Research Article

An Advance Computing Numerical Heuristic of Nonlinear SIR Dengue Fever System Using the Morlet Wavelet Kernel

Muhammad Umar,¹ Zulqurnain Sabir,¹ Muhammad Asif Zahoor Raja,² K. S. Al-Basyouni,³ S. R. Mahmoud,⁴ and Yolanda Guerrero Sánchez ⁵

¹Department of Mathematics and Statistics, Hazara University, Mansehra, Pakistan
²Future Technology Research Center, National Yunlin University of Science and Technology, 123 University Road, Section 3, Douliou, Yunlin 64002, Taiwan
³Mathematics Department, Faculty of Science, King Abdulaziz University, Jeddah, Saudi Arabia
⁴GRC Department, Faculty of Applied Studies, King Abdulaziz University, Jeddah, Saudi Arabia
⁵Department of Dermatology, Stomatology, Radiology and Physical Medicine, University of Murcia, Murcia, Spain

Correspondence should be addressed to Yolanda Guerrero Sánchez; yolanda.guerreros@um.es

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This study is associated to solve the nonlinear SIR dengue fever system using a computational methodology by operating the neural networks based on the designed Morlet wavelet (MWNNs), global scheme as genetic algorithm (GA), and rapid local search scheme as interior-point algorithm (IPA), i.e., GA-IPA. The optimization of fitness function based on MWNNs is performed for solving the nonlinear SIR dengue fever system. This MWNNs-based fitness function is accessible using the differential system and initial conditions of the nonlinear SIR dengue fever system. The designed procedures based on the MWNN-GA-IPA are applied to solve the nonlinear SIR dengue fever system to check the exactness, precision, constancy, and efficiency. The achieved numerical form of the nonlinear SIR dengue fever system via MWNN-GA-IPA was compared with the Runge–Kutta numerical results that verify the significance of MWNN-GA-IPA. Moreover, statistical reflections through different measures for the nonlinear SIR dengue fever system endorse the precision and convergence of the computational MWNN-GA-IPA.

1. Introduction

Dengue fever disease (DFD) is one of the epidemics, infectious, and serious diseases that disturbed about 2.5 billion individuals all over the world. DFD occurred in some main countries of Southeast Asia due to the hot seasons. The infectious DFD grows fast, when the environment alters and becomes dangerous because of the shortage of information amongst the individuals [1]. DFD is an epidemiologic transmittable fever formed by dengue infection (DI) that is conveyed through mosquitoes to humans and apes [2]. Some symptoms scientifically produced by DFD are headache, joint pain, skin rash, and DI. The global World Health Organization (WHO) classified DI into a hemorrhagic dengue fever before one decade [3]. DI is categorized into three organizational proteins, membrane protein (M), envelope protein (E), and capsid protein (C), while it has seven proteins that are nonstructural and their names are NS-I, NS-IIA, NS-III, NS-IIB, NS-IVA, NS-IVB, and NS-V [4]. According to envelope protein antigenicity, DI is ordered into four classes of microorganisms, which are infected as well as pathogenic [5].

DI has been reported in several zones of China and still any scientific report cannot prove it a confined endemic [6], whereas all other subgroups of DI have been imported; DI-I and DI-II are considered in China one of the main stereotypes endemic. In the Chinese province Guangdong, DV-I reported around 70% in 2014 and 80% in 2015. Presently,
an amino-based acid localizes transformation in NS-I of the Asian "Zika" inheritance, which has been described to grow the production of NS-I in the diseased host to make the virus convenient to Aedes type of mosquitoes, which is the focal wide spreading cause of virus "Zika" since 2015 [7].

Some epidemic infections such as COVID-19 are the existing transferred diseases, which have covered the whole world and the infection rate along with the number of deaths from COVID-19 increased steadily [8]. Another common disease is malaria, which is not directly transmitted from host to host. Protozoa is one of the transferred diseases, which spreads due to the anopheles of the female mosquito. According to the WHO report, almost 1/3 million individuals per year die from malaria. It distresses the children and pregnant women, mostly in the South African and American countries. Many chemical sprays have been widely implemented to control the mosquito population. Similarly, nonpolluting types of biological arrangements normally accomplished to emphasis the ecosystem of complicated kinds. The sterile insect apparatus (SIA) is a proficient nonpolluting scheme of insect control, which depends on the sterile male’s release. Consequently, adequately sterile males discharging causes the elimination of the wild’s population. Over a half century ago, SIA has been recognized in the Curacao Island to adjust the screwworm hover [9,10].

