# On the analysis of fractional diabetes model with exponential law 

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#### Abstract

In this work, we study the diabetes model and its complications with the Caputo-Fabrizio fractional derivative. A deterministic mathematical model pertaining to the fractional derivative of the diabetes mellitus is discussed. The analytical solution of the diabetes model is derived by exerting the homotopy analysis method, the Laplace transform and the Padé approximation. Moreover, existence and uniqueness of the solution are examined by making use of fixed point theory and the Picard-Lindelof approach. Ultimately, for illustrating the obtained results some numerical simulations are performed.


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## 1 Introduction

Nowadays, diabetes is world-wide a quiet epidemic heavily enlarging the charge of noncommunicable diseases and generally stimulated by reducing the levels of activity and enhancing pervasiveness of obesity. Mainly, two types of diabetes are studied: Type 1 diabetes, too familiar as insulin dependent diabetes mellitus, attacking people below the age of 40 and representing 10 to 15 percent of the diabetic population. Then, Type II diabetes, previously recognized as non-insulin dependent diabetes mellitus, describing the majority ( 85 to 90 percent). Although in all age groups with the spreading epidemic of obesity, it is anticipated that in 10 year's time, there will be additional children having Type II rather than having Type I; see [1-8]. In recent years many authors have studied different mathematical models to describe diabetes and related issues such as Boutayeb et al. [9], Mahataa et al. [10], Pandit et al. [11], Makroglou et al. [12] and others.

It has been demonstrated by many scientists and mathematicians that fractional extensions of mathematical models of integer order represent the natural fact in a very systematic way such as in the approaches of Caputo [13], Podlubny [14], Miller and Ross [15], Baleanu et al. [16], Singh et al. [17, 18], Kumar et al. [19] and others. Lately, Caputo and Fabrizio [20] suggested an innovative idea of non-integer order derivative possessing non-singular kernel, and the authors in [21] described the additional characteristics. In the present article, we employ a newly found arbitrary order derivative to the diabetes model. The principal aim of the present article is to use a novel non-integer order derivative to
study the diabetes model and present the details of uniqueness and existence of the solution of the diabetes model with the help of a fixed point theorem and the Picard-Lindelof approach.
In this work, we study the fractional diabetes model by applying the homotopy analysis transform method (HATM). The HATM is a new and well organized mixture of the HAM [22-24] and standard Laplace transform technique [25-29].
The development of the present article is given in the following sequential manner: In Sect. 2, the CF derivative of arbitrary order is explored. In Sect. 3, the diabetes model and exact solution associated to the newly CF arbitrary order derivative is described. We also discuss special solutions of the model in this portion. By exerting the fixed point theorem and the Picard-Lindelof approach, uniqueness and existence of the solutions of the system are analyzed in Sect. 4. Section 5 contains a stability analysis of an iterative scheme. In Sect. 6, the numerical simulations are graphically shown. Lately in Sect. 7, the conclusions are discussed.

## 2 Preliminaries

Definition 2.1 Let $l(\eta) \in H^{1}(x, y), y>x, \sigma \in[0,1]$ then the CF arbitrary order derivative [20] is given as

$$
\begin{equation*}
D_{\eta}^{\sigma}(l(\eta))=\frac{M(\sigma)}{1-\sigma} \int_{x}^{\eta} l^{\prime}(w) \exp \left[-\sigma \frac{\eta-w}{1-\sigma}\right] d w \tag{1}
\end{equation*}
$$

In Eq. (1) $M(\sigma)$ denotes a normalization of the function fulfilling the property $M(0)=$ $1=M(1)$ and its details can be found in a paper authored by Caputo and Fabrizio [20]. This derivative can be presented as follows, if $l \notin H^{1}(a, b)$ :

$$
\begin{equation*}
D_{\eta}^{\sigma}(g(\eta))=\frac{\sigma M(\sigma)}{1-\sigma} \int_{x}^{\eta}(l(\eta)-l(w)) \exp \left[-\sigma \frac{\eta-w}{1-\sigma}\right] d w . \tag{2}
\end{equation*}
$$

