Research Article

Ergodic Stationary Distribution of a Stochastic Hepatitis B Epidemic Model with Interval-Valued Parameters and Compensated Poisson Process

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Throughout the document, the Hepatitis B epidemic is studied, with a focus on developing mathematical models to understand its dynamics. The paper introduces a novel approach to modeling the epidemic, considering interval-valued parameters and stochastic processes. This approach allows for a more realistic representation of the disease's transmission dynamics, taking into account uncertainties and environmental factors.

1. Introduction

Hepatitis B is an enormous challenge and a significant global health issue caused by the Hepatitis B virus (HBV) [2]. Chronic HBV can be transmitted through sexual contact, through the touch, by impregnation with polluted blood, or by the direct transmission of Hepatitis B from the mother to a fetus during pregnancy (vertical transmission) [3]. According to the recent statistics of the World Health Organization (WHO) [4], about 350 million people worldwide have been infected and carrying HBV. This serious infection is responsible for approximately 600,000 deaths each year [5]. Because of the high severity of HBV infection and the large number of deaths associated with it, it is compulsory to control and prevent the spread of this virus. Mathematical models are a powerful tool to simulate and control the spread of the HBV infection. There exist many previous interesting works committed to studying Hepatitis B transmission. For example, Anderson and May [6] analyzed a straightforward mathematical model for illustrating the role of carrier individuals on the spread of HBV. In [7, 8], the authors developed the impact of vaccination and other controlling measures of HBV outbreak. They showed that the booster vaccine of Hepatitis B is very necessary and useful. Khan et al. [9] formulated the characteristics of HBV disease transmission and proposed the following deterministic Susceptible (S)—Infected (I)—Recovered (R) model:

\[
\begin{align*}
\dot{S}(t) &= A - \beta S(t)I(t) - (\mu + \theta)S(t), \\
\dot{I}(t) &= \beta S(t)I(t) - (\mu + \delta)I(t), \\
\dot{R}(t) &= \delta I(t) + \theta S(t) - \mu R(t),
\end{align*}
\]

with initial data \( S(0) = S_0 > 0, I(0) = I_0 > 0, \) and \( R(0) = R_0 > 0. \) The positive parameters of the deterministic model (1) are given in the following list. The deterministic model constructed above can be improved by taking into account the unpredictable biological conditions [10–14]. Also, environmental fluctuations have important effects on the growth and propagation of an epidemic disease [15, 16].
Khan et al. [9] discussed the dynamics of a stochastic Hepatitis B epidemic model with varying population size. They supposed that the effect of the random fluctuations is manifested as a perturbation in the Hepatitis B transmission rate. To confer the realistic aspect to our study and make it biologically reasonable, in this study, we extend the work of Khan et al. [9] to the case of Lévy noise perturbation. We take into consideration the effects due to some unexpected and severe environmental disturbances (tsunami, floods, earthquakes, hurricanes, whirlwinds, etc.) on the disease outbreak [17, 18]. Thus, we consider the following model:

\[
\begin{align*}
\text{d}S(t) &= (A - \beta S(t)I(t) - (\mu + \delta)S(t))\text{d}t - \sigma S(t)I(t)\text{d}W(t) \\
&\quad - \int_{\mathbb{R}} \eta(u)S(t^-)I(t^-)\mathcal{N}(\text{d}t, \text{d}u), \\
\text{d}I(t) &= (\beta S(t)I(t) - (\mu + \delta + r)I(t))\text{d}t + \sigma S(t^-)I(t^-)\text{d}W(t) \\
&\quad + \int_{\mathbb{R}} \eta(u)S(t^-)I(t^-)\mathcal{N}(\text{d}t, \text{d}u), \\
\text{d}R(t) &= (\delta I(t) + \beta S(t) - \mu R(t))\text{d}t,
\end{align*}
\]

(2)

where \(S(t^-)\) and \(I(t^-)\) are the left limits of \(S(t)\) and \(I(t)\), respectively. \(W(t)\) is a real-valued Brownian motion with intensity \(\sigma > 0\) defined on a complete probability space \((\Omega, \mathcal{F}, \mathbb{P})\) with a filtration \(\{\mathcal{F}_t\}_{t \geq 0}\) satisfying the usual conditions. \(N\) is a Poisson counting measure with the compensator \(\mathcal{N}\) and the characteristic measure \(\nu\) on a measurable subset \(Z\) of \((0, \infty)\) satisfying \(\nu(Z) < \infty\). \(W(t)\) is independent of \(N\). We assume that \(\nu\) is a Lévy measure such that \(\mathcal{N}(\text{d}t, \text{d}u) = N(\text{d}t, \text{d}u) - \nu(\text{d}u)\text{d}t\). The bounded function \(\eta: Z \times \Omega \rightarrow \mathbb{R}\) is \(\mathcal{B}(Z) \times \mathcal{F}_t\)-measurable and continuous with respect to \(\nu\).

In system (2), we assume that model parameters (see Table 1) are precisely known and constant. However, this hypothesis may not be validated due to the lack of data and errors of measurements. It is more realistic to study Hepatitis B dynamics with interval-valued parameters. Recently, Pal et al. [19] used interval-valued parameters to analyze the prey-predator model due to the lack of precise biological data such as prey and predator population growth rates. The same logic was applied for epidemic models. In [20], the authors treated a cholera epidemic model with uncertain parameters. They investigated the stability condition of equilibrium points. Bao et al. [21] studied a stochastic SIRS model that includes Lévy jumps and interval parameters. They established the stochastic threshold which determines the extinction and persistence in the mean of disease. In [22], the authors studied an imprecise SIR epidemic model. They solved the optimal control problem.

