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Construction and Analysis of a Nonstandard Computational Method for the Solution of SEIR Epidemic Model

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ABSTRACT

This paper is concerned with the numerical methods of susceptible exposed infectious recovered (SEIR) epidemic model of coronavirus disease 2019 (COVID-19). The model is explicated numerically with three numerical schemes, forward Euler, Runge-Kutta of order 4 (RK-4), and the proposed non-standard finite difference (NSFD) technique, respectively. In the epidemic model of infectious diseases, positivity is the main property of a consistent framework, since the negative value of a subpopulation is useless. The NSFD technique ends up being a more important and trustable numerical system than forward Euler and RK-4 techniques since it preserves positivity, stability, and convergence. On the contrary, forward Euler and RK-4 schemes do not hold these characteristics for some choices of step sizes. Numerical simulations confirmed the findings.

Keywords: convergence, NSFD method, SEIR model, stability

1. INTRODUCTION

In 1965, scientists identified the first human coronavirus case [\[1\]](#page-19-0). Later on, experts discovered a group of related human and animal illnesses caused by a family of viruses named coronaviruses after their crest-like appearance. People can contract seven types of coronavirus infections. COVID-19 sickness is a serious illness brought on by the SARS-CoV-2 infection [\[2\]](#page-20-0). The SARS-causing strain first appeared in southern China in 2002 and quickly spread to 29 different nations. By July 2003, 774 individuals had died and more than 8,000 had been infected [\[3\]](#page-20-1). Fever, headaches, and breathing problems, such as coughing and shortness of breath, are all brought on by coronavirus infection [\[4\]](#page-20-2).

Middle East Respiratory Syndrome (MERS) first appeared in Saudi Arabia. Many people who were exposed to it developed difficulty in breathing. It was difficult to treat the respiratory illnesses of such people

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and they could not recover without needing special care. People who have hidden illnesses, such as diabetes, cardiovascular disease, or risky development, are more likely to transfer recognized illnesses. Being educated about the condition and how the disease spreads is the greatest strategy to prevent and tone down its transmission.

On December 31, 2019, WHO received information regarding instances of pneumonia in Wuhan City, China with enigmatic explanations. On January 7, 2020, Chinese experts identified an original coronavirus as the cause and gave it the temporary designation '2019-nCoV' [\[5\]](#page-20-3). According to [\[6\]](#page-20-4), USA and Brazil are the two nations that are most affected by COVID-19. Dhandapani et. Al provided stiff, fuzzy IRD-14 day average transmission model of COVID-19 pandemic disease [\[7\]](#page-20-5). To examine the factors involved in the spread of the coronavirus pandemic in Pakistan and its potential controls, S. Ullah and M.A. Khan developed a numerical model [\[8\]](#page-20-6). Duccio and Fanelli focused on the ephemeral aspects of the coronavirus disease 2019 in China, Italy, and France for the time period January 22 - March 15, 2020 [\[9\]](#page-20-7). The focus of Xiao and Ruan [\[10\]](#page-20-8) remained on a pandemic model without a monotonic occurrence rate. The prescribed model depicts the psychological impact of specific illnesses on the local population as the number of infectious agents increase.

Since the spread of coronavirus has resulted in a pandemic, numerous mathematicians have conducted various analyses to create a model to predict its expansion $[11–29]$ $[11–29]$.

The NSFD scheme was introduced by Mickens [\[23\]](#page-22-1). The construction of the NSFD mathematical model for a two-layered differential condition was the focus of Cresson and Pierret [\[24\]](#page-22-2), who also investigated the model's various aspects, such as combination and solidness of the plan, among others. Selected mathematical models were established and tested using the RK method. Euler algorithm for requests 2 and 4 were completed. In order to account for a jungle fever model, Anguelov et al. [\[25\]](#page-22-3) constructed the NSFD plot and investigated its dependability and constituent parts. To resolve the nonlinear Riccati differential condition, Riaz et al. [\[26\]](#page-22-4) presented a very stable non-standard limited distinction (NSFD) strategy. By comparing the results to those of other mathematical operations, such as Euler and RK-4, the proposed model's accuracy and productivity were verified. A dynamic model for infection transmission components was proposed by Rafiq et al. [\[27\]](#page-22-5).