Note 1 If $\beta=\frac{1-\sigma}{\sigma} \in[0, \infty), \sigma=\frac{1}{1+\beta} \in[0,1]$, then Eq. (2) converts to the following form:

$$
\begin{equation*}
D_{\eta}^{\sigma}(l(\eta))=\frac{N(\beta)}{\beta} \int_{x}^{\eta} l^{\prime}(w) \exp \left[-\frac{\eta-w}{\beta}\right] d w, \quad N(0)=1=N(\infty) \tag{3}
\end{equation*}
$$

Furthermore,

$$
\begin{equation*}
\lim _{\beta \rightarrow 0} \frac{1}{\sigma} \exp \left[-\frac{\eta-w}{\beta}\right]=\delta(w-\eta) \tag{4}
\end{equation*}
$$

The corresponding integral was described by [21].

Definition 2.2 Let $0<\sigma<1$, then the non-integer order integral of $l(\eta)$ is presented as

$$
\begin{equation*}
I_{\eta}^{\sigma}(l(\eta))=\frac{2(1-\sigma)}{(2-\sigma) M(\sigma)} l(\eta)+\frac{2 \sigma}{(2-\sigma) M(\sigma)} \int_{0}^{\eta} l(s) d s, \quad \eta \geq 0 . \tag{5}
\end{equation*}
$$

By making use of Eq. (5), we have

$$
\begin{equation*}
\frac{2(1-\sigma)}{(2-\sigma) M(\sigma)}+\frac{2 \sigma}{(2-\sigma) M(\sigma)}=1 \tag{6}
\end{equation*}
$$

In view of the result (6), the authors of [21] introduced the modified CF derivative of order $0<\sigma<1$ expressed as

$$
\begin{equation*}
D_{\eta}^{\sigma}(l(t))=\frac{1}{1-\sigma} \int_{x}^{\eta} l^{\prime}(w) \exp \left[-\sigma \frac{\eta-w}{1-\sigma}\right] d w . \tag{7}
\end{equation*}
$$

## 3 HATM for fractional diabetes model and its complication

The classical diabetes model and its complications [1] are given by

$$
\begin{align*}
& \frac{d D}{d \eta}=I-(\lambda+\mu) D+\gamma C \\
& \frac{d C}{d \eta}=I+\lambda D-(\gamma+\mu+v+\delta) C \tag{8}
\end{align*}
$$

where $D$ indicates the no. of diabetics with no complications at time $\eta$. $C$ denotes the no. of diabetics having complications. I represents the occurrence of diabetes mellitus.
$\mu$ denotes the rate of natural mortality, $\lambda$ indicates the probability of a diabetic person spreading a complication, $\gamma$ shows the rate of healing complications, $v$ stands for the rate at which diabetic patients having complication converts critically disabled and $\delta$ denotes the mortality rate because of complications. $N(\eta)=C(\eta)+D(\eta)$ indicates the size of diabetics at the time $\eta$. Taking $N(\eta)=C(\eta)+D(\eta)$, we get

$$
\begin{align*}
& \frac{d C}{d \eta}=-(\lambda+\theta) C+\lambda N, \quad \eta>0,  \tag{9}\\
& \frac{d N}{d \eta}=I-(v+\delta) C-\mu N .
\end{align*}
$$

In Eq. (9), $\theta=\gamma+\mu+\nu+\delta$ with initial conditions

$$
\begin{equation*}
C(0)=C_{0}, \quad N(0)=N_{0} . \tag{10}
\end{equation*}
$$

The model (9) is not able to contain the entire memory effect of the biological system and the spreading of diabetes. To involve these discussed effects into the mathematical representation of the diabetes model, we modify the equation by changing the first order time derivative to the novel propounded CF arbitrary order derivative [20] as represented by

$$
\begin{align*}
& { }_{0}^{\mathrm{CF}} D_{\eta}^{\sigma} C=-(\lambda+\theta) C+\lambda N, \quad \eta>0,  \tag{11}\\
& { }_{0}^{\mathrm{CF}} D_{\eta}^{\sigma} N=I-(v+\delta) C-\mu N,
\end{align*}
$$

with initial conditions given in Eq. (10).
Solution. By performing the Laplace transform to Eq. (11),

$$
\begin{align*}
& \frac{s L[C]-C(0)}{s+\sigma(1-s)}=L[-(\lambda+\theta) C+\lambda N] \\
& \frac{s L[N]-N(0)}{s+\sigma(1-s)}=L[I-(v+\delta) C-\mu N] \tag{12}
\end{align*}
$$