The aim of this study is to solve the nonlinear SIR dengue fever system using a computational methodology by operating the MWNNs, global search GA, and rapid local search IPA, i.e., MWNN-GA-IPA. The stochastic approaches have been investigated normally to solve a number of applications directed to the differential linear/nonlinear systems [11, 12]. However, no one has applied the MWNNs to solve the nonlinear SIR dengue fever system. Few recent reported submissions of stochastic solvers are eye surgery model, functional singular system, Thomas–Fermi singular equation, HIV-based infection system, biological form of the prey-predator system, periodic singular problems, singular three-point differential model, COVID-19 SITR system, multifractional singular models, system of heat transmission in human head, and mosquito spreading in heterogeneous conditions [13–15]. The intention of this study is to solve the nonlinear SIR dengue fever system using the MWNN-GA-IPA. The literature form of nonlinear SIR dengue fever system is written as [16]

\[
\begin{align*}
\frac{dX(t)}{dr} &= \mu_h - \mu_h X(t) - \alpha X(t) Z(t), \quad X(0) = I_1, \\
\frac{dY(t)}{dr} &= \alpha X(t) Z(t) - \beta Y(t), \quad Y(0) = I_2, \\
\frac{dZ(t)}{dr} &= \gamma Y(t) - \gamma Y(t) Z(t) - \delta I Z(t), \quad Z(0) = I_3,
\end{align*}
\]

Section 4 describes the final remarks and future research reports.

2. Designed Procedure

The proposed structure of the MW-GA-IPA is used to solve the nonlinear SIR dengue fever system described in two phases as follows:

(i) An objective function using the MW is considered to activate the neural networks

(ii) Necessary clarifications are provided to enhance the merit function by applying the hybrid of GA-IPA

2.1. Designed Procedure Using MW Function. The mathematical design of the nonlinear SIR dengue fever system is described by using the achieved results of susceptible \(X(t)\), infected \(Y(t)\), and recovered \(Z(t)\) with the derivatives of these classes, written as [17]
The simplified form of system (2) using the above MW function is given as

\[
\begin{align*}
[\tilde{X}(\tau), \tilde{Y}(\tau), \tilde{Z}(\tau)] &= \left[ \sum_{i=1}^{m} v_{X,i} H\left(w_{X,i} \tau + u_{X,i}\right), \sum_{i=1}^{m} v_{Y,i} H\left(w_{Y,i} \tau + u_{Y,i}\right), \sum_{i=1}^{m} v_{Z,i} H\left(w_{Z,i} \tau + u_{Z,i}\right) \right], \\
[\tilde{X}^{(n)}(\tau), \tilde{Y}^{(n)}(\tau), \tilde{Z}^{(n)}(\tau)] &= \left[ \sum_{i=1}^{m} v_{X,i} H^{(n)}\left(w_{X,i} \tau + u_{X,i}\right), \sum_{i=1}^{m} v_{Y,i} H^{(n)}\left(w_{Y,i} \tau + u_{Y,i}\right), \sum_{i=1}^{m} v_{Z,i} H^{(n)}\left(w_{Z,i} \tau + u_{Z,i}\right) \right].
\end{align*}
\]

Where \( W \) is the unknown weight vector given as

\[
W = [W_X; W_Y; W_Z], \quad \text{for} \quad W_X = [v_X, \omega_X, u_X], \quad W_Y = [v_Y, \omega_Y, u_Y], \quad \text{and} \quad W_Z = [v_Z, \omega_Z, u_Z],
\]

