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Study of Heat and Mass Transfer in Electroosmotic Flow of Third Order Fluid through Peristaltic Microchannels

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Received: 18 April 2019; Accepted: 10 May 2019; Published: 27 May 2019



Abstract: An analysis is carried out to evaluate the effects of heat and mass transfer in an electro-osmotic flow of third order fluid via peristaltic pumping. Solutions are derived for small wave number and Peclet number. The emerging non-linear mathematical model is solved analytically and compared numerically by the built-in scheme of working software. The table is inserted for shear stress distribution and a graph for comparison of solution techniques and accuracy of obtained results. The effects of various parameters of interest on pumping, trapping, temperature, heat transfer coefficient, and concentration distribution have been studied graphically. Electro-osmotic exchange of energy and mass has a role in reservoir engineering, chemical industry, and in micro-fabrication technologies.

Keywords: electroosmosis; peristalsis; third order fluid; heat and mass transfer; perturbation; pumping and trapping

1. Introduction

In the past few decades, electroosmosis has found rich applications. The first microfluidic devices were developed in the 1980s with the name Micro-Electro-Mechanical Systems (MEMS). These devices have broad application prospects in a variety of fields including biological and medical-related industries, where microfluidic devices are referred to as Lab-on-chip (LOC) devices. LOC can be used in many applications including clinical diagnostics, and biological or chemical contamination. Today, the need for the reliable, fast and well-organized performance of the microfluidic system has created a rigorous demand for small, easy-to-operate and cheap systematic apparatus for drug delivery and DNA analysis.

Microfluidic devices typically use electroosmosis to pump fluid into the entire device. Electroosmosis describes the flow of electrolyte through a channel having a charged boundary due to an applied voltage. Electro-osmosis occurs in many biological, medical and industrial processes such as porous membranes, tubule/canalicular flow, botanical processes, fluid dialysis, transport in human skin and separation techniques. Haung et al. [1] explained the Electroosmotic flow (EOF) in capillaries by monitoring method. Gravesen et al. [2] reviewed the detailed applications of microfluidics. Haswell [3] scrutinized the developmental features of micro-flows on EOF basis. Patankar and Hu [4] predicted the simulation of EOF. Also, Kang et al. [5] examined the EOF in the annulus of capillaries.

The dynamics of peristaltic flow are triggered by its emergence of physiology (intestines, ureters, esophagus, bile ducts, catheters, granules, etc.) and industry (blood pumps, transport of corrosive liquids and hygienic fluid transport). Most of the research on this topic has been conducted on

Newtonian fluids. Non-Newtonian fluids cannot be elucidated by a single constitutive relation. Hence, several constitutive relations of these fluids have been proposed. The impact of viscosity in peristalsis for third-grade fluid was explored by Hayat and Abbasi [6]. Noreen [7] proposed the physiological motion of the third order fluid with a magnetic field and mixed convection. Moreover, Noreen [8] considered the peristaltic motion of MHD couple-stress fluid. Prakash et al. [9] worked on peristaltically induced third-grade fluid via the asymmetric conduit. Mallick and Misra [10] discussed Erying–Powell fluid with the electromagnetic field.

Mathematical simulations of peristaltic distribution in microfluidic devices have newly attracted attention. Peristaltic motion can be controlled by adding and resisting external electric fields. One such example is the addition in peristaltic flow by electroosmosis [11]. Tang et al. [12] explored the electroosmosis in the power-law fluid. Hadigol et al. [13] considered the microscopic mixing characteristics of the power-law fluid in electroosmosis. Yavari et al. [14] demonstrated an increase in EOF temperature of bio-fluids (non-Newtonian) via microchannel. Furthermore, Bandopadhyay et al. [15] studied electroosmosis for Oldroyd-B fluid and wall slip. Jhorar et al. [17] expounded the biomechanical transfer through asymmetric microconduit. Similarly, Tripathi et al. [18] inspected the microvascular blood transport via EOF. Francesko et al. [19] observed the progress of LOC and microfluidics in detail. Javavel et al. [20] illustrated the EOF of pseudoplastic nano liquids through peristaltic pumping. However, the above studies did not examine the electroosmotic effects in peristaltically flowing third order fluid.