On simplifying, we get

$$
\begin{align*}
& L[C]-\frac{C_{0}}{s}-\frac{s+\sigma(1-s)}{s} L[-(\lambda+\theta) C+\lambda N]=0, \\
& L[N]-\frac{N_{0}}{s}-\frac{s+\sigma(1-s)}{s} L[I-(v+\delta) C-\mu N]=0 . \tag{13}
\end{align*}
$$

The non-linear operator is defined as

$$
\begin{gather*}
P_{1}[\psi(\eta ; q)]=L[\psi(\eta ; q)]-\frac{C_{0}}{s}-\frac{s+\sigma(1-s)}{s} L[-(\lambda+\theta) \psi(\eta ; q)+\lambda \varphi(\eta ; q)]=0, \\
P_{2}[\varphi(\eta ; q)]=L[\varphi(\eta ; q)]-\frac{N_{0}}{s}-\frac{s+\sigma(1-s)}{s} L[I-(v+\delta) \psi(\eta ; q)-\mu \varphi(\eta ; q)]=0 \tag{14}
\end{gather*}
$$

and thus we have

$$
R_{1, k}\left(\vec{C}_{k-1}\right)=L\left[C_{k-1}(\eta)\right]-\frac{C_{0}}{s}\left(1-\chi_{k}\right)-\frac{s+\sigma(1-s)}{s} L\left[-(\lambda+\theta) C_{k-1}(\eta ; q)+\lambda N_{k-1}(\eta ; q)\right]
$$

and

$$
\begin{align*}
R_{2, k}\left[\vec{N}_{k-1}\right]= & L\left[N_{k-1}(\eta)\right]-\frac{N_{0}}{s}\left(1-\chi_{k}\right) \\
& -\frac{s+\sigma(1-s)}{s} L\left[I-(v+\delta) C_{k-1}(\eta ; q)-\mu N_{k-1}(\eta ; q)\right] \tag{15}
\end{align*}
$$

The $m$ th order deformation equation is presented as

$$
L\left[C_{k}(\eta)-\chi_{k} C_{k-1}(\eta)\right]=\hbar R_{1, k}\left(\vec{C}_{k-1}\right),
$$

and

$$
\begin{equation*}
L\left[N_{k}(\eta)-\chi_{k} N_{k-1}(\eta)\right]=\hbar R_{2, k}\left(\vec{N}_{k-1}\right) . \tag{16}
\end{equation*}
$$

On employing the inversion of the Laplace transform, we get

$$
C_{k}(\eta)=\chi_{k} C_{k-1}(\eta)+\hbar L^{-1}\left[R_{1, k}\left(\vec{C}_{k-1}\right)\right]
$$

and

$$
\begin{equation*}
N_{k}(\eta)=\chi_{k} N_{k-1}(\eta)+\hbar L^{-1}\left[R_{2, k}\left(\overleftarrow{N}_{k-1}\right)\right] \tag{17}
\end{equation*}
$$

On solving the above equations, for $k=1,2,3, \ldots$, we get

$$
C_{1}(\eta)=-\hbar\left[-(\lambda+\theta) C_{0}+\lambda N_{0}\right][1+\sigma(\eta-1)]
$$

i.e. $C_{1}(\eta)=-\hbar \alpha[1+\sigma(\eta-1)]$, where $\alpha=-q C_{0}+\lambda N_{0}$.

$$
N_{1}(\eta)=-\hbar\left[I-(v+\delta) C_{0}-\mu N_{0}\right][1+\sigma(\eta-1)]
$$

i.e. $N_{1}(\eta)=-\hbar \beta[1+\sigma(\eta-1)]$, where $\beta=I-m C_{0}-\mu N_{0}$, and so on.