The MWNN has not been implemented before to solve the nonlinear SIR dengue fever system. The mathematical form of MW function is written as [18]

\[
H(\tau) = \cos(1.75)e^{-0.5\tau^2}.
\]
An error function based on the merit function is given as

\[ e = \sum_{j=1}^{4} e_j, \]  

\[ e_1 = \frac{1}{N} \sum_{i=1}^{N} \left[ X_i - \mu_h \tilde{X}_i + \alpha X_i \tilde{Z}_i \right]^2, \]  

\[ e_2 = \frac{1}{N} \sum_{i=1}^{N} \left[ Y_i - \alpha \tilde{X}_i \tilde{Z}_i + \beta Y_i \right]^2, \]  

\[ e_3 = \frac{1}{N} \sum_{i=1}^{N} \left[ Z_i - \gamma \tilde{Y}_i + \gamma \tilde{Z}_i + \delta \tilde{Z}_i \right]^2, \]  

\[ e_4 = \frac{1}{3} \left[ (X_0 - I_1)^2 + (Y_0 - I_2)^2 + (Z_0 - I_3)^2 \right], \]  

where \( X_i = X(\tau_i), \ Y_i = Y(\tau_i), \ Z_i = Z(\tau_i), \) \( Nh = 1, \) and \( \tau_i = ih. \) \( \tilde{X}_i, \tilde{Y}_i, \) and \( \tilde{Z}_i \) show the proposed results of the susceptible class, infected class, and recovered class. Likewise, \( e_1, e_2, \) and \( e_3 \) denote the error function related to system (1), whereas \( e_4 \) denotes the error function on the basis of initial conditions.

2.2. Optimization: MWNN-GA-IPA. The optimization performance is presented for solving the nonlinear SIR dengue fever system using the MWNN-GA-IPA. The structure of the optimization per-

formance is presented for solving the nonlinear SIR dengue fever system using the MWNN-GA-IPA. f&_he structure of

optimization procedures. GA is pragmatic frequently to regulate the accurate population to solve several

complicated or stiff systems. To attain the best model out-

comes, GAs operate through the operators based on selection, reproduction, crossover, and mutation. Few existing GA’s applications are the hospitalization expenditure system [19], feature assortment in cancer microarray [20], organization of irregular magnetic character brain tumor imageries [21], vehicle routing system [22], prediction-based traffic flow system [23], radiation shielding optimizations in the bismuth-

borate spectacles [24], prediction of air blast [25], composition optimization of cloud service [26], task arrangement models in phased range radar [27], arrangement system of microarray cancer [28], system dynamics of monorail vehicle [17], and prediction system of liver disease [29].

IPA is known as an optimized local search approach, which is performed broadly in both types of models (con-

strained/unconstrained). IPA is used in the optimization of various complicated and nonstiff natured systems. Recently, IPA is executed for image restoration [30], multistage non-

linear nonconvex models [31], viscoplastic fluid flows [32], nonsmooth contact dynamics [33], power systems [34], and dynamic flux balance analysis models [35]. The hybridization process of GA-IPA is applied to remove the laziness of GA, i.e., global approach. The pseudocode based on the designed approach MWNN-GA-IPA is provided in Table 1.

2.3. Performance Measures. The mathematical measures using the statistical operators for variance accounted for (VAF), semi-interquartile (S.I) range, Theil’s inequality coefficient (T.I.C), and mean absolute deviation (M.A.D) along with the Global VAF (G-VAF), Global M.A.D (G-

M.A.D), and Global T.I.C to solve the nonlinear SIR dengue fever system which is given as

\[
\begin{align*}
[V.A.F_X, V.A.F_Y, V.A.F_Z] & = \left[ \frac{1 - \text{var}(X_r - \tilde{X}_r)}{\text{var}(X_r)} \right] \times 100, \left[ \frac{1 - \text{var}(Y_r - \tilde{Y}_r)}{\text{var}(Y_r)} \right] \times 100, \left[ \frac{1 - \text{var}(Z_r - \tilde{Z}_r)}{\text{var}(Z_r)} \right] \times 100, \\
\end{align*}
\]

\[
\begin{align*}
[E - V.A.F_X, E - V.A.F_Y, E - V.A.F_Z] & = \left[ 100 - V.A.F_X, 100 - V.A.F_Y, 100 - V.A.F_Z \right].
\end{align*}
\]

\[
\begin{align*}
\text{S.I Range} & = -0.5 \times (Q_1 - Q_3), \\
Q_1 & = 1^{\text{st}} \text{ quartile} & Q_3 & = 3^{\text{rd}} \text{ quartile}, \\
\end{align*}
\]

