Dynamical Analysis and Design of Computational Methods for Nonlinear Stochastic Leprosy Epidemic Model

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Dynamical analysis and design of computational methods for nonlinear stochastic leprosy epidemic model

Ali Raza · Jan Awrejcewicz · Muhammad Rafiq · Muhammad Mohsin

Abstract In this article, we present the dynamical analysis of the stochastic leprosy epidemic model. Positivity and boundedness are the criteria used in the deterministic model. A primary technique known as the Euler Maruyama is employed in the solution of the said model. Standard and non-standard computational methods are applied in evaluating the design stability and efficiency based on the chosen criteria. The standard computational methods like the Stochastic Euler and the Stochastic Runge Kutta fail to restore the essential features of biological problems. However, our proposed approach, the stochastic non-standard finite difference (NSFD), is used and found to be efficient, cost-effective, and accommodates all the desired feasible properties. Our method achieves all-time convergence against the backdrop of other classical techniques that perform conditionally or fail over a long period. In the end, a comparison between this scheme and the existing ones reviews the novelty of our approach.

Keywords Leprosy disease model · Stochastic differential equations · Computational methods · Stability analysis

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1 Introduction

In 2018, Giraldo et al. studied a mathematical model based on nonlinear ordinary differential equations. The model explains the transmission dynamics of paucibacillary leprosy (PB) and multibacillary leprosy (MB) [1]. In 2018, Varella et al. presented the mathematical model related to leprosy type infection: a case study of the spread of leprosy in Brazil [2]. In 2021, Marathi et al. developed a multi-dimensional mathematical model for the transmission of leprosy disease [3]. In 2016, Smith captured a compartmental model representing the present knowledge of susceptibility and pathogenesis to leprosy, adapted to different populations [4]. In 2007, Abubakar et al. implemented the Markov decision model to decide the best cost of drugs for the treatment of leprosy disease [5]. In 2012, Mushayabase et al. investigated the dynamics of leprosy on behalf of treatment and control measures [6]. In 2013, Ghayas et al. studied leprosy disease transmission dynamics: a case of Gilgit-Baltistan, Pakistan [7]. In 2013, Deps et al. linked the co-infection among leprosy and HIV [8]. In 2013, Chiyaka et al. developed numerical and theoretical analysis for transmission dynamics of leprosy disease [9]. Similarly, Arco et al., in 2016, diagnosed the leprosy symptoms and transmission modes [10], while Le et al., in 2018, investigated the control plans for the disease-like infections: a case study of monitoring and detection in China [11]. In 2017, Nobre et al. presented a high-risk analysis of leprosy infection like the multibacillary [12]. Ezenduka et al., in 2012, analyzed the cost-effectiveness of the three methods of detecting leprosy in Northern Nigeria to determine the most viable plan for locating the disease [13]. In 2009, Koba et al. analyzed the trends and patterns of native leprosy in Japan [14]. However, in the countries of the Amazon region, not until 2020 when Schaub et al. in [15] demonstrated the prevalence of distinct features of leprosy among humans and animal reservoirs. Kumar et al., in 2007, came out with a survey of leprosy disease among non-family contacts (NFC) and the family contacts (FC) of patients in Agra, India [16], while Fischer et al., in 2008, assessed the local distribution of leprosy cases in Bangladesh during 15-years of disease control program in Bangladesh [17].

The rest of this paper is organized based on the following sections: formulation of the deterministic leprosy epidemic model with fundamental properties is introduced in section 2. In section 3, the transition probabilities, positivity, boundedness and implementation of the well-known methods, including the NSFD, and their comparative analysis are exposed. Finally, section 4 offers the concluding remarks.

2 Formulation of the model

In this part, we studied the dynamics of a leprosy disease in humans using a mathematical model in which the human population is divided into four subpopulations, respectively. The total number is represented by \( N : [0, \infty) \rightarrow \mathbb{R} \) with the function of time \( t \geq 0 \). Furthermore, each of the subcomponents of
the population is denoted by a non-negative differentiable function $x, y, z, w : [0, \infty) \to \mathbb{R}$. The representation of each of this subdivision is described as follows: $x(t)$ depicts the susceptible individuals, $y(t)$ denotes the asymptomatic infected individuals, $z(t)$ represents the infected individuals by multibacillary leprosy, $w(t)$ shows the infected individuals by paucibacillary leprosy, and $N(t)$ represents the total population. The systematic flow of leprosy disease is presented in Fig. 1.

The physical interpretation of the model is presented as follows: $f$ represents a fraction of humans who developed multibacillary leprosy, $1-f$ portrays a fraction of humans who developed paucibacillary leprosy, $\rho$ describes the natural growth rate of humans, $\beta_p$ delineates the contact rate of paucibacillary leprosy individuals, $\beta_m$ shows the contact rate of multibacillary leprosy individuals, $\theta$ presents the rate at which asymptomatic individuals may be symptomatic individuals, $\mu_m$ depicts the death rate of the infected individuals by multibacillary leprosy, while $\mu$ is the natural death rate of individuals. The model of the differential equations is as given below:

$$\frac{dx(t)}{dt} = \rho - \left( \beta_m x(t) \frac{z(t)}{N(t)} + \beta_p x(t) \frac{w(t)}{N(t)} \right) - \mu x(t), \quad t \geq 0, \quad (1)$$

$$\frac{dy(t)}{dt} = \beta_m x(t) \frac{z(t)}{N(t)} + \beta_p x(t) \frac{w(t)}{N(t)} - \theta y(t) - \mu y(t), \quad t \geq 0, \quad (2)$$

$$\frac{dz(t)}{dt} = f \theta y(t) - \mu z(t), \quad t \geq 0, \quad (3)$$

$$\frac{dw(t)}{dt} = \theta (1-f) y(t) - \mu w(t), \quad t \geq 0, \quad (4)$$

$$\frac{dN(t)}{dt} = \rho - \mu (x(t) + w(t) + y(t)) - \mu z(t), \quad t \geq 0 \quad (5)$$

Fig. 1 Flow map of leprosy disease
with non-negative initial conditions \( x(0) \geq 0, \ y(0) \geq 0, \ \omega(0) \geq 0, \ z(0) \geq 0 \).

2.1 Model analysis

In this section, we discuss feasible region of the model. The variables \( x(t), \ y(t), \ \omega(t) \) and \( z(t) \) must be non-negative. Therefore, the obtainable results of the epidemic model remain positive and bounded in the region \( \mathcal{M} = \{(x, y, z, \omega) \in \mathbb{R}^4_+ : N(t) \leq \frac{\kappa}{\mu}, \ x \geq 0, y \geq 0, z \geq 0, \omega \geq 0 \} \) at any time \( t \geq 0 \).

2.1.1 Positivity of the model

**Theorem 1** The solution \( (x, y, \omega, z) \in \mathbb{R}^4_+ \) of the system (1)-(4) is positive at any time \( t \geq 0 \) given any non-negative initial conditions.

**Proof:** It is clear from equations (1)-(4) that:

\[
\left. \frac{dx}{dt} \right|_{x=0} = \kappa \geq 0, \quad \left. \frac{dy}{dt} \right|_{y=0} = (\beta_m x z + \beta_p x \omega) \geq 0, \quad \left. \frac{dz}{dt} \right|_{z=0} = \varrho y \geq 0
\]

and

\[
\left. \frac{d\omega}{dt} \right|_{\omega=0} = \theta (1 - f) y \geq 0,
\]

as desired.

