity period.

The initial condition of (1) is given as.

$$S(\eta) = \Phi_1(\eta), \quad I(\eta) = \Phi_2(\eta), \quad Q(\eta) = \Phi_3(\eta), \quad -\tau \leq \eta \leq 0, \quad (2)$$

Where  $\Phi = (\Phi_1, \Phi_2, \Phi_3)^T \in \mathbb{C}$  such that  $S(\eta) = \Phi_1(\eta) = \Phi_1(0) \geq 0$ ,  $I(\eta) = \Phi_2(\eta) = \Phi_2(0) \geq 0$ ,  $Q(\eta) = \Phi_3(\eta) = \Phi_3(0) \geq 0$ .

Let  $C$  denote the Banach space  $C([-\tau, 0], \mathbb{R}^3)$  of continuous functions mapping the interval  $[-\tau, 0]$  into  $\mathbb{R}^3$ . With a biological meaning, we further assume that  $\Phi_i(\eta) = \Phi_i(0) \geq 0$  for  $i = 1, 2, 3$ .

Consider the system without the parameter of emigrations. Hence system (1) can be rewritten as

$$\begin{cases} \dot{S}(t) = \rho - (\mu_1 + d)S(t) - \beta S(t)Q(t), \\ \dot{I}(t) = \beta S(t)Q(t) - (\mu_2 + d)I(t) - \gamma e^{-\mu_2\tau}S(t-\tau)Q(t-\tau), \\ \dot{Q}(t) = \gamma e^{-\mu_2\tau}S(t-\tau)Q(t-\tau) - (\mu_3 + d)Q(t), \end{cases} \quad (3)$$

With the initial conditions.

$$S(\eta) = \Phi_1(\eta), \quad I(\eta) = \Phi_2(\eta), \quad Q(\eta) = \Phi_3(\eta), \quad -\tau \leq \eta \leq 0, \quad (4)$$

Where  $\Phi_1(0) \geq 0, \Phi_2(0) \geq 0, \Phi_3(0) \geq 0, \quad -\tau \leq \eta < 0$ .

Since  $\dot{N}(t) \leq \rho - (\mu_1 + d)N(t)$ , and  $S(t) + I(t) + Q(t) \leq N(t)$ .

The region  $\Omega = \{(S, I, Q) \in \mathbb{R}_+^3, S + I + Q \leq N < \frac{\rho}{\mu_1 + d}\}$  is positively invariant set of (3).

This paper deals with the equilibrium and stability of system (3), precisely the global stability of endemic equilibrium by using Lyapunov functional technique, under certain conditions on the parameter this means that the disease persist in population.

Next, we introduce a Brownian motion to system (3) and we transform it into an Itô stochastic differential equation by using the contact rate a white noise. Finally study equilibrium of epidemic model with age.

The organization of this paper is as follows, in Section 1, Equilibrium and stability

of the model. In Section 2, Global asymptotic stability of endemic equilibrium. In

Section 3, stochastic stability. In Section 4 equilibrium and stability of the epidemic model with age.

## 2 Equilibrium and stability

An equilibrium point of system (3) satisfies

$$\begin{cases} \rho - (\mu_1 + d)S - \beta SQ = 0, \\ \beta kSQ - (\mu_2 + d)I - \gamma e^{-\mu_2\tau}S(t-\tau)Q(t-\tau) = 0, \\ \gamma e^{-\mu_2\tau}S(t-\tau)Q(t-\tau) - (\mu_3 + d)Q = 0, \end{cases} \quad (5)$$

We calculate the points of equilibrium in the absence and presence of infection.

In the absence of infection  $I = 0$ , the system (5) has a disease-free equilibrium  $E_0$ :

$$E_0 = \left( \hat{S}, \hat{I}, \hat{Q} \right)^T = \left( \frac{\rho}{\mu_1 + d}, 0, 0 \right)^T.$$

The eigenvalues can be determined by solving the characteristic equation of the linearization of (3) near  $E_0$  is

$$\det \begin{pmatrix} -(\mu_1 + d) - A & 0 & -\frac{\beta \rho}{\mu_1 + d} \\ 0 & -(\mu_2 + d) - A & \frac{\rho(\beta - \gamma e^{-\mu_2 \tau})}{\mu_1 + d} \\ 0 & 0 & \frac{\rho \gamma e^{-\mu_2 \tau}}{\mu_1 + d} - (\mu_3 + d) - A \end{pmatrix} = 0 \quad (6)$$

So the eigenvalues are

$$A_1 = -(\mu_1 + d), \quad A_2 = -(\mu_2 + d).$$

In order for  $\lambda_1, \lambda_2$ , to be negative, it is required that.

$$\frac{\rho \gamma e^{-\mu_2 \tau}}{\mu_1 + d} < \mu_3 + d \quad (7)$$

Then we define the basic reproduction number of the infection  $R_0$  as follows.