Consequently, the solution of Eq. (11) is given as

$$
C(\eta)=C_{0}+C_{1}+C_{2}+\cdots
$$

or

$$
\begin{equation*}
C(\eta)=C_{0}-\hbar \alpha[1+\sigma(\eta-1)]+\cdots \tag{18}
\end{equation*}
$$

and

$$
N(\eta)=N_{0}+N_{1}+N_{2}+\cdots
$$

or

$$
\begin{equation*}
N(\eta)=N_{0}-\hbar \beta[1+\sigma(\eta-1)]+\cdots, \tag{19}
\end{equation*}
$$

where $q=\lambda+\theta, m=v+\delta$ and $\theta=\gamma+\mu+\nu+\delta$.

## 4 Existence and uniqueness analysis

The fractional diabetic model (11) can be reduced to an integral equation of Volterra type when the integral related to the CF derivative is employed. We recall that the CaputoFabrizio non-integer order integral of $l(\eta)$ is the average of $l(\eta)$ and its Riemann-Liouville integral of fractional order.
Now using the integral operator of fractional order introduced by Losada and Nieto [21] on the system (11), we get

$$
\begin{align*}
C(\eta)-g_{1}(\eta)= & \frac{2(1-\sigma)}{(2-\sigma) M(\sigma)}\{-(\lambda+\theta) C(\eta)+\lambda N(\eta)\} \\
& +\frac{2 \sigma}{(2-\sigma) M(\sigma)} \int_{0}^{\eta}\{-(\lambda+\theta) C(s)+\lambda N(s)\} d s,  \tag{20}\\
N(\eta)-g_{2}(\eta)= & \frac{2(1-\sigma)}{(2-\sigma) M(\sigma)}\{I-(v+\delta) C(\eta)-\mu N(\eta)\} \\
& +\frac{2 \sigma}{(2-\sigma) M(\sigma)} \int_{0}^{\eta}\{I-(v+\delta) C(s)-\mu N(s)\} d s .
\end{align*}
$$

A possibility of reducing Eq. (20) to the iterative approach is presented as follows:

$$
\begin{equation*}
C_{0}(\eta)=g_{1}(\eta), \quad D_{0}(\eta)=g_{2}(\eta) . \tag{21}
\end{equation*}
$$

Now, we get the subsequent iterative algorithm

$$
\begin{align*}
C_{(n+1)}(\eta)= & \frac{2(1-\sigma)}{(2-\sigma) M(\sigma)}\left\{-(\lambda+\theta) C_{n}(\eta)+\lambda N_{n}(\eta)\right\} \\
& +\frac{2 \sigma}{(2-\sigma) M(\sigma)} \int_{0}^{\eta}\left\{-(\lambda+\theta) C_{n}(s)+\lambda N_{n}(s)\right\} d s,  \tag{22}\\
N_{(n+1)}(\eta)= & \frac{2(1-\sigma)}{(2-\sigma) M(\sigma)}\left\{I-(v+\delta) C_{n}(\eta)-\mu N_{n}(\eta)\right\} \\
& +\frac{2 \sigma}{(2-\sigma) M(\sigma)} \int_{0}^{\eta}\left\{I-(v+\delta) C_{n}(s)-\mu N_{n}(s)\right\} d s .
\end{align*}
$$

Here we assume that we can get the exact solution by taking the limit as $n$ tends to infinity.

### 4.1 Existence of the solution by Picard-Lindelof approach

Theorem 1 We discuss the existence of the solution by applying the Picard-Lindelof approach.

Proof We express the operators as

$$
\begin{align*}
& k_{1}(\eta, C)=-(\lambda+\theta) C+\lambda N,  \tag{23}\\
& k_{2}(\eta, N)=I-(v+\delta) C-\mu N,
\end{align*}
$$

where $k_{1}(\eta, C)$ and $k_{2}(\eta, N)$ are contractions with respect to $C$ and $N$ for the first and second functions, respectively.