A mathematical model is developed in the current study by comparing the results with prominent mathematical schemes including the Runge-Kutta approach and the Euler scheme. In order to investigate how insect vectors affect the development of plant diseases, Rafiq et al. [\[28\]](#page-22-6) proposed a one-of-a-kind, genuinely stable, non-standard finite differentiation (NSFD) numerical arrangement. Numerical assessments were supplied and the outcomes were distinguished. In order to plan the proposed model and determine the value of the limit boundary R_0 for the model, Rafiq et al. [\[29\]](#page-22-0) developed a model consisting of nonlinear differential conditions. To analyze the susceptible exposed infected recovered (SEIR) dynamic model of COVID-19 (coronavirus) with reference to bimodal infection transmission in vulnerable populations, Ahmed et al. [\[30\]](#page-22-7) proposed a construction protecting nonstandard limited distinction plan. The suggested mathematical framework produced workable arrangements for the perplexing bi-modular SITR nonlinear model, combined actually consistent states, and reflected dynamic consistency with the perpetual feel of the model. The model's analysis revealed that it maintained some stability at focused consistent states. For the mathematical setup of the SEIR response dispersion pandemic model, [\[31\]](#page-22-8) provided two innovative NSFD schemes. Moreover, [\[32\]](#page-23-0) proposed an SIR model in a fuzzy environment. Euler, Runge Kutta of order 4 (RK-4, and the NSFD methods respectively were developed with fuzzy extensions for the solution of the model. Dayan et al. presented rumor based fuzzy model and developed an NSFD scheme for its solution [\[33\]](#page-23-1). Some researchers studied fuzzy epidemic models using NSFD schemes [\[34,](#page-23-2) [35\]](#page-23-3). Ali et al. applied the NSFD scheme for the numerical solution of a cancer disease model [\[36\]](#page-23-4). Alsallami et al. used Euler, RK, and NSFD methods to solve a real-world problem [\[37\]](#page-23-5).

The remainder of this article is structured as follows. Model formulation is presented in Section 2. Section 3 discusses the model's equilibrium states, stability, and threshold analysis. Section 4 presents numerical simulation and discussion of the results, while Section 5 displays the numerical results. The novelty of the current work lies in the fact that the proposed model has not been analyzed before using the NSFD method.

2. SEIR MODEL FOR COVID-19

Consider the model

$$
\frac{dS}{dt} = \Lambda - \mu S - \frac{\beta SI}{N}
$$
 (2.1)

$$
\frac{dE}{dt} = \frac{\beta SI}{N} - (\mu + \epsilon)E\tag{2.2}
$$

$$
\frac{dI}{dt} = \epsilon E - (\gamma + \mu + \alpha)I
$$
\n(2.3)

$$
\frac{dR}{dt} = \gamma I - \mu R \tag{2.4}
$$

2.1. Theorem

The system $(1-4)$ has a DFE point $(S_0, E_0, I_0, R_0) = \left(\frac{\Lambda}{\mu}\right)$ $\frac{\Lambda}{\mu}$, 0 , 0 , 0) and an endemic equilibrium point $(S^*, E^*, I^*, R^*) = \left(\frac{\Lambda}{\Lambda}\right)^2$ $\mu + \frac{\beta I}{N}$ N , βSI N $\frac{N}{(\mu+\epsilon)}$, ϵE

 $\frac{\epsilon E}{\gamma + \mu + \alpha}$, $\frac{\gamma I}{\mu}$ $(\frac{\mu}{\mu})$, respectively.

2.2. Basic Reproductive Number

By incorporating the next generation matrix (NGM) approach, the value of R₀ is determined. Let $X = [S, E]^t$, then $\frac{dX}{dt} = y(x) - z(x)$, where

$$
Y(x) = \left(\frac{\beta SI}{N}\right)
$$

and

$$
Z(x) = \begin{pmatrix} (\mu + \epsilon)E \\ (+\gamma + \mu + \alpha)I - \epsilon E \end{pmatrix}.
$$

Y and Z are now the Jacobeans of the $Y(x)$ and $Z(x)$ respectively at DFEP. They are listed below.

$$
Y = \begin{pmatrix} 0 & \beta \frac{\beta \Lambda}{\mu N} \\ 0 & 0 \end{pmatrix}
$$

and

School of Science
 1011 - 91

$$
Z = \begin{pmatrix} (\mu + \epsilon) & 0 \\ -\epsilon & \gamma + \mu + \alpha \end{pmatrix}.
$$