\[
\begin{align*}
[T.I.C_X, T.I.C_Y, T.I.C_Z] & = \left[ \frac{\sqrt{1/(n-1) \sum_{r=1}^{n} (X_r - \tilde{X}_r)^2}}{\sqrt{1/(n-1) \sum_{r=1}^{n} X_r^2 + \sqrt{1/(n-1) \sum_{r=1}^{n} \tilde{X}_r^2}}} \right] \times \left[ \frac{\sqrt{1/(n-1) \sum_{r=1}^{n} (Y_r - \tilde{Y}_r)^2}}{\sqrt{1/(n-1) \sum_{r=1}^{n} Y_r^2 + \sqrt{1/(n-1) \sum_{r=1}^{n} \tilde{Y}_r^2}}} \right], \\
\end{align*}
\]

\[
\begin{align*}
[M.A.D_X, M.A.D_Y, M.A.D_Z] & = \left[ \sum_{r=1}^{n} |X_r - \tilde{X}_r| / \sum_{r=1}^{n} |X_r - \tilde{X}_r|, \sum_{r=1}^{n} |Y_r - \tilde{Y}_r| / \sum_{r=1}^{n} |Y_r - \tilde{Y}_r|, \sum_{r=1}^{n} |Z_r - \tilde{Z}_r| / \sum_{r=1}^{n} |Z_r - \tilde{Z}_r| \right],
\end{align*}
\]
The Problem
Formulation of MWNNs Biological Nonlinear SIR System

Optimization
Combination of global and local schemes, i.e., GA-IPA

Mathematical Model

Graphical illustration of GA-IPA
Trained weight vectors via MWNN for the proposed results
Comparative studies using the performance indices

Figure 1: Structure of the present approach to solve the nonlinear SIR dengue fever system.
Table 1: Optimization performance taking the MWNN-GA-IPA for the nonlinear SIR dengue fever system.

| Start of GA |
|---|
| Inputs: the chromosomes are characterized with the same system element as $W = [v, w, u]$ |
| Population: the chromosomes set is written as $W_X = [v_X, \omega_X, t_X], W_Y = [v_Y, \omega_Y, t_Y]$ and $W_Z = [v_Z, \omega_Z, t_Z]$ |
| Output: global values of the weight are represented as $W_{GA-Best}$ |
| Initialization: for the selection of chromosomes, select the weight vector values. |
| Fit evaluation: modify the values of fitness $e$ in population $P$ for each vector with the use of systems 4–8 |
| Stopping criteria: terminate when $[e = 10^{-20}]$, [Generations = 55], [StallLimit = 140], [PopSize = 285], and [TolFun = TolCon = $10^{-21}$] |
| Move to storage |
| Ranking: rank individual weight vector in population using the values of the fitness |
| Storage: save $W_{GA-Best}$, iterations, time, $e$, and count of function for the presence of GA |
| End of GA |

IPA starts |
| Inputs: start point: $W_{GA-Best}$ |
| Output: $W_{GA-IPA}$ shows the best weight values of GA-IPA |
| Terminate: stop when $[e = 10^{-20}]$, [Iterations = 750], [MaxFunEvals = 267000], [TolCon = TolX = $10^{-22}$], and [TolFun = $10^{-22}$] achieved. |
| Evaluation of fitness: compute $W$ and $e$ using equations (8)–(12) |
| Amendments: adjust "fmincon" for IPA, compute $e$ of better-quality of $W$ using systems 4–8 |
| Accumulate: transmute $W_{GA-IPA}$, $e$, function counts, iterations, and time for the existing IPA runs |

IPA process ends

where the approximate solutions are $\hat{X}$, $\hat{Y}$, and $\hat{Z}$, respectively.