In this paper, we consider the Hepatitis B epidemic model with stochastic transmissions and Lévy noise. To make our model more realistic, we consider imprecise biological parameters. To the best of our knowledge, the existence of a stationary distribution of system (2) with imprecise parameters remains not proved. In the next section, we propose a solution to the mentioned problem by considering an original method different from the Lyapunov approach described in [23]. Before proving the existence of a unique stationary distribution in Subsection 2.3, we demonstrate the well-posedness of the model (2) with interval-valued parameters in Subsection 2.2. Simulation examples are proposed in Subsection 2.4 to illustrate our theoretical study.

### 2. Main Results

#### 2.1. Imprecise Stochastic Hepatitis B Model

Before showing the main result of this paper, we first present some definitions of interval numbers and interval-valued functions which are used in our study. Then, we construct the imprecise stochastic Hepatitis B model.

**Definition 1** (see [19]). An interval number \(Z = [z, \bar{z}]\) is defined as \(Z = [z, \bar{z}] = \{x \mid z \leq x \leq \bar{z}, x \in \mathbb{R}\}\) where \(\mathbb{R}\) is the set of all real numbers and \(z\) and \(\bar{z}\) are the lower and upper limits of the interval numbers, respectively. Furthermore, any real number \(z\) can be represented in terms of interval number as \([z, \bar{z}]\).

**Definition 2** (see [19]). An interval-valued function for the interval \([x, y]\) can be represented by the following function:

\[
\psi(p) = x^{(1-p)y^p}, \quad \text{for } p \in [0, 1].
\]

**Theorem 1.** The following stochastic differential equation with interval-valued parameters

\[
\begin{align*}
\text{d}S(t) &= (\bar{A} - \bar{\beta}S(t)I(t) - (\bar{\mu} + \bar{\delta})S(t))\text{d}t \\
&\quad - \bar{\sigma}S(t^-)I(t^-)\text{d}W(t) - \int_{\mathbb{R}} \eta(u)S(t^-)I(t^-)\mathcal{N}(\text{d}t, \text{d}u), \\
\text{d}I(t) &= (\bar{\beta}S(t)I(t) - (\bar{\mu} + \bar{\delta} + \bar{r})I(t))\text{d}t \\
&\quad + \bar{\sigma}S(t^-)I(t^-)\text{d}W(t) + \int_{\mathbb{R}} \eta(u)S(t^-)I(t^-)\mathcal{N}(\text{d}t, \text{d}u), \\
\text{d}R(t) &= (\bar{\delta}I(t) + \bar{\beta}S(t) - \bar{\mu}R(t))\text{d}t,
\end{align*}
\]

(4)

where \(\bar{A} \in [A, \bar{A}], \bar{\beta} \in [\beta, \bar{\beta}], \bar{\mu} \in [\mu, \bar{\mu}], \bar{\delta} \in [\delta, \bar{\delta}], \bar{\sigma} \in [\sigma, \bar{\sigma}], \bar{r} \in [r, \bar{r}], \text{ and } \bar{\eta} \in [\eta, \bar{\eta}],\) is provided an interval-valued functional form of parameters by the following stochastic differential equation (SDE):

| Parameters | Interpretation |
|-----------|----------------|
| \(A\)     | The recruitment rate corresponding to births and immigration. |
| \(\mu\)   | The natural mortality rate. |
| \(\beta\) | The transmission rate. |
| \(\delta\) | The rate of individuals leaving \(I\) to \(R\). |
| \(r\)     | The disease-related death rate. |
| \(\theta\) | The successful vaccination rate. |
Theorem 2. For any initial value \( S_0, I_0, R_0 \in \mathbb{R}_+^3 \), there exists a unique positive solution \((S(t), I(t), R(t))\) of system (5) on the interval \([0, t)\), where \(t \geq \tau_0\) is the explosion time. To show that the solution is global, we only need to prove that \( \tau_0 = \infty \) a.s. Let \( \epsilon_0 \) be sufficiently large such that \( S_0, I_0, R_0 \) lying within the interval \([1/\epsilon_0, \epsilon_0]\). For each integer \( \epsilon \geq \epsilon_0 \), we define the following stopping time:

\[
\tau_\epsilon = \inf\{ t \in [0, \tau_0): \min\{S(t), I(t), R(t)\} \leq \epsilon \},
\]

where, throughout this paper, we set \( \inf\emptyset = \infty \) (as usual, \( \emptyset \) denotes the empty set). Clearly, \( \tau_\epsilon \) is increasing as \( \epsilon \to \infty \). Set \( \tau_\infty = \lim_{\epsilon \to \infty} \tau_\epsilon \) whence \( \tau_\infty \leq \tau_\epsilon \). If we can prove that \( \tau_\infty = \infty \) a.s., then \( \tau_\epsilon \leq \tau_\infty \) and the solution \((S(t), I(t), R(t))\) \( \in \mathbb{R}_+^3 \) for all \( t \geq 0 \) almost surely. Specifically, to complete the proof, all we need is only to prove that \( \tau_\infty = \infty \) a.s. If this statement is false, then there exists a pair of positive constants \( T > 0 \) and \( k \in (0, 1) \) such that

\[
P\{\tau_\infty \leq T\} > k.
\]

Hence, there is an integer \( \epsilon_1 \geq \epsilon_0 \) such that

\[
P\{\tau_\epsilon \leq T\} > k \quad \text{for all } \epsilon \geq \epsilon_1.
\]