2.1.2 Boundedness of the model

**Theorem 2** The solution of the system (1)-(4) is bounded and lie in the feasible region \( \mathcal{M} \)

**Proof:** Let us consider the population density function as follows:

\[
N(t) = x(t) + y(t) + \omega(t) + z(t),
\]

\[
\frac{dN}{dt} = \kappa - \mu N.
\]

From Gronwall’s inequality,

\[
N(t) = N(0) + \frac{\kappa}{\mu}, \quad t \geq 0
\]

\[
\lim_{t \to \infty} \sup N(t) \leq \frac{\rho}{\mu}.
\]

So, it is clear that the solution of (1)-(4) is bounded and lies in \( \mathcal{M} \).
2.1.3 Model equilibria

Considering equations (1)-(4) in which the state variables are constant, we have

\[ \kappa - (\beta_m x z + \beta_p x \omega) - \mu x = 0, \]  

(6)

\[ (\beta_m x z + \beta_p x \omega) - \mu y - \theta y = 0, \]  

(7)

\[ f \theta y - \mu z = 0, \]  

(8)

\[ (1 - f) \theta y - \mu \omega = 0. \]  

(9)

From equations (9), (8), (7) and (6), we get

\[ \omega^* = \frac{(1 - f) \theta y^*}{\mu}, \quad z^* = \frac{f \theta y^*}{\mu}, \quad x^* = \frac{(\mu + \theta)}{(\beta_m f \theta + \beta_p (1 - f) \theta)} \]

\[ \rho - (\beta_m x z + \beta_p x \omega) - \mu x = 0, \]

\[ \rho \left[ \frac{\beta_m f \theta + \beta_p (1 - f) \theta}{\mu (\mu + \theta)} \right] y + \mu = 0, \]

\[ \rho \left[ \frac{\beta_m f \theta + \beta_p (1 - f) \theta}{\mu (\mu + \theta)} \right] = y \left[ \frac{\beta_m f \theta + \beta_p (1 - f) \theta}{\mu} \right] + \mu, \]

\[ \frac{\rho}{\mu + \theta} - \frac{\mu}{[\beta_m f \theta + \beta_p (1 - f) \theta]} = y, \]

\[ g^* = \frac{\rho [\beta_m f \theta + \beta_p (1 - f) \theta] - \mu^2 (\mu + \theta)}{(\mu + \theta) [\beta_m f \theta + \beta_p (1 - f) \theta]}. \]

Thus, the endemic equilibrium (EE \( L_2 \)) = \( (x^*, y^*, \omega^*, z^*) \), where

\[ x^* = \frac{u (\theta + u)}{[\beta_m f \theta + \beta_p (1 - f) \theta]}, \quad y^* = \frac{[\beta_m f \theta + \beta_p (1 - f) \theta] \kappa - \mu^2 (\theta + u)}{(\theta + u) [\beta_m f \theta + \beta_p (1 - f) \theta]}, \]

\[ \omega^* = \frac{\theta (1 - f) [\beta_m f \theta + \beta_p (1 - f) \theta] \kappa - \mu^2 (\theta + u)}{u (\theta + u) [\beta_m f \theta + \beta_p (1 - f) \theta]}, \]

\[ z^* = \frac{f \theta [\beta_m f \theta + \beta_p (1 - f) \theta] \kappa - \mu^2 (\theta + u)}{u (\theta + u) [\beta_m f \theta + \beta_p (1 - f) \theta]}, \]

And also, the disease-free equilibrium (DFE \( L_1 \)) of the system (7)-(9) is as follows:

\[ x \neq 0, \quad y = 0, \quad z = 0, \quad \omega = 0. \]

Thus, the disease-free equilibrium (DFE \( L_1 \)) = \( (x^1, y^1, \omega^1, z^1) = (\kappa/\mu, 0, 0, 0) \).
2.2 Local stability

In this section, we present two well-known theorems for local stability. Again, consider the system (1)-(4) as a function of $F, G, H$ and $K$ as follows:

\[ F = \kappa - (\beta_m x + \beta_p x \omega) - \mu x, \quad G = (\beta_m x + \beta_p x \omega) \theta y - \mu y, \]
\[ H = f \theta y - \mu z, \quad K = \theta (1 - f) y - \mu \omega. \]

The partial derivatives of the function regarding state variables are as follows:

\[ \frac{\partial F}{\partial x} = - (\beta_m z + \beta_p \omega) - \mu, \quad \frac{\partial F}{\partial y} = 0, \quad \frac{\partial F}{\partial z} = -\beta_m x, \quad \frac{\partial F}{\partial \omega} = -\beta_p x \omega, \]
\[ \frac{\partial G}{\partial x} = (\beta_m z + \beta_p \omega), \quad \frac{\partial G}{\partial y} = \theta - \mu, \quad \frac{\partial G}{\partial z} = \beta_m x, \quad \frac{\partial G}{\partial \omega} = \beta_p x \omega, \]
\[ \frac{\partial H}{\partial x} = 0, \quad \frac{\partial H}{\partial y} = \theta f, \quad \frac{\partial H}{\partial z} = -\mu, \quad \frac{\partial H}{\partial \omega} = 0, \]
\[ \frac{\partial K}{\partial x} = 0, \quad \frac{\partial K}{\partial y} = \theta (1 - f), \quad \frac{\partial K}{\partial z} = 0, \quad \frac{\partial K}{\partial \omega} = -\mu. \]

Thus, the Jacobian matrix of the model takes the following form

\[
J = \begin{bmatrix}
- (\beta_m z + \beta_p \omega) - \mu & 0 & -\beta_m x & -\beta_p x \\
(\beta_m z + \beta_p \omega) & \theta - \mu & \beta_m x & \beta_p x \\
0 & f \theta & -\mu & 0 \\
0 & \theta (1 - f) & 0 & -\mu
\end{bmatrix}.
\]

**Theorem 3** The disease-free equilibrium $L_1 = (x^1, y^1, z^1, \omega^1) = \left( \frac{\kappa}{\mu}, 0, 0, 0 \right)$ is locally and asymptotically stable (LAS) if $R_0 < 1$. Otherwise, the system is unstable if $R_0 > 1$.

**Proof:** The Jacobian matrix evaluated at $L_1$ is as follows

\[
J_{L_1} = \begin{bmatrix}
- \mu & 0 & -\beta_m x & -\beta_p x \\
0 & \theta - \mu & \beta_m x & \beta_p x \\
0 & f \theta & -\mu & 0 \\
0 & \theta (1 - f) & 0 & -\mu
\end{bmatrix}.
\]

\[
|J - \lambda I| = \begin{vmatrix}
- \mu - \lambda & 0 & -\beta_m x & -\beta_p x \\
0 & \theta - \mu - \lambda & \beta_m x & \beta_p x \\
0 & f \theta & -\mu - \lambda & 0 \\
0 & \theta (1 - f) & 0 & -\mu - \lambda
\end{vmatrix} = 0,
\]