$$R_0 = \frac{\rho \gamma e^{-\mu_2 \tau}}{(\mu_1 + d)(\mu_3 + d)} \quad (8)$$

In the presence of infection  $I \neq 0$ , substituting in the system,  $\Omega$  also contains a unique positive, endemic equilibrium  $E_\tau^* = (S_\tau^*, I_\tau^*, Q_\tau^*)^T$  where

$$\begin{cases} S_\tau^* = \frac{\rho}{\mu_1 + d} \times \frac{1}{R_0}, \\ I_\tau^* = \frac{R_0 - 1}{\mu_2 + d} \left[ \frac{\rho}{R_0} - \frac{(\mu_1 + d)(\mu_3 + d)}{\beta} \right], \\ Q_\tau^* = \frac{\mu_1 + d}{\beta} (R_0 - 1) \end{cases} \quad (9)$$

Note that

$$N_\tau^* = \frac{\rho(\mu_2 - \mu_1)}{R_0(\mu_1 + d)(\mu_2 + d)} + \frac{\rho}{\mu_2 + d} + \frac{(R_0 - 1)(\mu_1 + d)(\mu_2 - \mu_3)}{\beta(\mu_2 + d)} \quad (10)$$

So  $E_\tau^* = (S_\tau^*, I_\tau^*, Q_\tau^*)^T$  is the unique positive endemic equilibrium point which exists if  $R_0 > 1$ .

**Theorem 1** *The disease-free equilibrium  $E_0$  is locally asymptotically stable if  $R_0 < 1$  and unstable if  $R_0 > 1$ .*

**Theorem 2** *With  $R_0 > 1$ , system (3) has a unique non-trivial equilibrium  $E_\tau^*$  is locally asymptotically stable.*

### 3 Global asymptotic stability of endemic equilibrium

Consider system (3), with introducing the variables,

$$x(t) = S(t) - S_\tau^*, \quad y(t) = I(t) - I_\tau^*, \quad z(t) = Q(t) - Q_\tau^*,$$

System (3) is centered at the endemic equilibrium  $E_\tau^* = (S_\tau^*, I_\tau^*, Q_\tau^*)^T$ , then

$$\begin{cases} \dot{x}(t) = [-(\mu_1 + d) - \beta Q_\tau^*] x + [-\beta S_\tau^*] z, \\ \dot{y}(t) = [(\beta - \gamma e^{-\mu_2 \tau}) Q_\tau^*] x + [-(\mu_2 + d)] y + [(\beta - \gamma e^{-\mu_2 \tau}) S_\tau^*] z, \\ \dot{z}(t) = [\beta \gamma e^{-\mu_2 \tau} Q_\tau^*] x + [\gamma e^{-\mu_2 \tau} S_\tau^* - (\mu_3 + d)] z \end{cases} \quad (11)$$

**Lemma 3** *Let*

$$S(s) = S(0) > 0, \quad I(s) = I(0) \geq 0 \text{ for all } s \in [-\tau, 0] \text{ and } Q(0) > 0.$$

$S(t)$ ,  $I(t)$  and  $Q(t)$  solutions of system (3) are positive for all  $t > 0$ .

**Proof.** For contradiction there exists the first time  $t_0$ , such that  $S(t_0)Q(t_0) = 0$ .

- Assume that  $S(t_0) = 0$ , then  $Q(t) \geq 0$  for all  $t \in [0, t_0]$ . With Eq 1 in the system (1) we have

$$\dot{S}(t_0) = \rho > 0.$$

For  $S(t_0) = 0$ ,  $S_0 > 0$ , we must have  $\dot{S}(t_0) < 0$  which is contradiction.

- Assume that  $I(t_0) = 0$ , then with Eq 2 in the system (1) we have

$$\dot{I}(t_0) = -\gamma e^{-\mu_2 \tau} S(t - \tau) Q(t - \tau)$$

$\dot{I}(t_0)$  is positive because  $S(t)$  and  $Q(t)$  solutions of system (1) are positive for all  $t > 0$ .

- For  $I(t_0) = 0$ ,  $I > 0$ , we must have  $\dot{I}(t_0) < 0$  which is contradiction.

- Assume that  $Q(t_0) = 0$ , then  $S(t) \geq 0$  for all  $t \in [0, t_0]$ .

Eq 3 in the system (3) we have

$$\begin{aligned} \dot{Q}(t_0) &= \gamma e^{-\mu_2 \tau} S(t - \tau) Q(t - \tau), \\ Q(t_0) &= \gamma \int_{t_0 - \tau}^{t_0} e^{-\mu_2(t_0 - s)} S(s) Q(s) ds. \end{aligned}$$

$S(s) > 0, S(s) > 0$  for all  $t \in [0, t_0]$ .

have  $\gamma \int_{t_0 - \tau}^{t_0} e^{-\mu_2(t_0 - s)} S(s) Q(s) ds > 0$ , and  $Q(t_0) = 0$ , which is contradiction.

■

**Lemma 4** *Let*

$S(s) = S_0 > 0$ ,  $Q(s) = Q_0 > 0$  for all  $s \in [-\tau, 0]$  and  $Q_0 > 0$ .