3. Simulations of the Results

The current work is associated to solve the nonlinear SIR dengue fever system shown in system (1). The relative presentation of the obtained results using the Runge–Kutta solutions is tested to form the correctness of MWNN-GA-IPA. Additionally, statistical operators indicate the precision and accuracy of MWNN-GA-IPA. The simplified measures of the nonlinear SIR dengue fever system using the suitable values are given as

$$
\begin{cases}
\frac{d\hat{X}}{d\tau} = 0.000046 - (0.000046 + 0.375\hat{Z}(\tau))\hat{X}(\tau), & X(0) = 0.9999, \\
\frac{d\hat{Y}}{d\tau} = 0.375\hat{X}(\tau)\hat{Z}(\tau) - 0.0323\hat{Y}(\tau), & Y(0) = 0.0006, \\
\frac{d\hat{Z}}{d\tau} = 0.328833 - (0.328833\hat{Y}(\tau) + 0.0001)\hat{Z}(\tau), & Z(0) = 0.0560.
\end{cases}
$$

(14)

The fitness function for system (14) is written as

$$
e = \frac{1}{N} \sum_{i=1}^{N} \left[ \left( \frac{\hat{X}_i' - (0.000046 + 0.375\hat{Z}_i)}{10000} \right)^2 + \left( \frac{\hat{Y}_i' - 0.375\hat{X}_i\hat{Z}_i + 0.0323\hat{Y}_i}{10000} \right)^2 \right]$$

$$+ \frac{1}{3} \left[ \left( \frac{\hat{X}_0 - 9999}{10000} \right)^2 + \left( \frac{\hat{Y}_0 - 6}{10000} \right)^2 + \left( \frac{\hat{Z}_0 - 56}{10000} \right)^2 \right].$$

(15)

The nonlinear SIR dengue fever system given in system (1) is optimized using the MWNN-GA-IPA for 100 trials to attain ANN model parameters for 10 neurons. Figure 1 is drawn using the best outputs of the weight vector, i.e., $W$ for the MWNN-GA-IPA. These best weights of the output are applied to solve the estimated outcomes of the nonlinear SIR.
Figure 2: Best weight vector set and result comparison for each class of nonlinear SIR dengue fever system. (a) Best weights of $X(\tau)$ for 10 neurons. (b) Best weights of $Y(\tau)$ for 10 neurons. (c) Best weights of $Z(\tau)$ for 10 neurons. (d) Comparison for $X(\tau)$ class. (e) Comparison for $Y(\tau)$ class. (f) Comparison for $Z(\tau)$ class.
The mathematical illustrations of these estimated results from MWNN-GA-IPA are given as

\[ \tilde{X}(\tau) = 0.067 \cos(1.75(0.2561\tau - 0.2972))e^{-0.5(0.2561\tau - 0.2972)^2} 
- 0.1262 \cos(1.75(-0.957\tau - 0.6639))e^{-0.5(-0.957\tau - 0.6639)^2} 
+ 0.0208 \cos(1.75(0.6513\tau - 0.0050))e^{-0.5(0.6513\tau - 0.0050)^2} 
- 0.3607 \cos(1.75(0.7574\tau - 0.8275))e^{-0.5(0.7574\tau - 0.8275)^2} 
+ 1.7204 \cos(1.75(-0.058\tau - 0.3235))e^{-0.5(-0.058\tau - 0.3235)^2} 
- 0.1039 \cos(1.75(-0.271\tau - 0.5232))e^{-0.5(-0.271\tau - 0.5232)^2} 
+ 1.5608 \cos(1.75(-0.3176\tau - 1.242))e^{-0.5(-0.3176\tau - 1.242)^2} 
+ 0.3663 \cos(1.75(-0.8974\tau - 0.693))e^{-0.5(-0.8974\tau - 0.693)^2} 
- 0.0041 \cos(1.75(-2.001\tau - 1.863))e^{-0.5(-2.001\tau - 1.863)^2} 
- 0.1883 \cos(1.75(0.612\tau + 0.813))e^{-0.5(0.612\tau + 0.813)^2}, \]

\[ (16) \]