$\lambda_1 = - \mu < 0,$

\[
\begin{vmatrix}
\theta - \mu - \lambda & \beta_m x & \beta_p x \\
f \theta & -\mu - \lambda & 0 \\
f \theta (1 - f) & 0 & -\mu - \lambda
\end{vmatrix} = 0.
\]
Letting $a_1 = \theta + \mu$, $a_2 = \beta_m \frac{z^*}{\mu}$, $a_3 = \beta p x^*$, $a_4 = \theta (1 - f)$, the following characteristic equation is obtained

$$
\begin{vmatrix}
-a_1 - \lambda & a_2 & a_3 \\
\theta f & -\mu - \lambda & 0 \\
-a_4 & 0 & -\mu - \lambda
\end{vmatrix} = 0
$$

which is transformed into its counterpart polynomial form as

$$
\lambda^3 + \lambda^2 (a_1 + 2\mu) + \lambda (-2a_1\mu - \mu^2 - a_2\theta - a_3a_4) + (a_4\mu^2 - a_2\theta\mu - a_3a_4) = 0.
$$

If we examine the coefficients of the characteristic equation above as

$$(a_1 + 2\mu) (a_4\mu^2 - a_2\theta\mu - a_3a_4) > 0$$

and

$$(a_4\mu^2 - a_2\theta\mu - a_3a_4) (-2a_1\mu - \mu^2 - a_2\theta - a_3a_4) > (a_1 + 2\mu).$$

The Routh-Hurwitz stability criterion for the 3rd-degree polynomial is satisfied, and hence the disease-free equilibrium is stable.

**Theorem 4** The endemic equilibrium $(EE - L_2)$, $L_2 = (x^*, y^*, z^*, w^*)$ is locally and asymptotically stable (LAS) if $R_0 > 1$.

**Proof:** The Jacobian matrix of the system at $L_2$ is as follows:

$$
J_{L_2} = 
\begin{bmatrix}
-(\beta_m z^* + \beta_p x^*) - \mu & 0 & -\beta_m x^* & -\beta px^* \\
(\beta_m z^* + \beta_p x^*) & \theta - \mu & \beta_m x^* & \beta px^* \\
0 & \theta f & -\mu & 0 \\
0 & \theta (1 - f) & 0 & -\mu
\end{bmatrix}
$$

Substituting $b_1 = (\beta_m z^* + \beta_p x^*) > 0$, $b_2 = \beta_m x^* > 0$, $b_3 = \beta px^* > 0$, $b_4 = \theta > 0$, $b_5 = f\theta > 0$, $b_6 = \mu > 0$ and $b_7 = \theta (1 - f)$ and proceeding in the same manner as before, we have:

$$
|J_{L_2} - \lambda I| = 
\begin{vmatrix}
-b_1 - \mu - \lambda & 0 & -b_2 & -b_3 \\
b_1 & -b_4 - \lambda & b_2 & b_3 \\
0 & b_5 & -b_6 - \lambda & 0 \\
0 & b_7 & 0 & -b_6 - \lambda
\end{vmatrix} = 0.
$$

Consequently, the characteristic equation becomes

$$
\lambda^4 + (b_1 + b_4 + 2b_6 + \mu) \lambda^3 + (b_3b_7 - b_1b_6 - b_6\mu - b_6^2 - b_4b_6 + b_2b_5)\lambda^2 + (b_3b_7b_6 + b_2b_5b_7 + b_2b_6b_5 - b_1b_6^2)\lambda + (b_3b_7b_6\mu + b_2b_5b_6\mu + b_1b_4b_6^2\mu) = 0,
$$
where
\[ n_0 = (b_1 + b_4 + 2b_6 + \mu), \]
\[ n_1 = (b_3b_7 - b_1b_6 - b_6\mu - b_6^2 - b_4b_6 - b_2b_5 + b_1b_6 - b_1b_4 - b_6\mu - b_4\mu - b_4b_6), \]
\[ n_2 = (b_3b_6b_7 + b_3\mu b_7 + b_2b_6b_5 - b_1b_6^2 - b_1b_4b_6 - \mu b_6^2 - b_2b_6^2 - b_4b_6\mu + b_2b_6\mu - b_1b_4b_6 - b_4b_6^2), \]
\[ n_3 = (b_3b_6b_7 + b_2b_6\mu + b_1b_4b_6^2 \mu). \]

We observe that the Routh-Hurwitz stability criterion for the 4th-degree polynomial is also satisfied, since
\[ n_0 > 0, \quad n_1 > 0, \quad n_1n_2 - n_0n_3 > 0, \]
\[ (n_1n_2 - n_0n_3)n_3 - n_1^2n_4 > 0 \quad \text{and} \quad n_4 > 0 \quad \text{only if} \quad R_0 > 1. \]

Hence, the endemic equilibrium is locally stable.

2.3 Reproduction number

In this section, we investigate the next-generation matrix method to find the transmission and transition matrices \((F\) and \(V\) respectively) of the system (1)-(4) after substituting the disease-free equilibrium as follows:
\[
\begin{bmatrix}
    y' \\
    z' \\
    u'
\end{bmatrix} = \begin{bmatrix}
    0 & \beta_n x & \beta_n x \\
    0 & 0 & 0 \\
    0 & 0 & 0
\end{bmatrix} \begin{bmatrix}
    y \\
    z \\
    w
\end{bmatrix} = \begin{bmatrix}
    \theta + u & 0 & 0 \\
    -f\theta & u & 0 \\
    -\theta (1 - f) & 0 & \mu
\end{bmatrix} \begin{bmatrix}
    y \\
    z \\
    w
\end{bmatrix},
\]
\[ F = \begin{bmatrix}
    0 & \beta_n \frac{z}{\mu} & \beta_n \frac{z}{\mu} \\
    0 & 0 & 0 \\
    0 & 0 & 0
\end{bmatrix}, \quad V = \begin{bmatrix}
    \theta + u & 0 & 0 \\
    -f\theta & \mu & 0 \\
    -\theta(1 - f) & 0 & \mu
\end{bmatrix}, \]
\[ |V| = \mu^2(\theta + u), \quad V^{-1} = \frac{\text{Adj}(V)}{|v|}, \]
\[ V^{-1} = \frac{1}{\mu^2(\theta + u)} \begin{bmatrix}
    \mu f \theta & 0 & 0 \\
    \mu f \theta & \mu(\theta + u) & 0 \\
    \mu \theta(1 - f) & 0 & \mu(\theta + u)
\end{bmatrix}, \]
\[ G = FV^{-1} = \begin{bmatrix}
    \beta_n \kappa f \theta + \beta_n \mu u \theta(1 - f) & \beta_n \kappa & \beta_n \kappa \\
    \mu^2(\theta + u) & 0 & 0 \\
    0 & 0 & 0
\end{bmatrix} \]
\[ |G - \lambda| = \begin{vmatrix}
    \beta_n \kappa f \theta + \beta_n \mu u \theta(1 - f) & 0 & 0 \\
    \mu^2(\theta + u) & -\lambda & 0 \\
    0 & 0 & -\lambda
\end{vmatrix} = 0, \]
\[
\left( \frac{\beta_m f \theta + \beta_p \theta (1 - f)}{\mu^2 \mu_m (\theta + u)} \right) - \lambda \right) \mu^2 = 0.
\]

The dominant eigenvalue of the matrix \( F V^{-1} \) is called the reproduction number and is denoted as

\[ R_0 = \left( \frac{\beta_m f \theta + \beta_p \theta (1 - f)}{\mu^2 (\theta + u)} \right) \kappa. \]

2.4 Stochastic model

Let us consider the vector \( C = [x, y, z, w]^T \) for the system (1)-(4). Transition probabilities are presented in Table 1 for the expectations \( E^*[\Delta C] \) and variance \( E^*[\Delta C \Delta C^T] \).