For \( t \geq \tau_\epsilon \) and each \( \epsilon \),

\[
d(S + I + R) = \left( (\bar{A})^{1-p}(\bar{A})^p - (\bar{\mu})^{1-p}(\bar{\mu})^p \right) (S + I + R) \]

\[
\quad - (\bar{\mu})^{1-p}(\bar{\mu})^p I dt
\]

\[
\leq \left( (\bar{A})^{1-p}(\bar{A})^p - (\bar{\mu})^{1-p}(\bar{\mu})^p \right) (S + I + R) dt.
\]

Then,

\[
S(t) + I(t) + R(t) \leq \left( \frac{(\bar{A})^{1-p}(\bar{A})^p}{(\bar{\mu})^{1-p}(\bar{\mu})^p} \right) S_0 + I_0 + R_0 - \left( \frac{(\bar{A})^{1-p}(\bar{A})^p}{(\bar{\mu})^{1-p}(\bar{\mu})^p} \right)
\]

\[
\leq \begin{cases} 
\frac{(\bar{A})^{1-p}(\bar{A})^p}{(\bar{\mu})^{1-p}(\bar{\mu})^p}, & \text{if } S_0 + I_0 + R_0 \leq \left( \frac{(\bar{A})^{1-p}(\bar{A})^p}{(\bar{\mu})^{1-p}(\bar{\mu})^p} \right) \\
S_0 + I_0 + R_0, & \text{if } S_0 + I_0 + R_0 > \left( \frac{(\bar{A})^{1-p}(\bar{A})^p}{(\bar{\mu})^{1-p}(\bar{\mu})^p} \right) 
\end{cases}
\]

\[
= C.
\]
Define the following Lyapunov $C^2$ function $V: \mathbb{R}^+_0 \rightarrow \mathbb{R}_+$ by
\[
V(S, I, R) = (S - 1 - \ln S) + (I - 1 - \ln I) + (R - 1 - \ln R).
\] (11)

Obviously, this function is nonnegative which can be seen from $x - 1 - \ln x > 0$ for $x > 0$.

For $0 \leq t \leq \tau_c \wedge T$, using Itô’s formula, we obtain that
\[
dV(S, I, R) = \mathcal{L}V(S, I, R)dt - (\tilde{\sigma})^{1-p} (\bar{\psi})^p SdW(t)
\] 
+ (\tilde{\sigma})^{1-p} (\bar{\psi})^p I\omega(t)
- \int_Z (\eta(u)SI + \ln(1 - \eta(u)I))\tilde{N}(dr, du)
+ \int_Z (\eta(u)SI - \ln(1 + \eta(u)S))\tilde{N}(dr, du),
\] (12)

where $\mathcal{L}$ is the differential operator, and
\[
\mathcal{L}V(S, I, R) = \left(1 - \frac{1}{R}\right)\left(\bar{S}^{1-p}(\bar{A})^p - (\tilde{\psi})^{1-p}(\bar{\psi})^p SI
\right)
- (\tilde{\psi})^{1-p}(\bar{\psi})^p S
\] 
+ \left(1 - \frac{1}{R}\right)(\tilde{\psi})^{1-p}(\bar{\psi})^p SI - (\tilde{\psi})^{1-p}(\bar{\psi})^p S
\] 
+ \left(1 - \frac{1}{R}\right)(\tilde{\psi})^{1-p}(\bar{\psi})^p I
\] 
+ \left(1 - \frac{1}{R}\right)(\tilde{\psi})^{1-p}(\bar{\psi})^p R
\] 
+ \frac{1}{2}(\tilde{\psi})^{2-2p}(\bar{\psi})^p S^2
\] 
- \int_Z (1 - \eta(u)I) + \eta(u)I)\nu(du)
\] 
- \int_Z (1 + \eta(u)S - \eta(u)S)\nu(du)
\] 
\leq \left(\bar{A}^{1-p}(\bar{A})^p + (\tilde{\psi})^{1-p}(\bar{\psi})^p C + (\tilde{\psi})^{1-p}(\bar{\psi})^p
\right)
+ \tilde{\psi}^{2-2p}(\bar{\psi})^p C^2
\] 
+ \tilde{\psi}^{1-p}(\bar{\psi})^p
+ \int_Z H_1\nu(du)
\] 
+ \int_Z H_2\nu(du),
\] (13)

where
\[
H_1 = -\ln(1 - \eta(u)I) - \eta(u)I,
\] 
\[
H_2 = -\ln(1 + \eta(u)S) + \eta(u)S.
\] (14)

By assumption 1, we have $1 - \eta(u)I > 0$. In addition, by Taylor–Lagrange’s formula, we show that
\[
H_1 = \eta(u)I - \eta(u)I + \frac{\eta^2(u)I^2}{2(1 - \kappa \eta(u)I)^2}
\] 
\leq \frac{\Gamma^2}{2(1 - \Gamma)^2}, \quad \kappa \in (0, 1).
\] (15)

Similarly, we get
\[
H_2 = -\eta(u)S + \eta(u)S + \frac{\eta^2(u)S^2}{2(1 + \kappa \eta(u)S)^2}
\] 
\leq \frac{\Gamma^2}{2(1 - \Gamma)^2}, \quad \kappa \in (0, 1).
\] (16)