Inverse of matrix Z is

$$
Z^{-1} = \begin{pmatrix} \frac{1}{(\mu + \epsilon)} & 0 \\ \frac{\epsilon}{(\mu + \epsilon)(\gamma + \mu + \alpha)} & \frac{1}{(\gamma + \mu + \alpha)} \end{pmatrix}.
$$

Computing the product YZ^{-1} we get,

$$
YZ^{-1} = \begin{pmatrix} 0 & \beta \frac{\beta \Lambda}{\mu N} \\ 0 & 0 \end{pmatrix} \begin{pmatrix} \frac{1}{(\mu + \epsilon)} & 0 \\ \frac{\epsilon}{(\mu + \epsilon)(\gamma + \mu + \alpha)} & \frac{1}{(\gamma + \mu + \alpha)} \end{pmatrix}
$$

$$
YZ^{-1} = \begin{pmatrix} \frac{\beta \epsilon}{(\mu + \epsilon)(\gamma + \mu + \alpha)} & \frac{\beta}{(\gamma + \mu + \alpha)} \\ 0 & 0 \end{pmatrix}.
$$

Calculating the spectral radius of YZ^{-1} we get,

$$
\lambda = \frac{\beta \epsilon}{(\mu + \epsilon)(\gamma + \mu + \alpha)}.
$$

The spectral radius of YZ^{-1} is equable to R₀ which is the maximal eigenvalue of YZ^{-1} .

Therefore,

92

$$
R_0 = \frac{\beta \epsilon}{(\mu + \epsilon)(\gamma + \mu + \alpha)}.
$$
\n(2.5)

2.3. Normalized Forward Sensitivity Index (NFSI)

Chitins developed the idea of sensitivity indices by calculating the sensitivity indices of R_0 [\[38\]](#page-23-6). A parameter's normalized forward sensitivity index (NFSI) is calculated as NFSI(ζ) = $\frac{\zeta \partial}{\zeta}$ $\frac{\varsigma \theta}{\mathrm{R}_o \partial \zeta}(\mathrm{R}_0).$

The NSFIs of β , ϵ , β , γ , and α are clalculated as follows:

$$
NFSI(\beta) = \frac{\beta \partial R_0}{R_o \partial \beta} = \frac{\beta}{\frac{\beta \epsilon}{(\mu + \epsilon)(\gamma + \mu + \alpha)}} \frac{\partial(\beta \epsilon)}{(\mu + \epsilon)(\gamma + \mu + \alpha)} = \frac{(\mu + \epsilon)(\gamma + \mu + \alpha)}{\epsilon} \frac{\epsilon}{(\mu + \epsilon)(\gamma + \mu + \alpha)} = 1.
$$

Scientific Inquiry and Review

Similarly,

$$
NFSI(\epsilon) = \frac{\epsilon}{\frac{\beta \epsilon}{(\mu + \epsilon)(\gamma + \mu + \alpha)}} \frac{\partial \beta \epsilon}{\partial \epsilon(\mu + \epsilon)(\gamma + \mu + \alpha)} = \frac{\mu}{(\mu + \epsilon)}.
$$

$$
NFSI(\mu) = -\frac{\gamma}{(\mu + \epsilon)(\gamma + \mu + \alpha)}, \text{ and}
$$

$$
NFSI(\alpha) = \frac{\alpha}{(\gamma + \mu + \alpha)}.
$$

These findings unambiguously demonstrate that β is the most sensitive parameter.

2.4. Stability Analysis

The Jacobean of the system (1-4) is

$$
J = \begin{pmatrix} \mu - \beta \frac{I}{N} & 0 & -\beta \frac{S}{N} & 0 \\ \beta \frac{I}{N} & -(\mu + \epsilon) & \beta \frac{S}{N} & 0 \\ 0 & -\epsilon & -(\gamma + \mu + \alpha) & 0 \\ 0 & 0 & \gamma & -\mu \end{pmatrix}
$$

$$
J(\xi_0) = \begin{pmatrix} \mu & 0 & -\beta & 0 \\ \beta \frac{I}{N} & -(\mu + \epsilon) & \beta & 0 \\ 0 & -\epsilon & -(\gamma + \mu + \alpha) & 0 \\ 0 & 0 & \gamma & -\mu \end{pmatrix}
$$

If the absolute eigenvalues of $J(\xi_0)$ are smaller than unity, that is, $|\lambda_i|$ < 1 , $i = 1, 2, 3$. The numerical scheme converges in all cases. The eigenvalue from the aforementioned Jacobean matrix were extracted, where $\lambda_1 = \lambda_4 =$ $-\mu$, $\lambda_2 = -(\mu + \epsilon)$, and $\lambda_3 = -(\gamma + \mu + \alpha)$.