**Table 1** Possible changes in the process of the model.

| Transition | Probabilities |
|------------|---------------|
| \( (\Delta C)_1 = [1 \ 0 \ 0 \ 0]^T \) | \( P_1 = \rho \Delta t \) |
| \( (\Delta C)_2 = [-1 \ 1 \ 0 \ 0]^T \) | \( P_2 = (\beta_m x \theta + \beta_p x \theta) \Delta t \) |
| \( (\Delta C)_3 = [-1 \ 0 \ 0 \ 0]^T \) | \( P_3 = \mu x \Delta t \) |
| \( (\Delta C)_4 = [0 \ -1 \ 1 \ 0]^T \) | \( P_4 = \theta y \Delta t \) |
| \( (\Delta C)_5 = [0 \ -1 \ 0 \ 0]^T \) | \( P_5 = \mu y \Delta t \) |
| \( (\Delta C)_6 = [0 \ 0 \ 1 \ -1]^T \) | \( P_6 = f \theta y \Delta t \) |
| \( (\Delta C)_7 = [0 \ 0 \ -1 \ 0]^T \) | \( P_7 = \mu z \Delta t \) |
| \( (\Delta C)_8 = [0 \ 0 \ 0 \ -1]^T \) | \( P_8 = \mu w \Delta t \) |

We have the following quantities:

(i) Expectation = \( E^*[\Delta C] = \sum_{i=1}^{8} P_i (\Delta c)_i = \left[ \begin{array}{c} \rho - (\beta_m x \theta + \beta_p x \theta) - \mu x \\ \beta_m x \theta + \beta_p x \theta - \theta y - \mu y \\ f \theta y - \mu z \\ \theta y - f \theta y - \mu w \end{array} \right] \Delta t; \)

(ii) Variance = \( E^*[\Delta C \Delta C^T] = \sum_{i=1}^{8} P_i [(\Delta C)_i][[(\Delta C)_i]^T = \\
\left[ \begin{array}{cccccccc} P_1 + P_2 + P_3 & -P_2 & 0 & 0 \\
-P_2 & P_2 + P_4 + P_5 & 0 & -P_4 \\
0 & 0 & P_6 + P_7 & -P_6 \\
0 & -P_4 & -P_6 & P_4 + P_6 + P_8 \end{array} \right] \Delta t = \\
\left[ \begin{array}{cccccccc} W_{11} & W_{12} & W_{13} & W_{14} \\
W_{21} & W_{22} & W_{23} & W_{24} \\
W_{31} & W_{32} & W_{33} & W_{34} \\
W_{41} & W_{42} & W_{43} & W_{44} \end{array} \right] \Delta t, \)
where:

\[
W_{11} = \rho + \left( \beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} \right) + \mu x, \quad W_{12} = - \left( \beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} \right),
\]

\[
W_{13} = 0, \quad W_{14} = 0,
\]

\[
W_{21} = - \left( \beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} \right), \quad W_{22} = \beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} + \theta y + \mu y,
\]

\[
W_{23} = 0, \quad W_{24} = - \theta y,
\]

\[
W_{31} = 0, \quad W_{32} = 0, \quad W_{33} = f \theta y + \mu z, \quad W_{34} = - f \theta y,
\]

\[
W_{41} = 0, \quad W_{42} = - \theta y, \quad W_{43} = - f \theta y, \quad W_{44} = \theta y + f \theta y + \mu w.
\]

The following additional quantities are defined:

\[
\text{Drift} = G(C, t) = \frac{E^* [\Delta C]}{\Delta t} = \begin{bmatrix}
\rho - \left( \beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} \right) - \mu x \\
\beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} - \theta y - \mu y \\
f \theta y - \mu z \\
\theta y - f \theta y - \mu w
\end{bmatrix}; \quad (10)
\]

\[
\text{Diffusion} = H(C, t) = \sqrt{E^* [\Delta C \Delta C^T]} = \begin{bmatrix}
W_{11} & W_{12} & W_{13} & W_{14} \\
W_{21} & W_{22} & W_{23} & W_{24} \\
W_{31} & W_{32} & W_{33} & W_{34} \\
W_{41} & W_{42} & W_{43} & W_{44}
\end{bmatrix}. \quad (11)
\]

Finally, the stochastic differential equation based on (10) and (11) takes the following form

\[
dC(t) = G(C, t) dt + H(C, t) dB(t)
\]

or equivalently

\[
d\begin{bmatrix}
x \\
y \\
z \\
w
\end{bmatrix} = \begin{bmatrix}
\rho - \left( \beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} \right) - \mu x \\
\beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} - \theta y - \mu y \\
f \theta y - \mu z \\
\theta y - f \theta y - \mu w
\end{bmatrix} dt +
\begin{bmatrix}
\rho \left( \beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} \right) + \mu x - \left( \beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} \right) \\
- \left( \beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} \right) \beta_m x \frac{z}{N} + \beta_p x \frac{w}{N} + \theta y + \mu y \\
0 - \theta y \\
0 - f \theta y - \theta y + f \theta y + \mu w
\end{bmatrix} dB(t) \quad (12)
\]
2.5 Euler Maruyama scheme

In this section, we utilize the Euler Maruyama scheme to determine the numerical solution of the stochastic differential equation (12). The employed model parameters are reported in Table 2. The following computational procedure holds:

\[
C_{n+1} = C_n + f(C_n, t) \Delta t + L(C_n, t) dB,
\]

\[
\begin{bmatrix}
  x^{n+1} \\
  y^{n+1} \\
  z^{n+1} \\
  w^{n+1}
\end{bmatrix}
= \begin{bmatrix}
  x^n \\
  y^n \\
  z^n \\
  w^n
\end{bmatrix} + \begin{bmatrix}
  \rho - (\beta_m x^n \frac{z^n}{N} + \beta_p x^n \frac{w^n}{N}) - \mu x^n \\
  \beta_m x^n \frac{z^n}{N} + \beta_p x^n \frac{w^n}{N} - \theta y^n - \mu y^n \\
  f \theta y^n - \mu z^n \\
  \theta y^n - f \theta y^n - \mu w^n
\end{bmatrix} \Delta t + \sqrt{\begin{bmatrix}
  \rho + (\beta_m x^n \frac{z^n}{N} + \beta_p x^n \frac{w^n}{N}) + \mu x^n \\
  - (\beta_m x^n \frac{z^n}{N} + \beta_p x^n \frac{w^n}{N}) \\
  0 \\
  0
\end{bmatrix} - \begin{bmatrix}
  \beta_m x^n \frac{z^n}{N} + \beta_p x^n \frac{w^n}{N} + \theta y^n + \mu y^n \\
  0 \\
  -\theta y^n \\
  -f \theta y^n + \theta y^n + f \theta y^n + \mu w^n
\end{bmatrix} \Delta B_n,
\]

where \( \Delta t \) stands for the discretization parameter.