Let

$$
\begin{equation*}
A_{1}=\sup _{C_{a, b_{1}}}\left\|K_{1}(\eta, C)\right\|, \quad A_{2}=\sup _{C_{a, b_{2}}}\left\|K_{2}(\eta, N)\right\|, \tag{24}
\end{equation*}
$$

where

$$
\begin{align*}
& C_{a, b_{1}}=|\eta-a, \eta+a| \times\left[C-b_{1}, C+b_{1}\right]=A_{1} \times B_{1}, \\
& C_{a, b_{2}}=|\eta-a, \eta+a| \times\left[N-b_{2,} N+b_{2}\right]=A_{1} \times B_{2} . \tag{25}
\end{align*}
$$

On consideration of the Picard operator, we have

$$
\begin{equation*}
\phi: C\left(A_{1}, B_{1}, B_{2}\right) \rightarrow C\left(A_{1}, B_{1}, B_{2}\right), \tag{26}
\end{equation*}
$$

given as follows:

$$
\begin{equation*}
\phi \xi(\eta)=\xi_{0}(\eta)+\Delta(\eta, \xi(\eta)) \frac{2(1-\sigma)}{(2-\sigma) M(\sigma)}+\frac{2 \sigma}{(2-\sigma) M(\sigma)} \int_{0}^{\eta} \Delta(s, \xi(s)) d s \tag{27}
\end{equation*}
$$

where $\xi(\eta)=\{C(\eta), N(\eta)\}, \xi_{0}(\eta)=\left\{g_{1}, g_{2}\right\}$ and $\Delta(\eta, \xi(\eta))=\left\{k_{1}\left(\eta, C(\eta), k_{2}(\eta, N(\eta)\}\right.\right.$.
Next we suppose the solution of the fractional diabetic model are bounded within a time period,

$$
\begin{align*}
&\|\xi(\eta)\|_{\infty} \leq \max \left\{B_{1}, B_{2}\right\},  \tag{28}\\
&\left\|\xi(\eta)-\xi_{0}(\eta)\right\|=\left\|\Delta(\eta, \xi(\eta)) \frac{2(1-\sigma)}{(2-\sigma) M(\sigma)}+\frac{2 \sigma}{(2-\sigma) M(\sigma)} \int_{0}^{\eta} \Delta(s, \xi(s)) d s\right\| \\
& \leq \frac{2(1-\sigma)}{(2-\sigma) M(\sigma)}\|\Delta(\eta, \xi(\eta))\|+\frac{2 \sigma}{(2-\sigma) M(\sigma)} \int_{0}^{\eta}\|\Delta(s, \xi(s))\| d s \\
& \leq\left(\frac{2(1-\sigma)}{(2-\sigma) M(\sigma)}+\frac{2 \sigma \eta_{0}}{(2-\sigma) M(\sigma)}\right) \max \left\{B_{1}, B_{2}\right\} \leq \rho B \leq \alpha, \tag{29}
\end{align*}
$$

where we demand that

$$
\begin{equation*}
\rho<\frac{\alpha}{B} . \tag{30}
\end{equation*}
$$

By employing the fixed point theorem pertaining to Banach space along with the metric, we obtain

$$
\begin{equation*}
\left\|\phi \xi_{1}-\phi \xi_{2}\right\|_{\infty}=\sup _{\eta \rightarrow A}\left|\xi_{1}-\xi_{2}\right| . \tag{31}
\end{equation*}
$$