\[ \tilde{Y}(\tau) = 0.0097 \cos(1.75(1.0108\tau + 0.8687))e^{-0.5(1.0108\tau + 0.8687)^2} 
+ 0.6808 \cos(1.75(-0.475\tau + 0.1533))e^{-0.5(-0.475\tau + 0.1533)^2} 
+ 1.3976 \cos(1.75(-0.107\tau + 1.5804))e^{-0.5(-0.107\tau + 1.5804)^2} 
- 0.5523 \cos(1.75(-0.066\tau - 0.3244))e^{-0.5(-0.066\tau - 0.3244)^2} 
+ 0.2293 \cos(1.75(0.3343\tau + 0.1346))e^{-0.5(0.3343\tau + 0.1346)^2} 
+ 0.8040 \cos(1.75(0.5024\tau + 0.8675))e^{-0.5(0.5024\tau + 0.8675)^2} 
+ 0.1466 \cos(1.75(-0.761\tau + 1.5040))e^{-0.5(-0.761\tau + 1.5040)^2} 
+ 0.9269 \cos(1.75(0.3945\tau - 1.1415))e^{-0.5(0.3945\tau - 1.1415)^2} 
+ 0.0374 \cos(1.75(0.9908\tau - 1.2051))e^{-0.5(0.9908\tau - 1.2051)^2} 
+ 0.1970 \cos(1.75(-0.764\tau + 0.2823))e^{-0.5(-0.764\tau + 0.2823)^2}, \]

\[ (17) \]

\[ \tilde{Z}(\tau) = 0.607 \cos(1.75(0.1112\tau + 1.3610))e^{-0.5(0.1112\tau + 1.3610)^2} 
- 0.0384 \cos(1.75(0.1343\tau + 1.1398))e^{-0.5(0.1343\tau + 1.1398)^2} 
+ 0.0150 \cos(1.75(0.7727\tau + 2.2949))e^{-0.5(0.7727\tau + 2.2949)^2} 
- 0.0024 \cos(1.75(-0.829\tau + 0.0417))e^{-0.5(-0.829\tau + 0.0417)^2} 
+ 0.3144 \cos(1.75(-0.365\tau + 1.4552))e^{-0.5(-0.365\tau + 1.4552)^2} 
- 1.1990 \cos(1.75(0.2513\tau - 1.1402))e^{-0.5(0.2513\tau - 1.1402)^2} 
+ 0.1784 \cos(1.75(0.0640\tau - 0.1835))e^{-0.5(0.0640\tau - 0.1835)^2} 
+ 1.3200 \cos(1.75(0.0134\tau - 0.5702))e^{-0.5(0.0134\tau - 0.5702)^2} 
- 0.0002 \cos(1.75(1.5840\tau - 1.8226))e^{-0.5(1.5840\tau - 1.8226)^2} 
- 0.8039 \cos(1.75(0.2124\tau + 0.2349))e^{-0.5(0.2124\tau + 0.2349)^2}. \]

\[ (18) \]

Systems (16)–(18) are implemented to solve the nonlinear SIR dengue fever system given in system (1) using the MWNN-GA-IPA and the acquired results are plotted in Figures 2–4. Figure 2 shows the set of best weights and comparison of the best obtained results with the Runge–Kutta numerical results. It is seen that the proposed and
reference results overlapped each other for $X(\tau)$, $Y(\tau)$, and $Z(\tau)$ classes to solve the nonlinear SIR dengue fever system. The plots of the AE for $X(\tau)$, $Y(\tau)$, and $Z(\tau)$ classes to solve the nonlinear SIR dengue fever system are reported in Figure 3. For the $X(\tau)$ class, $Y(\tau)$ class, and $Z(\tau)$ class, the AE best values lie about $10^{-6}$–$10^{-8}$, $10^{-3}$–$10^{-5}$, and $10^{-4}$–$10^{-6}$, and the AE mean values lie around $10^{-1}$–$10^{-2}$, $10^{-2}$–$10^{-3}$, and $10^{-3}$–$10^{-4}$, respectively. The performance
plots of the E-VAF and T.I.C indices for each class of the nonlinear SIR dengue fever system are plotted in Figure 4. For $X(τ)$ category, the best E-VAF, M.A.D, and T.I.C values lie around $10^{-10}$ to $10^{-8}$, $10^{-5}$ to $10^{-6}$, and $10^{-9}$ to $10^{-10}$. For $Y(τ)$ category, the best E-VAF, M.A.D, and T.I.C values lie around $10^{-5}$ to $10^{-6}$, $10^{-3}$ to $10^{-4}$, and $10^{-8}$ to $10^{-9}$. Similarly, for $Z(τ)$ class, the best E-VAF, M.A.D, and T.I.C values lie around $10^{-10}$ to $10^{-8}$, $10^{-5}$ to $10^{-6}$, and $10^{-9}$ to $10^{-10}$.
Figure 6: Convergence of M.A.D and E-VAF plots along with the histogram using MWNN-GA-IPA to solve each class of the nonlinear SIR dengue fever system. (a) Histogram for $X(\tau)$ class. (b) Histogram for $Y(\tau)$ class. (c) Histogram for $Z(\tau)$ class. (d) E-VAF for each class of the nonlinear SIR system. (e) Histogram for $X(\tau)$ class. (f) Histogram for $Y(\tau)$ class. (g) Histogram for $Z(\tau)$ class.
The graphical representations of the statistical trials along with the values of histograms are shown in Figures 5 and 6 for each class of nonlinear SIR dengue fever system. The convergence based on the E-VAF, M.A.D, and T.I.C operators is accomplished for independent trials to the nonlinear SIR dengue fever system. The achieved results from MWNN-GA-IPA are calculated satisfactorily based on the T.I.C, M.A.D, and E-VAF operators.