Then

$$S(t) \leq \max \left\{ \frac{\rho}{\mu_1 + d}, S_0 + I_0 + Q_0 \right\} = M$$

**Proof.** We have

$$\begin{aligned} N(t) &= S(t) + I(t) + Q(t), \\ \dot{N}(t) &\leq \mu - (\mu_1 + d) N(t). \end{aligned}$$

For  $R_0 < 1$  the solutions  $S(t)$ ,  $I(t)$  and  $Q(t)$  approach the disease free equilibrium as  $t \rightarrow \infty$ .

With Eq 2 in the system (3) we have  $\dot{I} \leq -(\mu_2 + d) I$ , hence if  $\mu_2 + d < 0$ ,

$$\lim_{t \rightarrow \infty} I(t) = 0,$$

With Eq 3 in the system (3) we have.

$$\lim_{t \rightarrow \infty} Q(t) = 0,$$

With Eq 1 in the system (3) we obtain  $\dot{S} = \rho - (\mu_1 + d) S$ .

$$\lim_{t \rightarrow \infty} S(t) = \frac{\rho}{\mu_1 + d},$$

Hence.

$$\lim_{t \rightarrow \infty} N(t) = \frac{\rho}{\mu_1 + d}.$$

From lemma1,  $S(t)$ ,  $I(t)$  and  $Q(t)$  solutions of system (1) are positive.

$$S(t) \leq \frac{\rho}{\mu_1 + d}, \text{ for all } t \leq 0.$$

Suppose that

$$N(0) \leq \frac{\rho}{\mu_1 + d}, \text{ then } N(t) \leq \frac{\rho}{\mu_1 + d}$$

On the contrary

If  $N(0) > \frac{\rho}{\mu_1 + d}$  then  $N(t) < N(0)$ , and  $S(t) < N(0)$  for all  $t > 0$ . ■

**Theorem 5** *Let*  $S(s) = S_0 > 0$ ,  $Q(s) = Q_0 > 0$  for all  $s \in [-\tau, 0]$  and  $Q_0 > 0$ .

$E_\tau^*$  is globally asymptotically stable for all  $\tau$

$$\tau > \max \left\{ \frac{1}{\gamma} \ln \frac{\omega M + 3\omega Q_\tau^*}{2\omega(\mu_1 + d)}, \frac{1}{\gamma} \ln \frac{\omega M + 3\omega Q_\tau^*}{2\omega(\mu_2 + d) + (\mu_2 + d) - \beta M}, \frac{1}{\gamma} \ln \frac{Q_\tau^* - 3\omega M}{2(\mu_3 + d) - \beta M} \right\}$$

Where

$$\begin{aligned} M &= \max \left\{ \frac{\rho}{\mu_1 + d}, S_0 + I_0 + Q_0 \right\}, \\ \omega &= \frac{\beta Q_\tau^*}{\mu_1 + \mu_2 + 2d} \end{aligned}$$

**Proof.** We consider system (3).

Let us introduce the functional

$$V(x, y, z) = \frac{1}{2} \omega (x + y)^2 + \frac{1}{2} (y^2 + z^2),$$

The derivative  $\dot{V}(x, y, z)$  is

$$\begin{aligned} \dot{V}(x, y, z) &= \omega (x + y) (\dot{x} + \dot{y}) + y\dot{y} + z\dot{z} \\ &= \omega (x + y) [(-(\mu_1 + d) - \beta Q_\tau^*)x - \beta S_\tau^* z + (\beta - \gamma e^{-\mu_2 \tau}) Q_\tau^* x - (\mu_2 + d)y + (\beta - \gamma e^{-\mu_2 \tau}) Q_\tau^* x - (\mu_2 + d)y + (\beta - \gamma e^{-\mu_2 \tau}) S_\tau^* z] + \\ &\quad + y [\beta \gamma e^{-\mu_2 \tau} Q_\tau^* x - (\mu_2 + d)y + (\beta - \gamma e^{-\mu_2 \tau}) S_\tau^* z] + \\ &\quad + z [\beta \gamma e^{-\mu_2 \tau} Q_\tau^* x + (\gamma e^{-\mu_2 \tau} S_\tau^* - (\mu_3 + d))z] \\ &= -\omega (\mu_1 + d) x^2 - [(\omega + 1)(\mu_2 + d)] y^2 - (\gamma e^{-\mu_2 \tau} S_\tau^* - (\mu_3 + d)) z^2 \\ &\quad + [\beta Q_\tau^* - \omega (\mu_1 + d) - \omega (\mu_2 + d)] xy + \beta S_\tau^* yz - [\omega Q_\tau^* \gamma e^{-\mu_2 \tau}] xx(t - \tau) \\ &\quad - (\omega + 1) Q_\tau^* \gamma e^{-\mu_2 \tau} yx(t - \tau) + Q_\tau^* \gamma e^{-\mu_2 \tau} zx(t - \tau) - \omega S_\tau^* \gamma e^{-\mu_2 \tau} xz(t - \tau) \\ &\quad - (\omega + 1) S_\tau^* \gamma e^{-\mu_2 \tau} yz(t - \tau) + S_\tau^* \gamma e^{-\mu_2 \tau} zz(t - \tau). \end{aligned}$$