Therefore,
\[
\mathcal{L}V(S, I, R) \leq (\tilde{\psi})^{1-p}(\bar{\psi})^p S + \tilde{\psi}^{1-p}(\bar{\psi})^p C + \tilde{\psi}^{1-p}(\bar{\psi})^p
\] 
+ \tilde{\psi}^{2-2p}(\bar{\psi})^p C^2
\] 
+ \tilde{\psi}^{1-p}(\bar{\psi})^p
+ \frac{\Gamma^2}{(1 - \Gamma)^2} \nu(Z)
\] 
\geq \tilde{C},
\] (17)

where $\tilde{C}$ is a positive constant. Integrating both sides of (12) from 0 to $\tau_c \wedge T$, and taking expectation, we get
\[
\mathbb{E}\mathcal{V}(S(\tau_c \wedge T), I(\tau_c \wedge T), R(\tau_c \wedge T)) \leq \mathcal{V}(S_0, I_0, R_0) + \tilde{C}T.
\] (18)

Setting $\Omega_\epsilon = \{ \tau_c \leq T \}$ for $\epsilon \geq \epsilon_0$ and by (8), we have $\mathbb{P}(\Omega_\epsilon) \geq k$. For $\omega \in \Omega_\epsilon$, there is some component of $S(\tau_c)$, $I(\tau_c)$, and $R(\tau_c)$ equals either $\epsilon$ or $1/\epsilon$. Hence, $\mathcal{V}(S(\tau_c), I(\tau_c), R(\tau_c))$ is not less than $\epsilon - 1 - \ln \epsilon$ or $(1/\epsilon) - 1 - \ln (1/\epsilon)$. Consequently,
\[
\mathcal{V}(S(0), I(0), R(0)) + \tilde{C}T \geq \mathbb{E}\left(\int_{\Omega_\epsilon} \mathcal{V}(S(\tau_c), I(\tau_c), R(\tau_c), w))\right)
\] 
\geq k\left((\epsilon - 1 - \ln \epsilon) + \left(1 - 1 - \ln \frac{1}{\epsilon}\right)\right).
\] (19)

Extending $\epsilon$ to $00$ a.s. which completes the proof of the theorem.

Remark 2.5. From mathematical and biological considerations, we can study the disease dynamics of the model (5) in the following bounded set:
\[
\Delta = \left\{(S, I, R) \in \mathbb{R}^+_0; \ S + I + R \leq \frac{\bar{A}^{1-p}(\bar{A})^p}{(\bar{\psi})^{1-p}(\bar{\psi})^p} \ a.s. \right\}.
\] (20)

Therefore, the region $\Delta$ is almost surely positively invariant set by system (5).
2.3. Existence and Uniqueness of a Stationary Distribution to System (5). Our aim in this subsection is to give the appropriate condition for the SDE model (5) which has a unique ergodic stationary distribution. To this end, we introduce the following lemma known as mutually exclusive possibilities. It was proved by Stettner [1].

**Lemma 1** (see [1]). Let $X(t) \in \mathbb{R}^n$ be a stochastic Feller process, then either an ergodic probability measure exists, or

$$\lim \sup_{t \rightarrow \infty} \frac{1}{t} \int_0^t \mathbb{P} \left( u, X_0, \Sigma \right) \nu(du) = 0,$$

for any compact set $\Sigma \in \mathbb{R}^n$.

where the supremum is taken over all initial distributions $\nu$ on $\mathbb{R}^d$ and $\mathbb{P} \left( t, X_0, \Sigma \right)$ is the probability for $X(t) \in \Sigma$ with $X(0) = X_0 \in \mathbb{R}^n$.

For convenience, we introduce the following notation. Let

$$\mathcal{R}^0 = \frac{1}{\left( (\bar{\mu})^{1-p} (\bar{\mu})^p + (\bar{\delta})^{1-p} (\bar{\delta})^p + (\bar{\gamma})^{1-p} (\bar{\gamma})^p \right)^p} \left( \frac{\bar{\beta} \lambda}{(\bar{\mu})^{1-p} (\bar{\mu})^p + (\bar{\delta})^{1-p} (\bar{\delta})^p} - \frac{\tilde{\sigma}^2 (\lambda)^2 (\lambda)^2}{2 (\mu)^2 (\mu)^2} \right), \tag{22}$$

where $\tilde{\sigma}^2 = (\bar{\sigma})^{2-2p} (\bar{\sigma})^2 + \int_\Gamma (\eta^2 (u) / (1 - \Gamma)^2) \nu(du)$.

For the ergodicity of system (5), we have the following result.

**Theorem 3.** If $\mathcal{R}^0 > 1$, the stochastic system (5) admits a unique stationary distribution and it has the ergodic property for any initial value $(S_0, I_0, R_0) \in \Delta$.

**Proof.** The following proof is divided into three steps:

Step I. Similar to the proof of Lemma 3.2 in [24] or Theorem 2.5 in [25], we briefly verify the Feller property of the SDE model (5). The main purpose of the next steps is to prove that (21) is impossible.

Step II. Define

$$\mathcal{W} = \ln I + \left( \frac{(\bar{\beta})^{1-p} (\bar{\beta})^p}{(\bar{\mu})^{1-p} (\bar{\mu})^p + (\bar{\delta})^{1-p} (\bar{\delta})^p} \right) S(t). \tag{23}$$