Now we have

$$
\begin{align*}
\left\|\phi \xi_{1}-\phi \xi_{2}\right\|= & \|\left\{\Delta\left(\eta, \xi_{1}(\eta)\right)-\Delta\left(\eta, \xi_{2}(\eta)\right)\right\} \frac{2(1-\sigma)}{(2-\sigma) M(\sigma)} \\
& +\frac{2 \sigma}{(2-\sigma) M(\sigma)} \int_{0}^{\eta}\left\{\Delta\left(s, \xi_{1}(s)\right)-\Delta\left(s, \xi_{2}(s)\right)\right\} d s \| \\
\leq & \frac{2(1-\sigma)}{(2-\sigma) M(\sigma)}\left\|\Delta\left(\eta, \xi_{1}(\eta)\right)-\Delta\left(\eta, \xi_{2}(\eta)\right)\right\| \\
& +\frac{2 \sigma}{(2-\sigma) M(\sigma)} \int_{0}^{\eta}\left\|\Delta\left(s, \xi_{1}(s)\right)-\Delta\left(s, \xi_{2}(s)\right)\right\| d s \\
\leq & \frac{2(1-\sigma)}{(2-\sigma) M(\sigma)} \beta\left\|\xi_{1}(\eta)-\xi_{2}(\eta)\right\|+\frac{2 \sigma \beta}{(2-\sigma) M(\sigma)} \int_{0}^{\eta}\left\|\xi_{1}(s)-\xi_{2}(s)\right\| d s \\
\leq & \left\{\frac{2(1-\sigma) \beta}{(2-\sigma) M(\sigma)}+\frac{2 \sigma \beta \eta_{0}}{(2-\sigma) M(\sigma)}\right\}\left\|\xi_{1}(\eta)-\xi_{2}(\eta)\right\| \\
\leq & \rho \beta\left\|\xi_{1}(\eta)-\xi_{2}(\eta)\right\|, \tag{32}
\end{align*}
$$

with $\beta<1$. Since $\xi$ is a contraction, we have $\rho \beta<1$, hence the defined operator $\phi$ is also a contraction. Therefore, the diabetic model involving CF derivative given in Eq. (11) has a unique set of solutions.

## 5 Stability analysis of iterative scheme

Let $(\wp,\|\cdot\|)$ indicate a Banach space and $W$ represent a self-map of $\wp$. Let $h_{n+1}=\varphi\left(W, h_{n}\right)$ be a specific iterative scheme. Let us consider $\aleph(W)$ to be the fixed point set of $W$ possessing at least one element and let $h_{n}$ be convergent and converge to $z \in \aleph(W)$. If $\left\{h_{n}\right\} \subseteq \wp$ and we define $a_{n}=\left\|h_{n+1}-\varphi\left(W, h_{n}\right)\right\|$, and if $\operatorname{Lim}_{n \rightarrow \infty} a^{n}=0$ yields the result $\operatorname{Lim}_{n \rightarrow \infty} h^{n}=z$, then the iteration algorithm $h_{n+1}=\varphi\left(W, h_{n}\right)$ is termed $W$-stable.

Theorem 2 ([30]) Let $(\wp,\|\cdot\|)$ be a Banach space and W be a self-map of the Banach


$$
\left\|W_{x}-W_{y}\right\| \leq H\left\|x-W_{x}\right\|+h\|x-y\|,
$$

$\forall x, y \in \wp$, where $H$ and $h$ s.t. $0 \leq H, 0 \leq h \leq 1$. Furthermore, it is supposed that Wis Picard $W$-stable.
We assume that the fractional diabetes model (11) is connected with the subsequent iterative formula

$$
\begin{align*}
& C_{n+1}(\eta)=C_{n}(\eta)+L^{-1}\left[\frac{s+\sigma(1-s)}{s} L\left[-(\lambda+\theta) C_{n}(\eta)+\lambda N_{n}(\eta)\right]\right] \\
& N_{n+1}(\eta)=N_{n}(\eta)+L^{-1}\left[\frac{s+\sigma(1-s)}{s} L\left[I-(v+\delta) C_{n}(\eta)-\mu N_{n}(\eta)\right]\right] . \tag{33}
\end{align*}
$$

In the above equation $\frac{s+\sigma(1-s)}{s}$ stands for the Lagrange multiplier.
Further, we develop the following result given as a theorem.