By lemma 2 we have  $S(t) \leq M$  for all  $t \geq 0$  and  $\omega$  is an arbitrary real constant choosing as follows

$$\omega = \frac{\beta Q_\tau^*}{\mu_1 + \mu_2 + 2d}$$

$$\begin{aligned} \dot{V}(x, y, z) &\leq -\omega (\mu_1 + d) x^2 - [(\omega + 1)(\mu_2 + d)] y^2 - (\mu_3 + d) z^2 \\ &\quad + \beta M yz - [\omega Q_\tau^* \gamma e^{-\mu_2 \tau}] xx(t - \tau) - (\omega + 1) Q_\tau^* \gamma e^{-\mu_2 \tau} yx(t - \tau) \\ &\quad + Q_\tau^* \gamma e^{-\mu_2 \tau} zx(t - \tau) - \omega M \gamma e^{-\mu_2 \tau} xz(t - \tau) \\ &\quad - (\omega + 1) M \gamma e^{-\mu_2 \tau} yz(t - \tau) + M \gamma e^{-\mu_2 \tau} zz(t - \tau). \end{aligned}$$

Applying Cauchy-Chwartz inequality; we obtain:

$$\begin{aligned}
\dot{V}(x, y, z) &\leq -\omega(\mu_1 + d)x^2 - [(\omega + 1)(\mu_2 + d)]y^2 - (\mu_3 + d)z^2 \\
&\quad - \frac{1}{2}\omega Q_\tau^* \gamma e^{-\mu_2 \tau} [x^2 + x^2(t - \tau)] - \frac{1}{2}(\omega + 1) Q_\tau^* \gamma e^{-\mu_2 \tau} [y^2 + x^2(t - \tau)] \\
&\quad + \frac{1}{2} Q_\tau^* \gamma e^{-\mu_2 \tau} [z^2 + x^2(t - \tau)] - \frac{1}{2}\omega M \gamma e^{-\mu_2 \tau} [x^2 + z^2(t - \tau)] \\
&\quad - \frac{1}{2}(\omega + 1) M \gamma e^{-\mu_2 \tau} [y^2 + z^2(t - \tau)] + \frac{1}{2} M \gamma e^{-\mu_2 \tau} [z^2 + z^2(t - \tau)] \\
&\quad + \frac{1}{2} \beta M [y^2 + z^2], \\
&\leq \left[ -\omega(\mu_1 + d) - \frac{1}{2}\omega \gamma e^{-\mu_2 \tau} (M + Q_\tau^*) \right] x^2 \\
&\quad + \left[ \frac{1}{2} \beta M - (\omega + 1)(\mu_2 + d) - \frac{1}{2}(\omega + 1) \gamma e^{-\mu_2 \tau} (M + Q_\tau^*) \right] y^2 \\
&\quad + \left[ \frac{1}{2} \beta M - (\mu_3 + d) + \frac{1}{2} \gamma e^{-\mu_2 \tau} (M + Q_\tau^*) \right] z^2 \\
&\quad - \left[ \omega Q_\tau^* \gamma e^{-\mu_2 \tau} \right] x^2(t - \tau) - \left[ \frac{1}{2} (3\omega + 1) \gamma M e^{-\mu_2 \tau} \right] z^2(t - \tau)
\end{aligned}$$

Choose the Lyapunov functional

$$V(x_t, y_t, z_t) = V(x, y, z) - \omega Q_\tau^* \gamma e^{-\mu_2 \tau} \int_{t-\tau}^t x^2(\theta) d\theta - \frac{1}{2} (3\omega + 1) \gamma M e^{-\mu_2 \tau} \int_{t-\tau}^t z^2(\theta) d\theta$$