Applying Itô’s formula gives

$$d\mathcal{W}(t) = \left( \frac{(\bar{\beta})^{1-p} (\bar{\beta})^p}{(\bar{\mu})^{1-p} (\bar{\mu})^p + (\bar{\delta})^{1-p} (\bar{\delta})^p} - I(t) dt + \right) \left( \ln (1 + \eta(u) S(t^-)) - \eta(u) S(t^-) \nu(du) \right) dt$$

$$+ (\bar{\sigma})^{1-p} (\bar{\sigma})^p S(t^-) dW(t) + \int_Z \ln (1 + \eta(u) S(t^-)) N(du, du)$$

$$+ \frac{(\bar{\beta} \lambda)^p}{(\bar{\mu})^{1-p} (\bar{\mu})^p + (\bar{\delta})^{1-p} (\bar{\delta})^p} dt - \left( \frac{(\bar{\beta})^{2-2p} (\bar{\beta})^2}{(\bar{\mu})^{1-p} (\bar{\mu})^p + (\bar{\delta})^{1-p} (\bar{\delta})^p} \right) S(t) I(t) dt$$

$$- \left( \frac{(\bar{\beta})^{1-p} (\bar{\beta})^p}{(\bar{\mu})^{1-p} (\bar{\mu})^p + (\bar{\delta})^{1-p} (\bar{\delta})^p} \right) \left( I(t) \right) \left( \bar{\sigma} \right)^{1-p} (\bar{\sigma})^p S(t^-) dW(t)$$

$$- \left( \frac{(\bar{\beta})^{1-p} (\bar{\beta})^p}{(\bar{\mu})^{1-p} (\bar{\mu})^p + (\bar{\delta})^{1-p} (\bar{\delta})^p} \right) \left( \bar{\sigma} \right)^{1-p} (\bar{\sigma})^p S(t^-) I(t^-) dW(t)$$

$$- \left( \frac{(\bar{\beta})^{1-p} (\bar{\beta})^p}{(\bar{\mu})^{1-p} (\bar{\mu})^p + (\bar{\delta})^{1-p} (\bar{\delta})^p} \right) \int_Z \eta(u) S(t^-) I(t^-) N(du, du).$$
Noting that $0 < S < (\tilde{A}^{1-p}\tilde{A})^p/(\tilde{\mu}^{1-p}\tilde{\mu})^p$, the equality (24) can be rewritten as follows:

$$
\mathbb{d}W(t) \succeq \left(\frac{(\tilde{\mu}^{1-p}(\tilde{\mu})^p}{(\tilde{\mu}^{1-p}(\tilde{\mu})^p + (\tilde{\theta})^{1-p}(\tilde{\theta})^p)} - \left(\tilde{\theta}^{1-p}(\tilde{\theta})^p + (\tilde{\theta})^{1-p}(\tilde{\theta})^p\right)S(t)I(t)\right) dt \\
+ \int_{Z} \ln(1 + \eta(u)S(t^-)) - \eta(u)S(t^-))\nu(du) dt \\
+ (\tilde{\theta})^{1-p}(\tilde{\theta})^p S(t^-) dW(t) + \int_{Z} \ln(1 + \eta(u)S(t^-))\bar{N}(dt, du) \\
\frac{(-\tilde{\theta}^{1-p}(\tilde{\theta})^p)}{(\tilde{\mu})^{1-p}(\tilde{\mu})^p + (\tilde{\theta})^{1-p}(\tilde{\theta})^p)} \bar{N}(dt, du) \\
\frac{(-\tilde{\theta}^{1-p}(\tilde{\theta})^p)}{(\tilde{\mu})^{1-p}(\tilde{\mu})^p + (\tilde{\theta})^{1-p}(\tilde{\theta})^p)} \int_{Z} \eta(u)S(t^-) I(t^-) \bar{N}(dt, du).
$$

Integrating the inequality (25) from 0 to $t$ leads to

$$
W(t) - W(0) \succeq \mathbb{E}\left[ \left(\frac{(\tilde{\mu}^{1-p}(\tilde{\mu})^p}{(\tilde{\mu}^{1-p}(\tilde{\mu})^p + (\tilde{\theta})^{1-p}(\tilde{\theta})^p)} - \left(\tilde{\theta}^{1-p}(\tilde{\theta})^p + (\tilde{\theta})^{1-p}(\tilde{\theta})^p\right)S(t)I(t)\right) dt \\
+ \int_{0}^{t} \left(\frac{(\tilde{\mu})^{1-p}(\tilde{\mu})^p}{(\tilde{\mu})^{1-p}(\tilde{\mu})^p + (\tilde{\theta})^{1-p}(\tilde{\theta})^p)} \bar{N}(ds, du) \\
+ K_1(t) + K_2(t) + K_3(t) + K_4(t),
$$

where

$$
K_1(t) = \int_{0}^{t} (\tilde{\theta})^{1-p}(\tilde{\theta})^p S(s^-) dW(s),
$$

$$
K_2(t) = \frac{(-\tilde{\theta}^{1-p}(\tilde{\theta})^p)}{(\tilde{\mu})^{1-p}(\tilde{\mu})^p + (\tilde{\theta})^{1-p}(\tilde{\theta})^p)} \int_{0}^{t} S(s^-) I(s^-) dW(s),
$$

$$
K_3(t) = \int_{0}^{t} \int_{Z} \ln(1 + \eta(u)S(s^-))\bar{N}(ds, du),
$$

$$
K_4(t) = \frac{(-\tilde{\theta}^{1-p}(\tilde{\theta})^p)}{(\tilde{\mu})^{1-p}(\tilde{\mu})^p + (\tilde{\theta})^{1-p}(\tilde{\theta})^p)} \cdot \int_{0}^{t} \int_{Z} \eta(u)S(s^-) I(s^-) \bar{N}(ds, du).
$$

The quadratic variation of $K_1$ is defined by $\langle K_1, K_1 \rangle_t = \int_{0}^{t} (\tilde{\theta}^{1-p}(\tilde{\theta})^p S^2(s) ds$. Therefore, we get

$$
\limsup_{t \to \infty} \frac{\langle K_1, K_1 \rangle_t}{t} = (\tilde{\theta}^{1-p}(\tilde{\theta})^p \leq \infty \quad a.s. \hspace{1cm} (28)
$$