Heat transfer in microfluidic devices is of great importance in cancerous tissues destruction, portable kits for diseases diagnostics, the examination of dilution techniques in blood flow and the micro-fabrication technologies. Sadeghi et al. [21] examined the effect of heat exchange in EOF for viscoelastic fluids. Babaie et al. [22] noticed the impact of heat flow for Power-law fluid in the microchannel. Further, Chen et al. [23] analyzed heat exchange for Non-Newtonian fluid suspension in the microchannel. Thermal exchange in blood through capillaries was incorporated by Sinha and Shit [24]. Shit et al. [25] also showed the EOF for heat exchange and MHD. Moreover, in medical operations, it is considered that the wavy walls in electroosmotic flow increase the mass transfer. Therefore, Bhatti et al. [26] described the heat and mass transmission through EDL effects. Also, Reddy et al. [27] revealed the heat transport in peristaltic pumping for Casson fluid through the microchannel. Peristaltic pumping with EDL was studied by Yadav et al. [28]. Narla et al. [29] examined the effect of heat in EOF through time. Moreover, Yang et al. [30] also depicted the transportation of heat in magneto-hydrodynamic EOF.

To our knowledge, there is currently no report on heat and mass transfer in EOF of third-order fluid altered by peristalsis. The present study fills the gap. The main purpose of the present investigation is to analyze the effect of heat transfer and mass transfer for EOF of a third order fluid with viscous dissipation in a microchannel. Note, third order fluid has outstanding significance for shear thickening and shear thinning properties. This investigation has been carried out by using lubrication approximation and Debye Hückel linearization. The non-linear governing equations are solved through perturbation method. Pressure rise and heat transfer coefficient at channel walls have been calculated numerically. Finally, the graphical results for physical quantities are drawn and discussed in detail.

2. Mathematical Model

2.1. Flow Analysis

Consider 2D electroosmotic flow (EOF) of third order fluid in a microfluidic channel having a width 2a, presented in Figure 1. A peristaltic wave at velocity c propagates along the microchannel wall. The temperature and concentration fields of the lower channel wall are maintained at T_0 and C_0 whereas upper at T_1 and C_1 , respectively.



Figure 1. Schematic of the geometry of electro-osmotically modulated peristaltic flow.

Mathematically, microchannel geometry is expressed as:

$$\widetilde{Y} = \widetilde{h}(\widetilde{X}, \widetilde{t}) = \left(a + b \sin\left(2\pi \frac{(\widetilde{X} - c\widetilde{t})}{\lambda}\right)\right)$$
(1)

Here \tilde{X} , *a*, *b*, *c*, λ and \tilde{t} are the axial coordinate, channel's half width, wave amplitude, wave speed, wavelength and time respectively.

2.2. Governing Equations

In the laboratory frame, governing equations for flow of third order fluid in electrohydrodynamic (EHD) environment are [17]:

Continuity Equation:

$$\frac{\partial \widetilde{U}}{\partial \widetilde{X}} + \frac{\partial \widetilde{V}}{\partial \widetilde{Y}} = 0, \tag{2}$$