Then

$$\begin{aligned}
\dot{V}(x_t, y_t, z_t) &= \dot{V}(x, y, z) - \omega Q_\tau^* \gamma e^{-\mu_2 \tau} x^2(t) + \omega Q_\tau^* \gamma e^{-\mu_2 \tau} x^2(t - \tau) \\
&\quad - \frac{1}{2} (3\omega + 1) \gamma M e^{-\mu_2 \tau} z^2(t) + \frac{1}{2} (3\omega + 1) \gamma M e^{-\mu_2 \tau} z^2(t - \tau) \\
&\leq \left[ -\omega (\mu_1 + d) - \frac{1}{2} \omega \gamma e^{-\mu_2 \tau} (M + Q_\tau^*) \right] x^2 \\
&\quad + \left[ \frac{1}{2} \beta M - (\omega + 1) (\mu_2 + d) - \frac{1}{2} (\omega + 1) \gamma e^{-\mu_2 \tau} (M + Q_\tau^*) \right] y^2 \\
&\quad + \left[ \frac{1}{2} \beta M - (\mu_3 + d) + \frac{1}{2} \gamma e^{-\mu_2 \tau} (M + Q_\tau^*) \right] z^2 \\
&\quad - [\omega Q_\tau^* \gamma e^{-\mu_2 \tau}] x^2 + \omega Q_\tau^* \gamma e^{-\mu_2 \tau} x^2(t - \tau) \\
&\quad - \left[ \frac{1}{2} (3\omega + 1) \gamma M e^{-\mu_2 \tau} \right] z^2 + \frac{1}{2} (3\omega + 1) \gamma M e^{-\mu_2 \tau} z^2(t - \tau) \\
&\leq \left[ -\omega (\mu_1 + d) - \frac{1}{2} \omega \gamma e^{-\mu_2 \tau} (M + Q_\tau^*) \right] x^2 \\
&\quad + \left[ \frac{1}{2} \beta M - (\omega + 1) (\mu_2 + d) - \frac{1}{2} (\omega + 1) \gamma e^{-\mu_2 \tau} (M + Q_\tau^*) \right] y^2 \\
&\quad + \left[ \frac{1}{2} \beta M - (\mu_3 + d) + \frac{1}{2} \gamma e^{-\mu_2 \tau} (M + Q_\tau^*) \right] z^2 \\
&\quad - [\omega Q_\tau^* \gamma e^{-\mu_2 \tau}] x^2 - \left[ \frac{1}{2} (3\omega + 1) \gamma M e^{-\mu_2 \tau} \right] z^2
\end{aligned}$$

Therefore

$$\begin{aligned}
\dot{V}(x_t, y_t, z_t) &\leq - \left[ \omega (\mu_1 + d) + \frac{1}{2} \omega \gamma e^{-\mu_2 \tau} (M + 3Q_\tau^*) \right] x^2 \\
&\quad - \left[ (\omega + 1) (\mu_2 + d) - \frac{1}{2} \beta M + \frac{1}{2} (\omega + 1) \gamma e^{-\mu_2 \tau} (M + Q_\tau^*) \right] y^2 \\
&\quad - \left[ (\mu_3 + d) - \frac{1}{2} \beta M - \frac{1}{2} \gamma e^{-\mu_2 \tau} (Q_\tau^* - 3\omega M) \right] z^2.
\end{aligned}$$

While the above inequality is always negative provided that

$$\tau > \max \left\{ \frac{1}{\gamma} \ln \frac{\omega M + 3\omega Q_\tau^*}{2\omega (\mu_1 + d)}, \frac{1}{\gamma} \ln \frac{\omega M + 3\omega Q_\tau^*}{2\omega (\mu_2 + d) + (\mu_2 + d) - \beta M}, \frac{1}{\gamma} \ln \frac{Q_\tau^* - 3\omega M}{2(\mu_3 + d) - \beta M} \right\}$$

With application of the Lyapunove-LaSalle type theorem in [10]

$$\lim_{t \rightarrow \infty} x(t) = 0, \quad \lim_{t \rightarrow \infty} y(t) = 0, \quad \lim_{t \rightarrow \infty} z(t) = 0.$$

■



## 4 Stochastic Stability

We limit ourselves here to perturbing only the contact rate so we replace  $\beta$  by  $\beta + \sigma W(t)$ ,

where  $W(t)$  is white noise (Brownian motion). The system (3) is transformed to the following Itô stochastic differential equations, with  $\gamma_0 = \gamma e^{-\mu_2 \tau}$

$$\begin{cases} dS = [\rho - (\mu_1 + d) S - \beta SQ] - \sigma SQ dW, \\ dI = [\beta SQ - (\mu_2 + d) I - \gamma_0 SQ] + \sigma SQ dW, \\ dQ = [\gamma_0 SQ - (\mu_3 + d) Q], \end{cases} \quad (12)$$

In this section, we will proof, under some conditions, that  $E_0$  is globally exponentially mean square and almost surely stable, and for this purpose, we need the following Theorem

**Theorem 6** *The set  $\Omega$  is almost surely invariant by the stochastic system (12). Thus if  $(S_0, I_0, Q_0) \in \Omega$ , then  $P[(S, I, Q) \in \Omega] = 1$ .*

**Proof.** The system (12) implies that  $dN \leq [\rho - (\mu_1 + d) N] dt$ , then we have

$$N(t) \leq \frac{\rho}{\mu_1 + d} + \left( N_0 - \frac{\rho}{\mu_1 + d} \right), \text{ for all } t \geq 0.$$

Since  $(S_0, I_0, Q_0) \in \Omega$ , then

$$N(t) \leq \frac{\rho}{\mu_1 + d}, \text{ for all } t \geq 0. \quad (13)$$

There exist  $\varepsilon_0 > 0$ , such that  $S_0 > \varepsilon_0 > 0$ ,  $I_0 > \varepsilon_0 > 0$  and  $Q_0 > \varepsilon_0 > 0$ . Considering

$$\begin{aligned} v_\varepsilon &= \inf \{ t \geq 0, S(t) \leq \varepsilon \text{ or } I(t) \leq \varepsilon \text{ or } Q(t) \leq \varepsilon, \}, \text{ for } \varepsilon \leq \varepsilon_0, \\ v &= \lim_{t \rightarrow 0} v_\varepsilon = \inf \{ t \geq 0, S(t) \leq 0 \text{ or } I(t) \leq 0 \text{ or } Q(t) \leq 0, \} \end{aligned} \quad (14)$$