Similarly, we have

$$
\limsup_{t \to \infty} \frac{\langle K_2, K_2 \rangle_t}{t} = \frac{(-\tilde{\theta}^{1-p}(\tilde{\theta})^p)}{(\tilde{\mu})^{1-p}(\tilde{\mu})^p + (\tilde{\theta})^{1-p}(\tilde{\theta})^p)} \cdot \limsup_{t \to \infty} \frac{1}{t} \int_{0}^{t} S(s^-) I(s^-) ds < \infty \quad a.s. \hspace{1cm} (29)
$$

By the assumption 1, we deduce that

$$
\ln(1 - \Gamma) \leq \ln(1 + \eta(u)S(s^-)) \leq \ln(1 + \Gamma). \hspace{1cm} (30)
$$

Then

$$
\limsup_{t \to \infty} \frac{\langle K_3, K_3 \rangle_t}{t} = \limsup_{t \to \infty} \frac{1}{t} \int_{0}^{t} \ln(1 + \eta(u)S(s^-))\nu(du) ds \\
\leq \max\left\{ \ln(1 + \Gamma)^2, (\ln(1 - \Gamma)^2 \right\} \nu(Z) < \infty \quad a.s., \hspace{1cm} (31)
$$

$$
\limsup_{t \to \infty} \frac{\langle K_4, K_4 \rangle_t}{t} \leq \frac{(-\tilde{\theta}^{1-p}(\tilde{\theta})^p)}{(\tilde{\mu})^{1-p}(\tilde{\mu})^p + (\tilde{\theta})^{1-p}(\tilde{\theta})^p)} \cdot \nu(Z) < \infty \quad a.s. \hspace{1cm} (32)
$$
According to the strong law of large numbers for local martingales [26], one can conclude that
\[
\lim_{t \to \infty} \frac{1}{t} K_i(t) = 0, \quad \text{a.s.,} \quad i = 1, 2, 3, 4. \tag{32}
\]

By using (16) and assumption 1, we get
\[
\frac{1}{t} \int_0^t \left( \ln(1 + \eta(u)S(s^*)) - \eta(u)S(s^*) \right) \nu(du) \, ds \geq - \frac{1}{2} \frac{(\bar{A})^{2-2p}(\bar{A})^{2p}}{(\bar{\mu})^{1-p}(\bar{\mu})^p} \int \eta^2(u) \, \nu(du). \tag{33}
\]

Let
\[
\bar{\sigma}^2 = (\bar{\sigma})^{2-2p}(\bar{\sigma})^{2p} + \int \frac{\eta^2(u)}{(1-\Gamma)^2} \, \nu(du). \tag{34}
\]

\[
\liminf_{t \to \infty} \frac{1}{t} \int_0^t (\bar{\mu})^{1-p}(\bar{\mu})^p S(I)(s) \, ds \geq \frac{(\bar{\mu})^{1-p}(\bar{\mu})^p + (\bar{\sigma})^{1-p}(\bar{\sigma})^p}{(\bar{\mu})^{1-p}(\bar{\mu})^p} \left( (\bar{\mu})^{1-p}(\bar{\mu})^p + (\bar{\sigma})^{1-p}(\bar{\sigma})^p \right)
\]
\[
- \left( (\bar{\mu})^{1-p}(\bar{\mu})^p + (\bar{\sigma})^{1-p}(\bar{\sigma})^p + (\bar{\sigma})^{1-p}(\bar{\sigma})^p \right)(\mathcal{R}^2_0 - 1) > 0 \quad \text{a.s.} \tag{35}
\]

Thus, we can derive that
\[
\liminf_{t \to \infty} \frac{1}{t} \int_0^t (\bar{\mu})^{1-p}(\bar{\mu})^p S(I)(s) \, ds \geq \frac{(\bar{\mu})^{1-p}(\bar{\mu})^p + (\bar{\sigma})^{1-p}(\bar{\sigma})^p}{(\bar{\mu})^{1-p}(\bar{\mu})^p} \left( (\bar{\mu})^{1-p}(\bar{\mu})^p + (\bar{\sigma})^{1-p}(\bar{\sigma})^p \right)(\mathcal{R}^2_0 - 1) > 0 \quad \text{a.s.} \tag{36}
\]

Step III. To continue our analysis, we need to set the following subsets:

\[
\Omega_1 = \{S, I, R \in \mathbb{R}^3 \mid S \geq \epsilon, \text{ and, } I \geq \epsilon \},
\]
\[
\Omega_2 = \{S, I, R \in \mathbb{R}^3 \mid S \leq \epsilon \},
\]
\[
\Omega_3 = \{S, I, R \in \mathbb{R}^3 \mid I \leq \epsilon \},
\]

where \( \epsilon > 0 \) is a positive constant to be determined later. It then follows from (36) that

\[
\liminf_{t \to \infty} \frac{1}{t} \int_0^t \mathbb{E}(\bar{\mu})^{1-p}(\bar{\mu})^p S(I)(u) \, du \geq \liminf_{t \to \infty} \frac{1}{t} \int_0^t \mathbb{E}(\bar{\mu})^{1-p}(\bar{\mu})^p S(I)(u) \, du
\]
\[
- \limsup_{t \to \infty} \frac{1}{t} \int_0^t \mathbb{E}(\bar{\mu})^{1-p}(\bar{\mu})^p S(I)(u) \, du
\]
\[
- \limsup_{t \to \infty} \frac{1}{t} \int_0^t \mathbb{E}(\bar{\mu})^{1-p}(\bar{\mu})^p S(I)(u) \, du
\]
\[
\geq \left( (\bar{\mu})^{1-p}(\bar{\mu})^p + (\bar{\sigma})^{1-p}(\bar{\sigma})^p \right)(\mathcal{R}^2_0 - 1)
\]
\[
- \frac{2(\bar{\mu})^{1-p}(\bar{\mu})^p \epsilon}{(\bar{\mu})^{1-p}(\bar{\mu})^p} \tag{38}
\]