2.4.1. Stability at Endemic Equilibrium Point. Jacobean matrix at the EE point is

$$
J(E^*) = \begin{pmatrix} \mu - \beta \frac{I^*}{N} & 0 & -\beta \frac{S^*}{N} & 0 \\ \beta \frac{I^*}{N} & -(\mu + \epsilon) & \beta \frac{S^*}{N} & 0 \\ 0 & -\epsilon & -(\gamma + \mu + \alpha) & 0 \\ 0 & 0 & \gamma & -\mu \end{pmatrix},
$$

where

School of Science
 193 UMT 93

$$
S^* = \frac{\Lambda}{\mu + \beta \frac{I}{N}} \text{ and } I^* = \frac{\epsilon E}{(\gamma + \mu + \alpha)}.
$$

Using the MATLAB database, the biggest eigenvalues of $J(E^*)$ were plotted. At the endemic equilibrium point, the Jacobean spectral radius has a maximum value that is smaller than unity, as is evident from the preceding figure, proving the intended assertion.

3. NUMERICAL MODELING OF SEIR MODEL FOR COVID-19

3.1. Forward Euler Method

$$
S^{n+1} = S^n + h[\Lambda - \mu S^n - S^n \frac{\beta I^n}{N}]
$$
\n(3.1)

$$
E^{n+1} = E^n + h \left[\frac{\beta S^n I^n}{N} - (\mu + \epsilon) E^n \right]
$$
\n(3.2)

$$
I^{n+1} = I^n + h \left[\epsilon E^n - (\gamma + \mu + \alpha)I^n \right] \tag{3.3}
$$

$$
R^{n+1} = R^n + h \left[\gamma I^n - \mu R^n \right] \tag{3.4}
$$

3.2. Fourth Order Runge-Kutta (RK-4) Scheme

3.2.1. Step 1

$$
k_1 = S^n + h[\Lambda - \mu S^n - S^n \frac{\beta I^n}{N}]
$$
\n(3.5)

$$
m_1 = E^n + h\left[\frac{\beta S^n I^n}{N} - (\mu + \epsilon) E^n\right]
$$
\n(3.6)

$$
n_1 = I^n + h \left[\epsilon E^n - (\gamma + \mu + \alpha)I^n \right] \tag{3.7}
$$

$$
p_1 = R^n + h \left[\gamma I^n - \mu R^n \right] \tag{3.8}
$$

3.2.2. Step 2

$$
k_2 = h[\Lambda - \mu(S^n + \frac{k_1}{2}) - \beta \frac{(S^n + \frac{k_1}{2})(I^n + \frac{n_1}{2})}{N}]
$$
(3.9)

$$
m_2 = h\left[\beta \frac{(S^n + \frac{k_1}{2})(I^n + \frac{n_1}{2})}{N} - (\mu + \epsilon)(E^n + \frac{m_1}{2})\right]
$$
(3.10)

$$
n_2 = h \left[\epsilon (E^n + \frac{m_1}{2}) - (\gamma + \mu + \alpha)(I^n + \frac{n_1}{2}) \right]
$$
 (3.11)

Scientific Inquiry and Review

94 — **SIR**

$$
p_2 = h \left[\gamma (I^n + \frac{n_1}{2}) - \mu (R^n + \frac{p_1}{2}) \right]
$$
 (3.12)

3.2.3. Step 3

$$
k_3 = h[\Lambda - \mu(S^n + \frac{k_2}{2}) - \beta \frac{(S^n + \frac{k_2}{2})(I^n + \frac{n_2}{2})}{N}]
$$
 (3.13)

$$
m_3 = h\left[\beta \frac{(S^n + \frac{k_2}{2})(I^n + \frac{n_2}{2})}{N} - (\mu + \epsilon)(E^n + \frac{m_2}{2})\right]
$$
(3.14)

$$
n_3 = h \left[\epsilon (E^n + \frac{m_2}{2}) - (\gamma + \mu + \alpha)(I^n + \frac{n_2}{2}) \right]
$$
 (3.15)

$$
p_3 = h \left[\gamma \left(I^n + \frac{n_2}{2} \right) - \mu \left(R^n + \frac{p_2}{2} \right) \right]
$$
 (3.16)