Let

$$V(t) = \log \frac{\rho}{(\mu_1 + d) S(t)} + \log \frac{\rho}{(\mu_1 + d) I(t)} + \log \frac{\rho}{(\mu_1 + d) Q(t)}.$$

Then, using Itô formula we have, for all  $t \geq 0$  and  $T \in [0, t \wedge v_\varepsilon]$ ,

$$\begin{aligned} dV(T) &= \left[ -\frac{\rho}{S} + (\mu_1 + d) + \beta Q + \frac{1}{2} I^2 \right] dT + \sigma Q dW \\ &\quad + \left[ (\gamma_0 - \beta) \frac{SQ}{I} + (\mu_2 + d) + \frac{1}{2} S^2 \right] dT + \sigma \frac{SQ}{I} dW \\ &\quad + [-\gamma_0 S + (\mu_3 + d)] dT, \end{aligned}$$

$$dV(T) \leq \left[ \mu_1 + \mu_2 + \mu_3 + 3d + \beta Q + \frac{1}{2} I^2 + \frac{1}{2} S^2 \right] dT + \sigma \frac{Q}{I} (I - S) dW \quad (15)$$

With (13), we have  $S, I$  and  $Q \in \left[0, \frac{\rho}{(\mu_1+d)}\right]$   
Let

$$\begin{aligned} L &= \mu_1 + \mu_2 + \mu_3 + 3d + \beta \frac{\rho}{\mu_1 + d} + \left(\frac{\rho}{\mu_1 + d}\right)^2, \\ f(I) &= \frac{Q}{I}, \end{aligned} \quad (16)$$

We remplace (16) into (15), we obtain

$$dV(T) \leq LdT + \sigma(I(T) - S(T))f(I(T))dW, \quad (17)$$

Then

$$V(T) \leq LT + \sigma \int_0^T f(I(u))(I(u) - S(u))dW(u), \quad (18)$$

With proposition 7.6 in [6],  $\sigma \int_0^T f(I(u))(I(u) - S(u))dW(u)$  is mean zero process then,

$$E(V(T)) \leq LT \quad (19)$$

for all  $t \geq 0$  and  $T \in [0, t \wedge v_\varepsilon]$ ,

$$S(t \wedge v_\varepsilon), I(t \wedge v_\varepsilon), \text{ and } Q(t \wedge v_\varepsilon) \in \left[0, \frac{\rho}{(\mu_1 + d)}\right],$$

Then

$$E(V(t \wedge v_\varepsilon)) \leq L(t \wedge v_\varepsilon) \leq Lt,$$

$$V(t \wedge v_\varepsilon) \geq 0,$$

$$E(V(t \wedge v_\varepsilon)) \geq E(V(t)) \times \mathcal{X}_{[v_\varepsilon \leq t]} \geq P(v_\varepsilon \leq t) \log \frac{\rho}{(\mu_1 + d)\varepsilon} \quad (20)$$

Where  $\mathcal{X}_{[v_\varepsilon \leq t]}$  is the indicator function of a subset  $[v_\varepsilon \leq t]$ ,  
Combining (19), and (20), we obtain

$$P(v_\varepsilon \leq t) \leq \frac{Lt}{\log \frac{\rho}{(\mu_1 + d)\varepsilon}}, \text{ for all } t \geq 0; \quad (21)$$

for all  $t \geq 0$ , and  $\varepsilon \rightarrow 0$ , we obtain  $P(v \leq t) = 0$ ;

From where

$$P(v \leq \infty) = 0$$

■

## 5 The Model with Age

The age distributions of the numbers in the classes are denoted by  $S(a, t)$ ,  $I(a, t)$ , and  $Q(a, t)$ , denote the sizes of the population susceptible to disease, and infectious members, quarantine members with the possibility of infection through temporary immunity, respectively of age  $a$ , at time  $t$ ,  $d(a)$  is the age-specific death rate,

The system of partial equations for the age distributions is

$$\begin{cases} \frac{\partial S}{\partial t} + \frac{\partial S}{\partial a} = -(\mu_1 + d(a)) S(a, t) + \beta_1(t) S(a, t), \\ \frac{\partial I}{\partial t} + \frac{\partial I}{\partial a} = -\beta_1(t) S(a, t) - (\mu_2 + d(a)) I(a, t) + \gamma_1(t - \tau) S(a, t - \tau), \\ \frac{\partial Q}{\partial t} + \frac{\partial Q}{\partial a} = -\gamma_1(t - \tau) S(a, t - \tau) - (\mu_3 + d(a)) Q(a, t), \end{cases} \quad (22)$$

With

$$\begin{aligned} \beta_1(t) &= -\beta \int Q(a, t) da \\ \gamma_1(t - \tau) &= -\gamma \int e^{-\mu_2 \tau} Q(a, t - \tau) da \end{aligned} \quad (23)$$