For the accurateness and precision measures, statistical studies are provided in Tables 2–4 to solve each class of the nonlinear SIR dengue fever using the operates minimum (Min), S.I range, maximum (Max), standard deviation (S.T.D), and median. The Min and Max standards show the best results and poorest results in the 100 executions. For the category X(τ), the Min, Max, median, S.I range, and S.T.D values lie around 10−3–10−4, 10−4–10−5, and 10−5–10−6.

The graph of Table 3 shows that the best E-VAF, M.A.D, and T.I.C values lie around 10−3–10−4, 10−4–10−5, and 10−5–10−6.
10\(^{-4}\)-10\(^{-5}\), 10\(^{-2}\)-10\(^{-3}\), 10\(^{-4}\)-10\(^{-5}\), 10\(^{-6}\)-10\(^{-7}\), and 10\(^{-3}\)-10\(^{-4}\), respectively. Likewise, the Min, Max, median, S.I range, and S.T.D values for the category \(Z(t)\) lie around 10\(^{-5}\)-10\(^{-11}\), 10\(^{-2}\)-10\(^{-3}\), 10\(^{-4}\)-10\(^{-7}\), 10\(^{-6}\)-10\(^{-7}\), and 10\(^{-2}\)-10\(^{-3}\), respectively. These calculated presentations found the worth and value of the proposed MWNN-GA-IPA to solve the nonlinear SIR dengue fever system. One can establish through the achieved results that the MWNN-GA-IPA is stable and precise.

The global performance of the operators [G-M.A.D], [G-T.I.C], and [G-E.VAF] for 100 trials of MWNN-GA-IPA is plotted in Table 5 to solve each category of the nonlinear SIR dengue fever system. The global-based mean [G-M.A.D], [G-T.I.C], and [G-E.VAF] values are found to be 10\(^{-2}\)-10\(^{-3}\), 10\(^{-3}\)-10\(^{-5}\), whereas the global values of the S.I Range lie in the interval 10\(^{-6}\)-10\(^{-7}\), 10\(^{-10}\)-10\(^{-11}\), and 10\(^{-6}\)-10\(^{-7}\) for each category of the nonlinear SIR dengue fever system. The close optimal outcomes acquired by the global measures approve the accurateness, correctness, and precision of MWNN-GA-IPA.

### 4. Conclusions

The current investigations are linked to design a neural network based on Morlet wavelet (MWNN) function for solving the nonlinear SIR dengue fever system based on dengue infection using the optimization procedures of global and local search approaches, i.e., GA-IPA. The nonlinear SIR dengue fever system is capable to evaluate through GA-IPA using the layer arrangement of Morlet wavelet neural networks taking 10 neurons. The overlapped results through MWNN-GA-IPA and the reference results show the good accuracy level to solve the nonlinear SIR dengue fever system based on dengue infection. The performance measures based on T.I.C, M.A.D, and E-VAF have been calculated satisfactorily. The statistical assessments for 100 independent trials using MWNN-GA-IPA in terms of minimum, S.I range, median, standard deviation, and maximum operatives further validate the worth and correctness of the proposed MWNN-GA-IPA. Furthermore, statistics analysis has been performed in the case of SIR dengue fever model based on dengue infection.