3.2.4. Step 4

$$
k_4 = h[\Lambda - \mu(S^n + k_3) - \beta \frac{(S^n + k_3)(I^n + n_3)}{N}]
$$
\n(3.17)

$$
m_4 = h[\beta \frac{(S^n + k_3)(I^n + n_3)}{N} - (\mu + \epsilon)(E^n + m_3)] \tag{3.18}
$$

$$
n_4 = h \left[\epsilon (E^n + m_3) - (\gamma + \mu + \alpha)(I^n + n_3) \right]
$$
 (3.19)

$$
p_4 = h \left[\gamma (I^n + n_3) - \mu (R^n + p_3) \right]
$$
 (3.20)

3.2.5. Step 5

$$
S^{n+1} = S^n + \frac{1}{6} \left[k_1 + 2k_2 + 2k_3 + k_4 \right] \tag{3.21}
$$

$$
E^{n+1} = E^n + \frac{1}{6} [m_1 + 2m_2 + 2m_3 + m_4]
$$
 (3.22)

$$
I^{n+1} = I^n + \frac{1}{6} [n_1 + 2n_2 + 2n_3 + n_4]
$$
 (3.23)

$$
R^{n+1} = R^n + \frac{1}{6} [p_1 + 2p_2 + 2p_3 + p_4]
$$
 (3.24)

3.3. Non-Standard Finite Difference (NSFD) Scheme

To develop an explicit NSFD scheme, consider the above system as

95 School of Science **1967** School of Science **1968** UMT 95

$$
\frac{df(t)}{dS} = \frac{f(t+h) - f(t)}{h} + 0(h) \qquad h \to 0.
$$

 S^n , E^n , I^n , and Rⁿ are approximations. Here, h is the time step size.

$$
\frac{S^{n+1} - S^n}{h} = \Lambda - \mu S^{n+1} - S^{n+1} \frac{\beta I^n}{N}
$$
 (3.25)

$$
S^{n+1} = \frac{S^n + h\Lambda}{1 + h\mu + h\frac{\beta I^n}{N}}
$$
(3.26)

$$
E^{n+1} = \frac{E^n + h \frac{\beta S^n I^n}{N}}{1 + h(\mu + \epsilon)}
$$
(3.27)

$$
I^{n+1} = \frac{I^n + h \epsilon E^n}{1 + h \left(\gamma + \mu + \alpha\right)}\tag{3.28}
$$

$$
R^{n+1} = \frac{R^n + h\,\gamma I^n}{1 + h\mu} \tag{3.29}
$$

3.4. Stability Analysis of the NSFD Technique

The NSFD scheme of the SEIR model is carried out in this section at DFEP $(S_0, E_0, I_0, R_0) = (\frac{\Lambda}{\mu})$ $\frac{\pi}{\mu}$, 0, 0, 0). Consider the **Eq.** (3.25) to (3.28),

$$
F = S^{n+1} = \frac{S + h\Lambda}{1 + h\mu + h\frac{\beta I}{N}}
$$
(3.30)

$$
G = E^{n+1} = \frac{E + h \frac{\beta S I}{N}}{1 + h(\mu + \epsilon)}
$$
(3.31)

$$
H = I^{n+1} = \frac{I + h \epsilon E}{1 + h (\gamma + \mu + \alpha)}
$$
(3.32)

$$
K = R^{n+1} = \frac{R + h\gamma I}{1 + h\mu}
$$
 (3.33)

Jacobean matrix of the **Eq.** (**3.30**) to (**3.33**) at the DFEP is as follows:

.

$$
J(\xi_0) = \begin{pmatrix} \frac{1}{1+h\mu} & 0 & -\frac{\left(\frac{\Lambda}{\mu} + h\Lambda\right)\left(\frac{h\beta}{N}\right)}{\left[1+h\mu\right]^2} & 0\\ 0 & \frac{1}{1+h(\mu+\epsilon)} & \frac{h\beta\Lambda}{\mu N} & 0\\ 0 & \frac{h\epsilon}{1+h(\gamma+\mu+\alpha)} & \frac{1}{1+h(\gamma+\mu+\alpha)} & 0\\ 0 & 0 & \frac{h\gamma}{1+h\mu} & \frac{1}{1+h\mu} \end{pmatrix}
$$