### 5.1 Equilibrium and stability

Assume that sub population does not depend on the time when the system (22) is written as follows

$$\begin{cases} \frac{dS}{da} = (\beta_1 - \mu_1 - d(a)) S(a), \\ \frac{dI}{da} = (\gamma_1 - \beta_1) S(a) - (\mu_2 + d(a)) I(a), \\ \frac{dQ}{da} = -\gamma_1 S(a) - (\mu_3 + d(a)) Q(a), \end{cases} \quad (24)$$

The initial condition of (24) is given as

$$S(0) = S_1, \quad I(0) = I_1, \quad Q(0) = Q_1 \quad (25)$$

Differential equations of the system (24) are solved with different methods of resolutions and with (25), so

$$S(a) = S_1 e^{-(\mu_1 - \beta_1)a} \Phi(a), \quad (26)$$

$$I(a) = I_1 \Phi(a) e^{-\mu_2 a} - \frac{(\gamma_1 - \beta_1) S_1 \Phi(a)}{\mu_1 - \beta_1 - \mu_2} \left( e^{-(\mu_1 - \beta_1)a} - e^{-\mu_2 a} \right), \quad (27)$$

$$Q(a) = Q_1 \Phi(a) e^{-\mu_3 a} - \frac{\gamma_1 S_1 \Phi(a)}{\mu_1 - \beta_1 - \mu_3} \left( e^{-(\mu_1 - \beta_1)a} - e^{-\mu_3 a} \right) \quad (28)$$

Where

$$\Phi(a) = \exp \left( -\int d(a) da \right) \quad (29)$$

The system (24) has the unique positive equilibrium point  $P_1$ ,

$$P_1 = \left( \hat{S}_1, \hat{I}_1, \hat{Q}_1 \right)^T = (0, 0, 0)^T.$$

We calculate the Jacobian matrix according to the system (24) with  $P_1$

$$J(P_1) = \begin{bmatrix} \beta_1 - \mu_1 - d(a) & 0 & 0 \\ \lambda - \gamma_0 & -(\mu_2 + d(a)) & 0 \\ -\gamma_0 & 0 & -(\mu_3 + d(a)) \end{bmatrix}$$

The epidemic is locally asymptotically stable if and only if all eigenvalues of the Jacobian matrix  $J(P_1)$  have negative real part. The eigenvalues can be determined by solving the characteristic equation of the linearization of (25) near  $P_1$  is

$$\det \begin{pmatrix} \beta_1 - \mu_1 - d(a) - A & 0 & 0 \\ \lambda - \gamma_0 & -(\mu_2 + d(a)) - A & 0 \\ -\gamma_0 & 0 & -(\mu_3 + d(a)) - A \end{pmatrix} = 0 \quad (31)$$

So the eigenvalues are

$$A_1 = \beta_1 - \mu_1 - d(a), \quad A_2 = -(\mu_2 + d(a)), \quad A_3 = -(\mu_3 + d(a))$$

In order to  $A_1, A_2,$  and  $A_3$  will be negative, it is required that

$$\beta_1 < \mu_1 + d(a)$$

The basic reproduction number  $R_0$  is defined as the total number of infected population in the resulting sub-infected population where almost all of the uninfected. The basic reproduction number of the infection  $R_0$  is defined as follows:

$$R_0 = \frac{\beta_1}{\mu_1 + d(a)} \quad (32)$$

The time during which people remain infective is defined as

$$T = \frac{1}{\mu_1 + d(a)}$$

The doubling time  $t_d$  of the epidemic can be obtained as

$$t_d = \frac{(\ln 2) T}{R_0 - 1} \quad (33)$$

**Theorem 1** The disease-free equilibrium  $P_1$  is locally asymptotically stable if  $R_0 < 1$  and unstable if  $R_0 > 1$ .

Let (26), so if  $R_0 < 1$  then  $\mu_1 - \beta_1 > 0$ , so  $S(a)$  converges to zero.

Let (27), so

$$I(a) \leq \left[ I_1 \Phi(a) - \frac{(\gamma_1 - \beta_1) S_1 \Phi(a)}{\mu_1 - \beta_1 - \mu_2} \right] e^{-m_1 a}, \quad m_1 = \min \{\mu_1 - \beta_1, \mu_2\} \quad (34)$$

If  $R_0 < 1$ ,  $i(a)$  converges to zero.

Let (28), so

$$Q(a) = \left[ Q_1 \Phi(a) - \frac{\gamma_1 S_1 \Phi(a)}{\mu_1 - \beta_1 - \mu_3} \right] e^{-m_2 a}, \quad m_1 = \min \{\mu_1 - \beta_1, \mu_3\} \quad (35)$$

If  $R_0 < 1$ ,  $Q(a)$  converges to zero.

This paper addresses a SIQ model with temporary immunity, whenever the quarantine individuals will return to the susceptible. The endemic equilibrium is globally asymptotically stable, then under some conditions, study the stochastic stability. Finally, the equilibrium and stability of the epidemic model with age.