Theorem 3 Consider $G$ to be a self-map expressed in the subsequent manner:

$$
\begin{align*}
& G\left(C_{n}(\eta)\right)=C_{n+1}(\eta)=C_{n}(\eta)+L^{-1}\left[\frac{s+\sigma(1-s)}{s} L\left[-(\lambda+\theta) C_{n}(\eta)+\lambda N_{n}(\eta)\right]\right] \\
& G\left(N_{n}(\eta)\right)=N_{n+1}(\eta)=N_{n}(\eta)+L^{-1}\left[\frac{s+\sigma(1-s)}{s} L\left[I-(v+\delta) C_{n}(\eta)-\mu N_{n}(\eta)\right]\right] \tag{34}
\end{align*}
$$

then the iterations are G-stable in $L^{2}(a, b)$ if $\left(1-(\lambda+\theta) \varpi_{1}(\eta)+\lambda \varpi_{2}(\eta)\right)<1$ and $(1-(v+$ $\left.\delta) \varpi_{3}(\eta)-\mu \varpi_{4}(\eta)\right)<1$.

Proof First of all it will be shown that $G$ has a fixed point. To prove the above theorem, we derive the result given below for $(n, m) \in N \times N$.

$$
\begin{align*}
& \left\|G\left(C_{n}(\eta)\right)-G\left(C_{m}(\eta)\right)\right\| \\
& \quad \leq\left\|C_{n}(\eta)-C_{m}(\eta)\right\| \\
& \quad+\left\|L^{-1}\left[\frac{s+\sigma(1-s)}{s} L\left[-(\lambda+\theta)\left(C_{n}(\eta)-C_{m}(\eta)\right)+\lambda\left(N_{n}(\eta)-N_{m}(\eta)\right)\right]\right]\right\| \tag{35}
\end{align*}
$$

and

$$
\begin{align*}
& \left\|G\left(N_{n}(\eta)\right)-G\left(N_{m}(\eta)\right)\right\| \\
& \quad \leq\left\|N_{n}(\eta)-N_{m}(\eta)\right\| \\
& \quad+\left\|L^{-1}\left[\frac{s+\sigma(1-s)}{s} L\left[-(v+\delta)\left(C_{n}(\eta)-C_{m}(\eta)\right)-\mu\left(N_{n}(\eta)-N_{m}(\eta)\right)\right]\right]\right\| \tag{36}
\end{align*}
$$

We consider that the solutions play some role, i.e.,

$$
\begin{equation*}
\left\|C_{n}(\eta)-C_{m}(\eta)\right\| \cong\left\|N_{n}(\eta)-N_{m}(\eta)\right\| . \tag{37}
\end{equation*}
$$

Further, on using the result (37) in (35) and (36),

$$
\begin{equation*}
\left\|G\left(C_{n}(\eta)\right)-G\left(C_{m}(\eta)\right)\right\| \leq\left(1-(\lambda+\theta) \varpi_{1}(\eta)+\lambda \varpi_{2}(\eta)\right)\left\|C_{n}(\eta)-C_{m}(\eta)\right\| \tag{38}
\end{equation*}
$$

and

$$
\begin{equation*}
\left\|G\left(N_{n}(\eta)\right)-G\left(N_{m}(\eta)\right)\right\| \leq\left(1-(v+\delta) \varpi_{3}(\eta)-\mu \varpi_{4}(\eta)\right)\left\|N_{n}(\eta)-N_{m}(\eta)\right\|, \tag{39}
\end{equation*}
$$

where $\varpi_{1}, \varpi_{2}, \varpi_{3}$ and $\varpi_{4}$ are functions occurring from $L^{-1}\left[\frac{s+\sigma(1-s)}{s} L(\bullet)\right]$ with $(1-(\lambda+$ $\left.\theta) \varpi_{1}(\eta)+\lambda \varpi_{2}(\eta)\right)<1$ and $\left(1-(\nu+\delta) \varpi_{3}(\eta)-\mu \varpi_{4}(\eta)\right)<1$.

Consequently, the self-map $G$ possesses a fixed point. Further, we will show that the conditions associated with Theorem 2 are satisfied by G. Let Eqs. (38) and (39) hold, thus
on putting

$$
h=0, \quad H=\left\{\begin{array}{l}
\left(1-(\lambda+\theta) \varpi_{1}(\eta)+\lambda \varpi_{2}(\eta)\right),  \tag{40}\\
\left(1-(v+\delta) \varpi_{3}(\eta)-\mu \varpi_{4}(\eta)\right) .
\end{array}\right.
$$

Then from Eq. (40), we can notice that for the mapping $G$ along with the condition associated with Theorem 2 holds. Consequently for the considered mapping $G$ and the conditions pertaining to Theorem 2 are satisfied, hence $G$ is Picard G-stable. Thus, the proof of Theorem 3 is finished.