Table 2 Value of parameters.

| Parameters | Values       |
|------------|--------------|
| \( f \)    | 0.7          |
| \( \mu \)  | 0.5          |
| \( \theta \) | 0.19         |
| \( \kappa \) | 0.5         |
| \( \beta_p \) | 2 (DFE)  |
| \( \beta_m \) | 2.3 (EE) |
| \( \beta_p \) | 2 (DFE)  |

2.6 Parametric perturbation of the model

In what follows, we introduce parametric perturbation in the system (1)-(4) as follows

\[
\beta_m dt = \beta_m dt + \sigma_1 dB(t), \quad \beta_p dt = \beta_p dt + \sigma_2 dB(t),
\]

and we get

\[
dx(t) = \left(\rho - \frac{\beta_m x(t) z(t)}{N} - \frac{\beta_p x(t) w(t)}{N} - \mu x(t)\right) dt - \frac{\sigma_1 \beta_m x(t) z(t)}{N} dB(t) - \frac{\sigma_2 \beta_p x(t) w(t)}{N} dB(t), \quad t \geq 0
\]

(13)
\[ dy(t) = \left( \frac{\beta_m x(t) z(t)}{N} + \frac{\beta_p x(t) w(t)}{N} - \theta y(t) - \mu y(t) \right) dt + \frac{\sigma_1 \beta_m x(t) z(t)}{N} dB(t) + \frac{\sigma_2 \beta_p x(t) w(t)}{N} dB(t), \quad t \geq 0 \]  
(14)

\[ dz(t) = (f \theta y(t) - \mu z(t)) dt, \quad t \geq 0, \]  
(15)

\[ dw(t) = (\theta (1 - f) y(t) - \mu w(t)) dt, \quad t \geq 0 \]  
(16)

where \( \sigma_i, i = 1, 2 \) are the randomness of the model and \( B(t) \) is the Brownian motion.

2.7 Positivity and boundedness of the stochastic model

Consider a complete probability space which is denoted by \((\Omega, F, P)\) and with filtration represented by \(\{F_t\}_{t \in \mathbb{R}}\). The initial set \(F_0\) contains all \(F\) null sets satisfying the conditions – right continuous and increasing. Let \(U(t) = (x(t), y(t), z(t), w(t))\). Its norm follows

\[ |U(t)| = \sqrt{x^2(t) + y^2(t) + z^2(t) + w^2(t)}. \]  
(17)

In addition, let \(C^{2,1}_r (\mathbb{R}_+^4 : \mathbb{R}_+)\) be the families of all non-negative function \(V(U, t)\) defined on \(\mathbb{R}_+^4 \times (0, \infty)\), being twice differentiable in \(U\) and once in "\(t\".

We define the differential operator \(L\) associated with four-dimensional SDEs of the form

\[ dU(t) = H_1(U, t) + K_1(U, t) dB(t). \]  
(18)

We have

\[ L = \frac{\partial}{\partial t} + \sum_{i=1}^{3} H_{1i}(U, t) \frac{\partial}{\partial U_i} + \frac{1}{2} \sum_{i,j=1}^{3} \left( K_{1i}^T(U, t) K_1(U, t)_{i,j} \times \frac{\partial^2}{\partial U_i \partial U_j} \right). \]

The action of the operator \(L\) on a function \(U^* \in C^{2,1}_r (\mathbb{R}_+^4 : \mathbb{R}_+)\) yields

\[ LU^*(U, t) = U^*_{t} (U, t) + U^*_U (U, t) H_1(U, t) + \frac{1}{2} \text{Trace} \left( K_1^T(U, t) U^*_{UU}(U, t) K_1(U, t) \right), \]

where \(T\) means transportations.

**Theorem 5** For system (13)-(16) and any given initial conditions \((x(0), y(0), z(0), w(0)) \in \mathbb{R}_+^4\), there is a unique solution \((x(t), y(t), z(t), w(t)), t \geq 0\) which remain \(\mathbb{R}_+^4\) with probability one.
Proof: By Itô’s formula, (13)-(16) admits a positive solution in the sense of unique local on \([0, \tau_e]\) where explosion time is denoted \(\tau_e\) due to the local Lipschitz coefficients of the model.

Next, we shall prove that the equations (13)-(16) assume this solution in a global sense – meaning that \(\tau_e = \infty\). Let \(m_0 = 0\) be sufficiently large for \(x(0), y(0), z(0)\) and \(w(0)\) lying with the interval \(\left\{\frac{1}{m_0}, m_0\right\}\).

For each integer \(m \geq 0\), define a sequence that is so-called stopping time as
\[
\tau_m = \inf \left\{ \tau \in [0, \tau_e]: x(t) \left(\frac{1}{m}, m\right) \text{ or } y(t) \left(\frac{1}{m}, m\right) \text{ or } z(t) \left(\frac{1}{m}, m\right) \text{ or } w(t) \left(\frac{1}{m}, m\right) \right\},
\]
where we set \(\varphi = \infty\) (\(\varphi\) is an empty set).

Since \(\tau_m\) is non-decreasing as \(m \to \infty\), we get
\[
\tau_\infty = \lim_{m \to \infty} \tau_m,
\]
which implies \(\tau_\infty \leq \tau_e\). Now, we wish to show \(\tau_\infty = \infty\), as desired. If this statement is violated, then there exist \(\tau > 0\) and \(a_1 \in (0, 1)\) such that
\[
P(\tau_\infty \leq T) > a_1, \quad \forall m \geq m_1,
\]
\[
P(\tau_m \leq T) > a_1, \quad \forall m \geq m_1.
\]

Let us define a function \(f : \mathbb{R}_+^4 \to \mathbb{R}_+\) as
\[
f(x, y, z, w) = (x - 1 - \ln x) + (y - 1 - \ln y) + (z - 1 - \ln z) + (w - 1 - \ln w).
\]

By applying Itô’s formula on (23), we have
\[
df(x, y, z, w) = \left(1 - \frac{1}{x}\right) dx + \left(1 - \frac{1}{y}\right) dy + \left(1 - \frac{1}{z}\right) dz + \left(1 - \frac{1}{w}\right) dw + \frac{\sigma_1^2 + \sigma_2^2}{2} dt,
\]
\[
df(x, y, z, w) = \left(1 - \frac{1}{x}\right) \left[\rho - \beta_{mxz} x - \frac{\beta_{pxw} x}{N} - \mu x\right] dt - \frac{\sigma_1 \beta_{mxz}}{N} dB(t) - \frac{\sigma_2 \beta_{pxw}}{N} dB(t) + \left(1 - \frac{1}{y}\right) \left[\frac{\beta_{mxz}}{N} + \frac{\beta_{pxw}}{N} - \theta y - \mu y\right] + \left(1 - \frac{1}{z}\right) \left[f \theta y - \mu z\right] + \left(1 - \frac{1}{w}\right) \left[\theta (1 - f) y - \mu w\right]
\]
\[ df(x, y, z, w) \leq \left[ \rho + \theta + \mu + \frac{\sigma_1^2 + \sigma_2^2}{2} \right] dt + \frac{\sigma_1 \beta_{m,xz}}{N} dB(t) - \frac{\sigma_2 \beta_{p,xw}}{N} dB(t). \tag{24} \]

For simplicity, let \( N_1 = \rho + \theta + \mu + \frac{\sigma_1^2 + \sigma_2^2}{2} \). Then equation (24) will be written as
\[ df(x, y, z, w) \leq N_1 dt + \left[ \frac{\sigma_1 \beta_{m,xz}}{N} + \frac{\sigma_2 \beta_{p,xw}}{N} \right] dB(t), \tag{25} \]
where \( N_1 \) stands for a positive constant.