## References

- [1] Anderson R. M and Medley R M and Jhonson A K. A Preliminary Study of the Transmission Dynamics of the Human Immunodeficiency Virus (HIV), the Causative Agent of AIDS. IMA. J. Math. Appl. Med. Biol 3, 229-263.(1986).
- [2] Abta A and Kaddar A and Talibi H. A. Global Stability for Delay SIR and SEIR Epidemic Models With Saturated Incidence Rates. Electronic Journal of Differential Equations, 23,1-13.(2012).
- [3] Bailey. N.T.J. Some Stochastic Models for Small Epidemics in Large Population. Appl. Statist.13, 9-19.(1964).
- [4] Bailey. N.T.J. The Mathematical Theory of Infection Diseases and its Application. Applied Statistics, 26, N1, 85-87.(1977).
- [5] Batiha, M. S. M. Noorani and I. Hashim. Numerical solutions of the non-linear integro-differential equations, Int. J. Open Probl. Compt. Math, 34-42.(2008).
- [6] Becker.N.G. The Uses of Epidemic Models. *Biometrics* **35**, 295-305. (1979).
- [7] Billard.L. A Stochastic General Epidemic in m Sub-Population. *J. Appl. Prob.* **13**, 567-572. (1976).
- [8] Jinliang W, Xinxin Tian. Global Stabily of a Delay Differential Equation Of Hepatitis B Virus Infection With Immune Response. Electronic Journal of Differential Equations,94,1-11. (2013).
- [9] Jin. Z, Zhien. M and Maoan. H. Globale stability of an SIRS epidemic model with delay, Acta Matimatica Scientia. 26 B. 291-306.(2006).

- [10] Kuang Y. Delay-Differential Equations with application in population biology. Academic Press, new york.(1993) .
- [11] Lounes. R and Arazoza. H. Modeling HIV Epidemic Under Contact Tracing. The Cuban Case. Journal of theoritical Medecine Vol 2, 267-274.(2000) .
- [12] Lounes. R, Arazoza. H. A Non-Linear Model for a Sexually Transmitted Disease with contact tracing. IMA. J. MJath. Appl. Med. Biol.19, 221-234.(2002) .
- [13] Lahrouz A and El Maroufy H. Qualitative Behaviour of a Model of an SIRS Epidemic:Stability and Permanence. Applied Mathematics & Information Sciences. An International Journal 5 (2), 220-238.(2011).
- [14] Luo Q and Mao X. Stochastic population dynamics under regime switching. J. Math. Anal. Appl.334, 69-84.(2007)
- [15] Michael Steel J. Stochastic calculus and finantial applications. Springer-Verlag. (2003) .
- [16] Naresh R, and Omar S. An epidemic model for the transmission dynamics of HIV/AIDS and another infection. International Journal of Mathematical Archive-1(3) , 68-72. (2010) .
- [17] Perto. L. Differential Equations and Dynamical Systems. 2nd edition, Springer, New York.(1996) .
- [18] Ray Waston. A useful Random Time-Seal Transformation For The Standard Epidemic Model. J. Appl. Prob.17, 324-332. (1980) .
- [19] Ray Waston, 1980. On The Size Distribution For Some Epidemic Models. J. Appl. Prob.17, 912-921.(1980) .
- [20] Robert N and May. Population Biology of infectious diseases I. International centre of theoritical physics.1-9. (1982) .
- [21] Ruoyan Sun. Global stability of the endemic equilibrium of multigroup SIR models with nonlinear incidence. Computers and Mathematics with Applications 60. 2286-2291. (2010) .
- [22] Takeuchi and W. Ma. Stability analysis on a delayed SIR epidemic model with density dependent birth process, Dy-nam. Contin. Discrete Impuls. Systems, 5 . 171-184.(1999).
- [23] Volodymyr Makarov, Denis Dragunov. A numeric-analytical method for solving the Cauchy problem for ordinary diferential equations. Applied Mathematics and Computation,1-26. (2010) .

- [24] W. Ma, Y. Takeuchi, T. Hara and E. Beretta, Permanence of are SIR epidemic model with distributed time delays, *Tohoku Math. J.* 54, 581-591.(2002)
- [25] W. Wang, Global behavior of an SEIR epidemic model with time delay, *Appl. Math. Letters*.15, 423-428.(2002).
- [26] Wen L and Yang X. Global stability of a delayed SIRS model with temporary immunity. *Chaos, Solitons and Fractals* 38, 221-226. (2008)
- [27] Xiao, L Chen and F. ven den Bosch, Dynamical behavior for a stage-structured SIR infectious disease model, *Nonlinear Anal. Real World Appl* 3,175-190.(2002).
- [28] Z. Ma, J. Liu and J. Li, Stability analysis for differential infectivity epidemic models, *Nonlinear Anal. Real World Appl* 4, 841-856. (2003).
- [29] Zhang F and Zhen Li and Zhang F. Global stability of an SIRepidemic model with constant infectious period. *Applied Mathematics and Computation* 199, 285-291.(2008).