We can choose
\[
\epsilon \leq \frac{(\bar{\mu})^{1-p}(\bar{\mu})^p + (\bar{\sigma})^{1-p}(\bar{\sigma})^p)(\bar{\mu})^{1-p}(\bar{\mu})^p}{4(\bar{\mu})^{1-p}(\bar{\mu})^p (\bar{A})^{1-p}(\bar{A})^p}
\]
\[
(\bar{\mu})^{1-p}(\bar{\mu})^p + (\bar{\sigma})^{1-p}(\bar{\sigma})^p + (\bar{\sigma})^{1-p}(\bar{\sigma})^p (\mathcal{R}^2_0 - 1), \tag{39}
\]

then, we obtain
\[
\liminf_{t \to +\infty} \frac{1}{t} \int_0^t \mathbb{E}\left[ (\tilde{\beta})^{1-p} (\tilde{\beta})^p S(u)I(u)1_{\Omega_1} \right] du \\
= \frac{1}{2} \frac{(\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p}{(\tilde{\beta})^{1-p} (\tilde{\beta})^p} + (\tilde{\rho})^{1-p} (\tilde{\rho})^p} \\
\cdot (R_0^* - 1) > 0 \quad \text{a.s.}
\] (40)

Let \( a \) and \( b \) two real numbers greater than 1 such that \((1/a) + (1/b) = 1\). By utilizing Young inequality \( xy \leq (x^a/a) + (y^b/b) \) for all \( x, y > 0 \), we get
\[
\liminf_{t \to +\infty} \frac{1}{t} \int_0^t \mathbb{E}\left[ (\tilde{\beta})^{1-p} (\tilde{\beta})^p S(u)I(u)1_{\Omega_1} \right] du \\
\leq \liminf_{t \to +\infty} \frac{1}{t} \int_0^t \mathbb{E}\left[ a^{-1}(\tilde{\beta})^{1-p} (\tilde{\beta})^p S(u)I(u) \right] du + b^{-1} \cdot 1_{\Omega_1} du \\
\leq a^{-1}(\tilde{\beta})^{1-p} (\tilde{\beta})^p a^{-1} \left( (\tilde{\lambda})^{1-p} (\tilde{\lambda})^p \right)^{-2a} \\
+ \liminf_{t \to +\infty} \frac{1}{t} \int_0^t \mathbb{E}(b^{-1} \cdot 1_{\Omega_1}) du.
\] (41)

where \( \omega \) is a positive constant satisfying
\[
\omega^a \leq a^{-4} \left( (\tilde{\beta})^{1-p} (\tilde{\beta})^p \right)^{-2a} \left( (\tilde{\lambda})^{1-p} (\tilde{\lambda})^p \right)^{-2a} \\
\cdot (\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p (\tilde{\rho})^{1-p} (\tilde{\rho})^p \\
+ (\tilde{\rho})^{1-p} (\tilde{\rho})^p \cdot (R_0^* - 1).
\] (42)

From (41), we deduce that
\[
\liminf_{t \to +\infty} \frac{1}{t} \int_0^t \mathbb{E}\left[ 1_{\Omega_1} \right] du \geq \frac{(\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p) b \omega b}{4(\tilde{\beta})^{1-p} (\tilde{\beta})^p} \\
\cdot (\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p + (\tilde{\rho})^{1-p} (\tilde{\rho})^p \cdot (R_0^* - 1) > 0 \quad \text{a.s.}
\] (43)

Setting
\[
\Omega_4 = \{(S, I, R) \in \mathbb{R}_+^3 | S \geq \zeta, \text{or}, I \geq \zeta \}, \\
\Omega_5 = \{(S, I, R) \in \mathbb{R}_+^3 | e \leq S \leq \zeta, \text{and}, e \leq I \leq \zeta \},
\] (44)

where \( \zeta > 0 \) is a positive constant to be explained in the following. By using the Tchebychev inequality, we can observe that
\[
\mathbb{E}\left[ 1_{\Omega_4} \right] \leq \mathbb{P}(S(t) \geq \zeta) + \mathbb{P}(I(t) \geq \zeta) \leq \frac{1}{\zeta} \mathbb{E}(S(t) + I(t)) \\
\leq \frac{1}{\zeta} \frac{\tilde{\lambda}^{1-p} (\tilde{\lambda})^p}{(\tilde{\mu})^{1-p} (\tilde{\mu})^p}.
\] (45)

Computational and Mathematical Methods in Medicine

Choosing
\[
\frac{1}{\xi} \leq \frac{(\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p) b \omega b}{8(\tilde{\beta} \tilde{A})^{1-p} (\tilde{\beta} \tilde{A})^p} \\
\cdot (\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p + (\tilde{\rho})^{1-p} (\tilde{\rho})^p \cdot (R_0^* - 1).
\] (46)

We thus obtain
\[
\limsup_{t \to +\infty} \frac{1}{t} \int_0^t \mathbb{E}(1_{\Omega_4}) du \leq \frac{(\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p) b \omega b}{8(\tilde{\beta} \tilde{A})^{1-p} (\tilde{\beta} \tilde{A})^p} \\
\cdot (\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p + (\tilde{\rho})^{1-p} (\tilde{\rho})^p \cdot (R_0^* - 1).
\] (47)