Integration of equation (25) yields
\[ \int_0^{\tau_m A} df(x, y, z, w) \leq N_1 \int_0^{\tau_m A} ds + \int_0^{\tau_m A} \left[ \frac{\sigma_1 \beta_{m,xz}}{N} + \frac{\sigma_2 \beta_{p,xw}}{N} \right] dB(s), \tag{26} \]
where \( \tau_m A = \min(\tau_m, T) \). Therefore, we assume
\[ EU^*(x(\tau_m A), y(\tau_m A), z(\tau_m A), w(\tau_m A)) \leq U^*(x(0), y(0), z(0), w(0)) + N_1 T. \tag{27} \]

Setting \( \chi_m = \{ \tau_m \leq T \} \) for \( m > m_1 \) and taking into account equation (20), we have \( P(\chi_m \geq a_1) \).

For every \( \chi_1 \in \chi_m \) there are some \( I \) such that \( U_i(\tau_m, \chi_1) \) equals either \( m \) or \( \frac{1}{m} \) for \( i = 1, 2, 3, 4 \). Hence, we have
\[ U = U^*(x(\tau_m, \chi_1), y(\tau_m, \chi_1), z(\tau_m, \chi_1), w(\tau_m, \chi_1)). \]

If it is less than \( \min(m - 1 - \ln m, \frac{1}{m} - 1 - \ln \frac{1}{m}) \), then we obtain
\[ U^*(x(0), y(0), z(0), w(0)) + N_1 T \geq E(I_{\chi_m} U^*(x(\tau_m), y(\tau_m)), z(\tau_m), w(\tau_m)) \geq \left\{ \min \left\{ m - 1 - \ln m, \frac{1}{m} - 1 - \ln \frac{1}{m} \right\} \right\} \tag{28} \]

The indicator function is represented by \( I_{\chi} \) of \( \chi_m \). Letting \( m \to \infty \) leads to the contradiction \( \infty = U^*(x(0), y(0), z(0), w(0)) + N_1 T < \infty \), as desired.

### 3 Computational methods

This section discusses well-known methods like stochastic Euler, stochastic Runge Kutta, and stochastic non-standard finite difference method with non-negative initial conditions.
3.1 Stochastic Euler

This approach (also known as the forward Euler method) is a first-order numerical approximation for solving stochastic differential equations (SDEs). The discretization of the system (13)-(16) under the rules of this scheme is as follows:

\[
\frac{x^{n+1} - x^n}{h} = \rho \frac{x^n}{N} - \beta \frac{z^n}{N} - \mu x^n - \frac{\sigma_1 \beta m x^n z^n}{N} \Delta B_n - \frac{\sigma_2 \beta p x^n w^n}{N} \Delta B_n,
\]

\[
x^{n+1} = x^n + h \left[ \rho \frac{x^n}{N} - \beta \frac{z^n}{N} - \mu x^n - \frac{\sigma_1 \beta m x^n z^n}{N} \Delta B_n - \frac{\sigma_2 \beta p x^n w^n}{N} \Delta B_n \right], \tag{29}
\]

\[
\frac{y^{n+1} - y^n}{h} = \frac{\beta m x^n z^n}{N} + \frac{\beta p x^n w^n}{N} - \theta y^n - \mu y^n + \frac{\sigma_1 \beta m x^n z^n}{N} \Delta B_n + \frac{\sigma_2 \beta p x^n w^n}{N} \Delta B_n,
\]

\[
y^{n+1} = y^n + h \left[ \frac{\beta m x^n z^n}{N} + \frac{\beta p x^n w^n}{N} - \theta y^n - \mu y^n + \frac{\sigma_1 \beta m x^n z^n}{N} \Delta B_n + \frac{\sigma_2 \beta p x^n w^n}{N} \Delta B_n \right], \tag{30}
\]

\[
\frac{z^{n+1} - z^n}{h} = (f \theta y^n - \mu z^n), \quad z^{n+1} = z^n + h (f \theta y^n - \mu z^n), \tag{31}
\]

\[
\frac{w^{n+1} - w^n}{h} = \theta (1 - f) y^n - \mu w^n,
\]

\[
w^{n+1} = w^n + h (\theta (1 - f) y^n - \mu w^n), \tag{32}
\]

where \( h \) is any discretization parameter and \( n \geq 0 \). Using the values of the parameters as presented in Table 2, the graphs of stochastic Euler’s scheme for disease-free equilibrium (DFE) and endemic equilibrium (EE) plotted with MATLAB software are shown in Figures 2-5.
Fig. 2 Combined behavior of sub-populations at DFE for $h = 0.1$.

Fig. 3 Combined behavior of sub-populations at DFE for $h = 1$.

Fig. 4 Combined behavior of sub-populations at EE for $h = 0.1$. 
3.2 Stochastic Runge-Kutta method

This technique is also known as Runge Kutta of order 4th method. In numerical analysis, it is of the family of an implicit and explicit iterative process. In 1900 two German mathematicians Carl Runge and Wilhelm Kutta, developed this method. The stochastic Runge-Kutta scheme for equations (13)-(16) can be established according to the following steps.

**First Step**

\[ K_1 = h \left[ \rho - \frac{\beta_m x^n z^n}{N} - \frac{\beta_p x^n w^n}{N} - \mu x^n - \frac{\sigma_1 \beta_m x^n z^n}{N} \Delta B_n - \frac{\sigma_2 \beta_p x^n w^n}{N} \Delta B_n \right] \]

\[ L_1 = h \left[ \frac{\beta_m x^n z^n}{N} + \frac{\beta_p x^n w^n}{N} - \theta y^n - \mu y^n + \frac{\sigma_1 \beta_m x^n z^n}{N} \Delta B_n + \frac{\sigma_2 \beta_p x^n w^n}{N} \Delta B_n \right] , \]

\[ M_1 = h [f \theta y^n - \mu z^n] , \quad N_1 = h [\theta (1 - f) y^n - \mu w^n] . \]

**Second Step**

\[ K_2 = h \left[ \rho - \frac{\beta_m (x^n + K_1^2)}{N} \left( z^n + \frac{M_1}{2} \right) - \frac{\beta_p (x^n + K_1^2)}{N} \left( w^n + \frac{N_1}{2} \right) - \right] \]

\[ \mu \left( x^n + \frac{K_1}{2} \right) - \frac{\sigma_1 \beta_m (x^n + K_1^2)}{N} \left( z^n + \frac{M_1}{2} \right) \Delta B_n - \]

Fig. 5 Combined behavior of sub-populations at EE for \( h = 1 \).
\( L_2 = h \left[ \frac{\beta_m (x^n + \frac{K_1}{2})}{N} \left( z^n + \frac{M_1}{2} \right) + \frac{\beta_p (x^n + \frac{K_1}{2})}{N} \left( w^n + \frac{N_1}{2} \right) \right] \),

\( \theta \left( y^n + \frac{L_1}{2} \right) - \mu \left( y^n + \frac{L_1}{2} \right) + \frac{\sigma_1 \beta_m (x^n + \frac{K_2}{2})}{N} \left( z^n + \frac{M_2}{2} \right) \Delta B_n + \frac{\sigma_2 \beta_p (x^n + \frac{K_2}{2})}{N} \left( w^n + \frac{N_2}{2} \right) \Delta B_n \),

\( M_2 = h \left[ f \theta \left( y^n + \frac{L_1}{2} \right) - \mu \left( z^n + \frac{M_1}{2} \right) \right] \),

\( N_2 = h \left[ \theta \left( 1 - f \right) \left( y^n + \frac{L_1}{2} \right) - \mu \left( w^n + \frac{N_1}{2} \right) \right] \),