In future, the proposed MWNN-GA-IPA is proficient to solve the biological nonlinear systems, singular higher order nonlinear systems, and fluid dynamic systems.

### Data Availability

This work is not based on any data.

### Conflicts of Interest

The authors declare that they have no conflicts of interest.
delay differential model,” *Mathematical Problems in Engineering*, vol. 2020, Article ID 7359242, 9 pages, 2020.

[16] M. Umar, Z. Sabir, M. A. Zahoor Raja, and Y. G. Sánchezd, “A stochastic numerical computing heuristic of SIR nonlinear model based on dengue fever,” *Results in Physics*, vol. 19, Article ID 103585, 2020.

[17] Y. Jiang, P. Wu, J. Zeng, Y. Zhang, Y. Zhang, and S. Wang, “Multi-parameter and multi-objective optimisation of articulated monorail vehicle system dynamics using genetic algorithm,” *Vehicle System Dynamics*, vol. 58, no. 1, pp. 74–91, 2020.

[18] Z. Sabir, H. A. Wahab, M. Umar, M. G. Sakar, and M. A. Z. Raja, “Novel design of Morlet wavelet neural network for solving second order Lane-Emden equation,” *Mathematics and Computers in Simulation*, vol. 172, pp. 1–14, 2020.

[19] Z. Tao, L. Huiling, W. Wenwen, and Y. Xia, “GA-SVM based feature selection and parameter optimization in hospitalization expense modeling,” *Applied Soft Computing*, vol. 75, pp. 323–332, 2019.

[20] S. Sayed, M. Nassef, A. Badr, and I. Farag, “A nested genetic algorithm for feature selection in high-dimensional cancer microarray datasets,” *Expert Systems with Applications*, vol. 121, pp. 233–243, 2019.

[21] D. Jude Hemanth and J. Anitha, “Modified genetic algorithm approaches for classification of abnormal magnetic resonance brain tumour images,” *Applied Soft Computing*, vol. 75, pp. 21–28, 2019.

[22] M. A. Mohammed, M. K. Abd Ghani, R. I. Hamed, S. A. Mostafa, M. S. Ahmad, and D. A. Ibrahim, “Solving vehicle routing problem by using improved genetic algorithm for optimal solution,” *Journal of Computational Science*, vol. 21, pp. 255–262, 2017.

[23] X. Luo, L. Niu, and S. Zhang, “An algorithm for traffic flow prediction based on improved sarima and GA,” *KSCE Journal of Civil Engineering*, vol. 22, no. 10, pp. 4107–4115, 2018.

[24] M. Wilson, “Optimization of the radiation shielding capabilities of bismuth-borate glasses using the genetic algorithm,” *Materials Chemistry and Physics*, vol. 224, pp. 238–245, 2019.

[25] D. Jahed Armaghani, M. Hasanianpanah, A. Mahdiyar, M. Z. Abd Majid, H. Bakshandeh Amnieh, and M. M. D. Tahiri, “Airblast prediction through a hybrid genetic algorithm-ANN model,” *Neural Computing & Applications*, vol. 29, no. 9, pp. 619–629, 2018.

[26] Y. Yang, B. Yang, S. Wang, F. Liu, Y. Wang, and X. Shu, “A dynamic ant-colony genetic algorithm for cloud service composition optimization,” *International Journal of Advanced Manufacturing Technology*, vol. 102, no. 1–4, pp. 355–368, 2019.

[27] H. Motieghader, A. Najafi, B. Sadeghi, and A. Masoudi-Nejad, “A hybrid gene selection algorithm for microarray cancer classification using genetic algorithm and learning automata,” *Informatics in Medicine Unlocked*, vol. 9, pp. 246–254, 2017.

[28] H. Zhang, J. Xie, J. Ge, Z. Zhang, and B. Zong, “A hybrid adaptively genetic algorithm for task scheduling problem in the phased array radar,” *European Journal of Operational Research*, vol. 272, no. 3, pp. 868–878, 2019.

[29] M. Hassoon, M. S. Kouhi, M. Zomorodi-Moghadam, and M. Abdar, “Rule Optimization of Boosted C5.0 classification using genetic algorithm for liver disease prediction,” in *Proceedings of the 2017 International Conference on Computer and Applications (ICCA)*, September 2017.