## 6 Numerical results and discussion

In the present portion, we compute the numerical results for $C(\eta)$ and $N(\eta)$ at $\sigma=0.70, \sigma=$ 0.85 and $\sigma=1$. The special solution of the non-integer diabetes model is derived by applying HATM and the Pade approximation [31] with as distinct parameters the number of $v=0.05, \delta=0.05, \mu=0.02, \gamma=0.08, \lambda=0.02$ and $I=60,000$. Fig. 1 depicts the nature of the number of diabetics having complications corresponding to the time for distinct order of fractional derivative. Figure 2 represents the impact of order of CF derivative on the size of the diabetics corresponding to time. Figure 3 indicates the effect of the rate at which diabetic patients having complications converted into the number of critically disabled diabetics having complications with respect to time. Figure 4 depicts the effect of the rate at which diabetic patients having complications converted into the size of critically disabled diabetics with respect to time. Figure 5 shows the effect of the mortality rate because of complications on the number of diabetics having complications with respect to time. Figure 6 indicates the effect of the mortality rate because of complications on the size of diabetics with respect to time. Figure 7 presents the effect of the natural mortality rate on the number of diabetics having complications with respect to time. Figure 8 presents the effect of the natural mortality rate on the size of diabetics with respect to time. Graphical behavior reveals that the model is significantly dependent on the arbitrary order. Figures $1-8$ demonstrate the clear difference at $\sigma=1, \sigma=0.85$ and $\sigma=0.70$. The model represents a new feature of $\sigma=0.85$ and $\sigma=0.70$, that was invisible when modeling with $\sigma=1$.

## 7 Conclusions

In this study, we have applied the novel introduced Caputo-Fabrizio arbitrary order derivative to the diabetes model and sloved the fractional problem via HATM and Padé approximation. The HATM is a pioneering and strong computational scheme to solve noninteger order differential equations. The HATM carries an auxiliary parameter $\hbar$, which is very useful to modify and manage the convergence of the series solutions. Hence, it can be determined that HATM is manageable, straightforward to employ, and a stronger computational procedure for examining linear and non-linear problems. The uniqueness and existence of the solutions are explored by exerting fixed point theory and the Picard-Lindelof approach. Some numerical results are analyzed to describe the effect of the arbitrary order. The effects of various parameters on the number of diabetics having complications and the size of diabetics with respect to time are shown graphically. By numerical simulation


Figure 1 Response of $C(\eta)$ w.r.t. time $\eta$ for different values of $\sigma$


Figure 2 Nature of $N(\eta)$ w.r.t. time $\eta$ for various values of $\sigma$


Figure 3 Nature of $C(\eta)$ w.r.t. time $\eta$ for different values of $v$


Figure 4 Behavior of $N(\eta)$ w.r.t. time $\eta$ for different values of $v$


Figure 5 Response of the solution $C(\eta)$ with respect to time $\eta$ for various values of $\delta$


Figure 6 Characteristic of $N(\eta)$ w.r.t. time $\eta$ for separate values of $\delta$


Figure 7 Response of $C(\eta)$ w.r.t. time $\eta$ for different values of $\mu$


Figure 8 Response of $N(\eta)$ w.r.t. time $\eta$ for distinct values of $\mu$
we can observe that when $\sigma \rightarrow 1$ the Caputo-Fabrizio non-integer order derivative reveals more absorbing characteristics. The results of this study are very helpful for medical practitioners dealing with diabetes and related issues. Thus, we have concluded that the CF derivative is useful in the description of physical, chemical, biological, medical, social, and engineering processes.

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## Authors' contributions

JS, DK and DB designed the study, developed the methodology, collected the data, performed the analysis, and wrote the manuscript. All authors read and approved the final manuscript.

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