The eigenvalues of the aforementioned Jacobean matrix are $\lambda_1 = \frac{1}{1+i}$ $\frac{1}{1+h\mu}$ < 1 and $\lambda_2 = \frac{1}{1+t}$ $\frac{1}{1+h\mu}$ < 1. The following matrix is used to compute the remaining eigenvalues.

$$
J(\xi_0) = \begin{pmatrix} \frac{1}{1 + h(\mu + \epsilon)} & \frac{h\beta\Lambda}{\mu N} \\ \frac{h\epsilon}{1 + h(\gamma + \mu + \alpha)} & \frac{1}{1 + h(\gamma + \mu + \alpha)} \end{pmatrix}.
$$

The other eigenvalues are quite complicated to be calculated algebraically. Therefore, they have been calculated numerically (shown in Figure 1).

The spectral radius of the Jacobean matrix at the DFE point has a maximum value that is less than unity, as is evident from the following figure, proving the intended assertion.

School of Science
 1977 School of Science
 1978 School of Sc

4. NUMERICAL SIMULATION OF THE SEIR MODEL

The findings are discussed in detail in this section. Additionally, Euler, RK-4, and NSFD methods in the model are compared in terms of their use.

Figure 3. Infected Population using Euler Scheme at h = 0.1

Figure 5. Infected Population using Euler Scheme at $h = 10$

99 UMT 99 Volume 7 Issue 1, 2023

Figure 6. Infected Population using RK-4 Method at h = 0.01

Figure 7. Infected Population using RK-4 Method at h =0.1

100

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Figure 8. Infected Population using RK-4 Method at h =1

Figure 9. Infected Population using RK-4 Method at $h = 10$

1011 School of Science **1011** School of Science **1011** 301

Figure 10. Infected Population using NSFD Scheme at h = 0.01

Figure 11. Infected Population using NSFD Scheme at h = 0.1

102—**NIR**

Figure 12. Infected Population using NSFD Scheme at h =1

Figure 13. Infected Population using NSFD Scheme at h =10

Figure 14. Comparison of Euler, RK-4, and NSFD Scheme at h=0.01

Figure 15. Comparison of Euler, RK-4, and NSFD Scheme at h=0.1

104⁻

Scientific Inquiry and Review

Figure 16. Comparison of Euler, RK-4, and NSFD Scheme at h=1

Figure 17. Comparison of Euler, RK-4, and NSFD Scheme at h=10

105 School of Science **105** School of Science **105**

The behavior of the SEIR model is presented in the above graphs. Figure 2 and Figure 3 show that the Euler technique behaves favorably and converges at small step sizes ($h = 0.01$ and $h = 0.1$). In Figure 4 and Figure 5, the Euler method shows divergence at slightly large step sizes ($h = 1$ and $h =$ 10). In Figure 6 and Figure 7, the RK-4 method shows convergence at small step sizes ($h = 0.01$ and $h = 0.1$). The method shows divergence as the step size increases, as shown in Figure 8 and Figure 9, respectively. InFigure 10-13, the NSFD method converges to the same equilibrium points at different step sizes. In Figure 14-17, a comparison of Euler's method, RK-4 method, and NSFD method is depicted. It is clear from the above graphs that all models show similar behaviors at small step sizes and converge to the same equilibrium points. The Euler and RK-4 methods show divergence and negative behavior as step size increases, while the NSFD method converges to the same point. Graphical depiction of behaviors demonstrates that the Euler and RK-4 methods only provide convergence solutions for small step sizes and remain inconclusive for large step sizes. The NSFD scheme, on the other hand, exhibits good behavior and also provides a convergent solution for step sizes that are extremely large.

5. CONCLUSION

The mathematical study of the SEIR model for the spread of COVID-19 was carried out in this research. For this purpose, the reproduction number R_0 , sensitivity of R_0 , and equilibrium points of the model were determined. It was shown that both equilibrium points have similar stability properties. In order to solve the investigated model, Euler, RK-4, and NSFD algorithms were deployed. Different time step sizes were used for numerical experiments at DFE and EE locations. The collected results were examined and compared. From the said examination, the current study concludes that the suggested method yields findings that converge to true stable states for any time step size. However, the Euler and RK-4 methods do not hold for large time step sizes. Moreover, the NSFD method is bounded, dynamically consistent, and preserves the positivity of the solution which are important requirements when modeling a prevalent disease.

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Scientific Inquiry and Review

108—**NER**

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