**Third Step**

\( K_3 = h \left[ \rho - \beta_m \left( x^n + \frac{K_2}{2} \right) - \beta_p \left( x^n + \frac{K_2}{2} \right) \right] \),

\( \mu \left( x^n + \frac{K_2}{2} \right) - \frac{\sigma_1 \beta_m (x^n + \frac{K_2}{2})}{N} \left( z^n + \frac{M_2}{2} \right) \Delta B_n - \frac{\sigma_2 \beta_p (x^n + \frac{K_2}{2})}{N} \left( w^n + \frac{N_2}{2} \right) \Delta B_n \),

\( L_3 = h \left[ \beta_m \left( x^n + \frac{K_2}{2} \right) \left( z^n + \frac{M_2}{2} \right) + \beta_p \left( x^n + \frac{K_2}{2} \right) \left( w^n + \frac{N_2}{2} \right) \right] \),

\( \theta \left( y^n + \frac{L_2}{2} \right) - \mu \left( y^n + \frac{L_2}{2} \right) + \frac{\sigma_1 \beta_m (x^n + \frac{K_2}{2})}{N} \left( z^n + \frac{M_2}{2} \right) \Delta B_n + \frac{\sigma_2 \beta_p (x^n + \frac{K_2}{2})}{N} \left( w^n + \frac{N_2}{2} \right) \Delta B_n \),

\( M_3 = h \left[ f \theta \left( y^n + \frac{L_2}{2} \right) - \mu \left( z^n + \frac{M_2}{2} \right) \right] \),

\( N_3 = h \left[ \theta \left( 1 - f \right) \left( y^n + \frac{L_2}{2} \right) - \mu \left( w^n + \frac{N_2}{2} \right) \right] \),

**Fourth Step**
\[ K_4 = h \left[ \rho - \frac{\beta_m (x^n + K_3) (z^n + M_3)}{N} - \frac{\beta_p (x^n + K_3) (w^n + N_3)}{N} - \frac{\mu (x^n + K_3)}{N} + \frac{\mu (y^n + L_3)}{N} + \frac{\sigma_1 \beta_m (x^n + K_3) (z^n + M_3)}{N} \Delta B_n - \frac{\sigma_2 \beta_p (x^n + K_3) (w^n + N_3)}{N} \Delta B_n \right], \]

\[ L_4 = h \left[ \frac{\beta_m (x^n + K_3) (z^n + M_3)}{N} + \frac{\beta_p (x^n + K_3) (w^n + N_3)}{N} - \theta (y^n + L_3) - \frac{\mu (y^n + L_3)}{N} - \frac{\sigma_1 \beta_m (x^n + K_3) (z^n + M_3)}{N} \Delta B_n + \frac{\sigma_2 \beta_p (x^n + K_3) (w^n + N_3)}{N} \Delta B_n \right], \]

\[ M_4 = h[f \theta(y^n + L_3) - \mu(z^n + M_3)], \]

\[ N_4 = h[f \theta(1 - f)(y^n + L_3) - \mu(w^n + N_3)]. \]

**Final Step**

\[
\begin{align*}
  x^{n+1} &= x^n + \frac{1}{6} [K_1 + 2K_2 + 2K_3 + K_4] \\
  y^{n+1} &= y^n + \frac{1}{6} [L_1 + 2L_2 + 2L_3 + L_4] \\
  z^{n+1} &= z^n + \frac{1}{6} [M_1 + 2M_2 + 2M_3 + M_4] \\
  w^{n+1} &= w^n + \frac{1}{6} [N_1 + 2N_2 + 2N_3 + N_4]
\end{align*}
\]

(33)

where \( h \) is any discretization parameter and \( n \geq 0 \). The graphs of the stochastic Runge Kutta scheme for disease-free equilibrium (DFE) and endemic equilibrium (EE) plotted with MATLAB software are presented in Figures 6-9 for the parameters from Table 2.
Fig. 6 Combined behavior of sub-populations at DFE for \( h = 0.1 \).

Fig. 7 Combined behavior of sub-populations at DFE for \( h = 2 \).

Fig. 8 Combined behavior of sub-populations at EE for \( h = 0.1 \).
3.3 Stochastic NSFD

The NSFD method is also known as the implicitly driven explicit method. From equations (13)-(16), we established the computational scheme in the following manner.

Equation (13) yields

\[ x^{n+1} = x^n + h \left[ \rho - \frac{\beta_m x^{n+1} z^n}{N} - \frac{\beta_p x^{n+1} w^n}{N} - \mu x^{n+1} - \frac{\sigma_1 \beta_m x^{n+1} z^n}{N} \Delta B_n - \frac{\sigma_2 \beta_p x^{n+1} w^n}{N} \Delta B_n \right], \]

\[ x^{n+1} = x^n + h \left[ \frac{\sigma_1 \beta_m x^{n+1} z^n}{N} \Delta B_n - \frac{\sigma_2 \beta_p x^{n+1} w^n}{N} \Delta B_n \right], \tag{34} \]

Equation (14) gives

\[ y^{n+1} = y^n + h \left[ \beta_m x^n z^n + \beta_p x^n w^n - \theta y^{n+1} - \mu y^{n+1} + \frac{\sigma_1 \beta_m x^n z^n}{N} \Delta B_n + \frac{\sigma_2 \beta_p x^n w^n}{N} \Delta B_n \right], \]

\[ y^{n+1} = y^n + h \left[ \frac{\beta_m x^n z^n}{N} + \frac{\beta_p x^n w^n}{N} + \frac{\sigma_1 \beta_m x^n z^n}{N} \Delta B_n + \frac{\sigma_2 \beta_p x^n w^n}{N} \Delta B_n \right], \tag{35} \]

Equation (15) leads to

\[ z^{n+1} = z^n + h \left( f \theta y^n - \mu z^{n+1} \right), \]

\[ z^{n+1} = \frac{z^n + h f \theta y^n}{1 + h \mu}. \tag{36} \]
Equation (16) produces

\[ w^{n+1} = w^n + h \left( \theta (1 - f) y^n - \mu w^{n+1} \right) \]

\[ w^{n+1} = \frac{w^n + h\theta (1 - f) y^n}{1 + h\mu}, \]  \hspace{1cm} (37)

where \( h \) is any discretization parameter and \( n \geq 0 \).

3.3.1 Stability analysis

**Theorem 6** Theorem: For any \( n \geq 0 \), the proposed SNSFD method is stable if the eigenvalues of the system (33)-(37) lie in the unit circle for \( R_0 < 1 \).