According to (43), one can derive that
\[
\liminf_{t \to +\infty} \frac{1}{t} \int_0^t \mathbb{E}\left[ 1_{\Omega_5} \right] du \geq \liminf_{t \to +\infty} \frac{1}{t} \int_0^t \mathbb{E}(1_{\Omega_4}) du \\
\leq \limsup_{t \to +\infty} \frac{1}{t} \int_0^t \mathbb{E}(1_{\Omega_4}) du \\
\geq \frac{(\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p) b \omega b}{8(\tilde{\beta} \tilde{A})^{1-p} (\tilde{\beta} \tilde{A})^p} \\
\cdot (\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p + (\tilde{\rho})^{1-p} (\tilde{\rho})^p \cdot (R_0^* - 1) \quad a.s.
\] (48)

Based on the above analysis, we have determined a compact domain \( \Sigma \subset \mathbb{R}_+^3 \) such that
\[
\liminf_{t \to +\infty} \frac{1}{t} \int_0^t \mathbb{E}(u, (S_0, I_0, R_0), \Sigma) du \\
\geq \frac{(\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p) b \omega b}{8(\tilde{\beta} \tilde{A})^{1-p} (\tilde{\beta} \tilde{A})^p} \\
\cdot (\tilde{\mu})^{1-p} (\tilde{\mu})^p + (\tilde{\theta})^{1-p} (\tilde{\theta})^p + (\tilde{\rho})^{1-p} (\tilde{\rho})^p \cdot (R_0^* - 1) \quad a.s.
\] (49)

Applying similar arguments to those in [24], we show the uniqueness of the ergodic stationary distribution of our model (5), denoted by \( \pi(\cdot) \). This completes the proof.

2.4. Numerical Simulations. In this subsection, in order to show different dynamical results of the stochastic model (2) under imprecise parameter values, we present some numerical simulations. We use Milstein’s method to simulate the trajectories of the stochastic model (5). The parameters values are given in the following list. For the purpose of showing the effects of imprecise parameters and Lévy noise on Hepatitis B dynamics, we have realized the simulation.
10000 times. We assume that $\eta(u) = 0.03$, $Z = (0, \infty)$, and $\nu(Z) = 1$. Then, we obtain the following results: noticing that the assumption 1 is always held with parameters’ value in Table 2. From Figures 1–3, we show the existence of the unique stationary distributions for $S(t)$, $I(t)$, and $R(t)$ of model (5) at $t = 300$, where the smooth curves are the probability density functions of $S(t)$, $I(t)$, and $R(t)$, respectively. It can be obviously observed that the

Table 2: Parameters’ value used in numerical simulations.

| Notation | Value | Source | Notation | Value |
|----------|-------|--------|----------|-------|
| $\alpha$ | 0.4   | [9]    | $\alpha$ | 0.6   |
| $\mu$    | 0.09  | [9]    | $\mu$    | 0.2   |
| $\beta$  | 0.1   | [9]    | $\beta$  | 0.2   |
| $\delta$ | 0.3   | [9]    | $\delta$ | 0.5   |
| $\gamma$ | 0.1   | [9]    | $\gamma$ | 0.3   |
| $\bar{\delta}$ | 0.2 | Assumed | $\bar{\delta}$ | 0.3 |
| $\bar{\sigma}$ | 0.08 | Assumed | $\bar{\sigma}$ | 0.1 |

Figure 1: The trajectories and histogram of solution of model (5) with initial value $(S_0, I_0, R_0) = (0.5, 0.3, 0.2)$ and $p = 1$. 

| Time | Density |
|------|---------|
| 0    | 0.4     |
| 1    | 0.45    |
| 2    | 0.5     |
| 3    | 0.6     |
| 4    | 0.7     |
| 5    | 0.8     |
| 6    | 0.9     |

| Density |
|---------|
| 0.35    |
| 0.4     |
| 0.45    |
| 0.5     |
| 0.55    |
| 0.6     |
| 0.65    |

| Density |
|---------|
| 0.08    |
| 0.1     |
| 0.12    |
| 0.14    |
| 0.16    |
| 0.18    |
| 0.2     |
solution of the SDE model (5) persists in the mean. Furthermore, different values of the parameter imprecision $p$ can also crucially affect the persistence of Hepatitis B (see Table 3).

3. Discussion

In the study of the dynamics of stochastic systems, the existence of an ergodic stationary distribution is one of the most important and significant characteristics. For this purpose, we have used the Feller property and mutually exclusive possibilities lemma to establish the sharp and optimal condition for the existence of the stationary distribution without employing the classical Lyapunov method. To ensure the realistic aspect of our model, we replaced constant parameters in the model (2) by imprecise ones.

Based on Theorem 4.2 in [23], for any $\pi$-integrable function $g: \mathbb{R}_+ \rightarrow \mathbb{R}$,

$$\mathbb{P}\left( \lim_{t \to \infty} \frac{1}{t} \int_0^t g(X(s)) ds = \int_{\mathbb{R}_+} g(x) \pi(x) dx \right) = 1.$$  

(50)

The ergodic property for HBV means that the stochastic model has a unique stationary distribution which predicts the survival of the infected population in the future. That means the HBV persists for all time regardless of the initial conditions [27]. Furthermore, the ergodic property grants a reason why the integral average of a solution of system (5) converges to a fixed point whilst the system may fluctuate around as time goes by.
Data Availability

The theoretical data used to support the findings of this study are included within the article.

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this paper.

Authors’ Contributions

The authors declare that the study was conducted in collaboration with the same responsibility. All authors read and approved the final manuscript.

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