**Proof:** Consider the right-hand sides of the system of equations (33)-(37) as functions \( A, B, C \) and \( D \) with the assumption that \( \Delta B_n = 0 \), as follows:

\[ A = \frac{x + h\rho}{1 + h\left( \frac{\beta_{\mu}^z}{N} + \frac{\beta_{\mu}^w}{N} + \mu \right)}, \]

\[ B = \frac{y + h \left( \frac{\beta_{\mu}^x z}{N} + \frac{\beta_{\mu}^x w}{N} \right)}{1 + h\theta + h\mu}, \]

\[ C = \frac{z + hf\theta y}{1 + h\mu}, \]

\[ D = \frac{w + h\theta y (1 - f)}{1 + h\mu}. \]

The elements of the Jacobian matrix are

\[ \frac{\partial A}{\partial x} = \frac{1}{1 + h\mu}, \quad \frac{\partial A}{\partial y} = 0, \quad \frac{\partial A}{\partial z} = -\frac{(x + h\rho) \left( \frac{h\beta_{\mu}^x}{N} \right)}{\left( 1 + \frac{h\beta_{\mu}^z}{N} + \frac{h\beta_{\mu}^w}{N} + h\mu \right)^2}, \]

\[ \frac{\partial B}{\partial x} = \frac{\frac{h\beta_{\mu}^z}{N} + \frac{h\beta_{\mu}^w}{N}}{1 + h\theta + h\mu}, \quad \frac{\partial B}{\partial y} = \frac{1}{1 + h\theta + h\mu}, \]

\[ \frac{\partial B}{\partial z} = \frac{\frac{h\beta_{\mu}^x x}{N}}{1 + h\theta + h\mu}, \quad \frac{\partial B}{\partial w} = \frac{\frac{h\beta_{\mu}^x}{N}}{1 + h\theta + h\mu}, \]

\[ \frac{\partial C}{\partial x} = 0, \quad \frac{\partial C}{\partial y} = \frac{hf\theta}{1 + h\mu}, \quad \frac{\partial C}{\partial z} = \frac{1}{1 + h\mu}, \quad \frac{\partial C}{\partial w} = 0, \]

\[ \frac{\partial D}{\partial x} = 0, \quad \frac{\partial D}{\partial y} = \frac{h\theta (1 - f)}{1 + h\mu}, \quad \frac{\partial D}{\partial z} = 0, \quad \frac{\partial D}{\partial w} = \frac{1}{1 + h\mu}. \]
and we have

\[
J = \begin{bmatrix}
\frac{1}{1+h\mu} & 0 & -\frac{(x+h\rho)(\frac{\mu}{\beta \mu})}{(1+h\theta+h\mu)^2} & -\frac{(x+h\rho)(\frac{\lambda}{\beta \mu})}{(1+h\theta+h\mu)^2} \\
\frac{h_f \theta}{1+h\mu} & \frac{1}{1+h\mu} & \frac{h_f \theta}{1+h\mu} & \frac{h_f \theta}{1+h\mu} \\
0 & \frac{h_f \theta (1-f)}{1+h\mu} & 0 & \frac{1}{1+h\mu} \\
0 & \frac{h_f \theta (1-f)}{1+h\mu} & 0 & \frac{1}{1+h\mu}
\end{bmatrix}.
\]

At the disease-free equilibrium (DFE- \( L_1 \)) = \( \left( \frac{\mu}{\beta}, 0, 0, 0 \right) \), we get

\[
J \left( \frac{\mu}{\beta}, 0, 0, 0 \right) = \begin{bmatrix}
\frac{1}{1+h\mu} & 0 & -\frac{(\frac{\mu}{\beta}+h\rho)(\frac{h_{\beta \mu} \rho}{(1+h\mu)^2})}{N(1+h\theta+h\mu)} & -\frac{(\frac{\mu}{\beta}+h\rho)(\frac{h_{\beta \mu} \rho}{(1+h\mu)^2})}{N(1+h\theta+h\mu)} \\
\frac{1}{1+h\theta+h\mu} & \frac{h_{\beta \mu} \rho}{N(1+h\theta+h\mu)} & \frac{h_{\beta \mu} \rho}{N(1+h\theta+h\mu)} & \frac{h_{\beta \mu} \rho}{N(1+h\theta+h\mu)} \\
0 & \frac{h_f \theta}{1+h\mu} & \frac{1}{1+h\mu} & 0 \\
0 & \frac{h_f \theta (1-f)}{1+h\mu} & 0 & \frac{1}{1+h\mu} - \lambda
\end{bmatrix}
\]

Hence

\[
\begin{vmatrix}
\frac{1}{1+h\mu} & 0 & -\frac{(\frac{\mu}{\beta}+h\rho)(\frac{h_{\beta \mu} \rho}{(1+h\mu)^2})}{N(1+h\theta+h\mu)} & -\frac{(\frac{\mu}{\beta}+h\rho)(\frac{h_{\beta \mu} \rho}{(1+h\mu)^2})}{N(1+h\theta+h\mu)} \\
\frac{1}{1+h\theta+h\mu} & \frac{h_{\beta \mu} \rho}{\mu N(1+h\theta+h\mu)} & \frac{h_{\beta \mu} \rho}{\mu N(1+h\theta+h\mu)} & \frac{h_{\beta \mu} \rho}{\mu N(1+h\theta+h\mu)} \\
0 & \frac{h_f \theta}{1+h\mu} & \frac{1}{1+h\mu} - \lambda & 0 \\
0 & \frac{h_f \theta (1-f)}{1+h\mu} & 0 & \frac{1}{1+h\mu} - \lambda
\end{vmatrix} = 0,
\]

and \( \lambda_1 = \frac{1}{1+h\mu} \) < 1.

We also have

\[
\begin{vmatrix}
f_1 - \lambda & f_2 & f_3 & 0 \\
f_4 & f_5 - \lambda & 0 & f_3 \\
f_6 & 0 & f_5 - \lambda & 0 \\
\end{vmatrix} = 0,
\]

where \( f_1 = \frac{1}{1+h\theta+h\mu}, f_2 = \frac{h_{\beta \mu} \rho}{\mu N(1+h\theta+h\mu)}, f_3 = \frac{h_{\beta \mu} \rho}{\mu N(1+h\theta+h\mu)}, f_4 = \frac{h_f \theta}{1+h\mu}, f_5 = \frac{h_f \theta (1-f)}{1+h\mu}, f_6 = \frac{h_f \theta (1-f)}{1+h\mu} \).

The polynomial form of the characteristic equation reads

\[
\lambda^3 - \lambda^2 (f_1 + 2f_5) - \lambda (f_3f_6 - 2f_1f_5 - f_5^2 + f_2f_4) - (f_1f_5^2 - f_3f_5f_6 - f_2f_4f_5) = 0.
\]
Hence, using the Mathematica software, all the eigenvalues of the above Jacobian matrix were found to lie within the unit circle for $R_0 < 1$. Thus, the system (33)-(37) is stable.

3.4 Results comparison

Here, a comparison of the stochastic NSFD method with other stochastic numerical methods was carried out. We can easily conclude that some of the examined stochastic numerical approaches conditionally converge or diverge with larger time step values by looking at the numerical experimentations. However, the NSFD method remains convergent for all time step sizes.

**Fig. 10** Infected class by paucibacillary leprosy at EE for $h = 0.1$.

**Fig. 11** Infected class by paucibacillary leprosy at EE for $h = 1$. 
4 Concluding remarks

Stochastic modelling is a reliable and efficient technique to handle highly non-linear problems close to nature, and it may be understood as the extension of deterministic modelling. In this paper, we applied the parametric perturbation technique to the formulated model. We focused on the NSFD scheme that is dynamically consistent and positively bounded. Contrary to the assertions in some literature that most epidemiologists used Euler Maruyama, stochastic Euler, and stochastic Runge Kutta algorithms, we have demonstrated that the novelty of the stochastic NSFD scheme is also satisfactory. We have shown that the other techniques are conditionally convergent and even fail in the long run. However, the investigated system unveils the NSFD method in a stochastic form as always being convergent. Furthermore, in the future, we
will extend this idea to other types of modelling, including spatial-temporal, fractional, fractal fractional and delay problems of dynamical systems.

Conflict of interest

There is no challenging interest in this article.

Availability of data and materials

All of the necessary data and the implementation details have been included in the manuscript.

Authors’ contributions

The authors have equal contributions.

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