Equation of motion:

$$\rho \left(\frac{\partial}{\partial \widetilde{t}} + \widetilde{U} \frac{\partial}{\partial \widetilde{X}} + \widetilde{V} \frac{\partial}{\partial \widetilde{Y}} \right) \widetilde{U} = -\frac{\partial \widetilde{P}}{\partial \widetilde{X}} + \left(\frac{\partial \widetilde{S}_{\widetilde{X}\widetilde{X}}}{\partial \widetilde{X}} + \frac{\partial \widetilde{S}_{\widetilde{X}\widetilde{Y}}}{\partial \widetilde{Y}} \right) + \rho_e E_x, \tag{3}$$

$$\rho \left(\frac{\partial}{\partial \widetilde{t}} + \widetilde{U} \frac{\partial}{\partial \widetilde{X}} + \widetilde{V} \frac{\partial}{\partial \widetilde{Y}} \right) \widetilde{V} = -\frac{\partial \widetilde{P}}{\partial \widetilde{Y}} + \left(\frac{\partial \widetilde{S}_{\widetilde{Y}\widetilde{X}}}{\partial \widetilde{X}} + \frac{\partial \widetilde{S}_{\widetilde{Y}\widetilde{Y}}}{\partial \widetilde{Y}} \right).$$
(4)

Energy equation with viscous dissipation [9]:

$$\rho c_p \left(\frac{\partial}{\partial \widetilde{t}} + \widetilde{U} \frac{\partial}{\partial \widetilde{X}} + \widetilde{V} \frac{\partial}{\partial \widetilde{Y}} \right) \widetilde{T} = k \left(\frac{\partial^2}{\partial \widetilde{X}^2} + \frac{\partial^2}{\partial \widetilde{Y}^2} \right) \widetilde{T} + \left[\frac{\partial \widetilde{U}}{\partial \widetilde{X}} \widetilde{S}_{\widetilde{X}\widetilde{X}} + \frac{\partial \widetilde{V}}{\partial \widetilde{Y}} \widetilde{S}_{\widetilde{Y}\widetilde{Y}} + \left(\frac{\partial \widetilde{U}}{\partial \widetilde{Y}} + \frac{\partial \widetilde{V}}{\partial \widetilde{X}} \right) \widetilde{S}_{\widetilde{X}\widetilde{Y}} \right].$$
(5)

Concentration equation:

$$\left(\frac{\partial}{\partial \widetilde{t}} + \widetilde{U}\frac{\partial}{\partial \widetilde{X}} + \widetilde{V}\frac{\partial}{\partial \widetilde{Y}}\right)\widetilde{C} = D_m \left(\frac{\partial^2}{\partial \widetilde{X}^2} + \frac{\partial^2}{\partial \widetilde{Y}^2}\right)\widetilde{C} + \frac{k_t D_m}{T_m} \left(\frac{\partial^2}{\partial \widetilde{X}^2} + \frac{\partial^2}{\partial \widetilde{Y}^2}\right)\widetilde{T}.$$
(6)

Here \widetilde{U} is the velocity along \widetilde{X} direction, \widetilde{V} along \widetilde{Y} direction, ρ_e is electrical charge density, \widetilde{P} is pressure, ρ is density of fluid, E_x is axial electric field, c_p is the specific heat, k is thermal conductivity, k_t is ratio of thermal diffusion, D_m is mass diffusivity coefficient, T_m is average temperature, \widetilde{T} is temperature and \widetilde{C} is concentration field, which are dimensional.

Stress tensor for fluid [6] is defined as

$$\widetilde{S} = \left(\mu + \beta_3 \operatorname{tr}(\widetilde{A}_1^2)\right) \widetilde{A}_1 + \alpha_1 \widetilde{A}_2 + \alpha_2 \widetilde{A}_1^2 + \beta_1 \widetilde{A}_3 + \beta_2 \left(\widetilde{A}_1 \widetilde{A}_2 + \widetilde{A}_2 \widetilde{A}_1\right)$$
(7)

where μ , $\alpha_1, \alpha_2, \beta_1, \beta_2$ and β_3 are the material constants. Similarly, Revilin Erickson tensors A_n are

$$\widetilde{A}_{1} = \left(\nabla \widetilde{V}\right) + \left(\nabla \widetilde{V}\right)^{\mathrm{T}}, \text{ for } \widetilde{V} = [\widetilde{U}(\widetilde{X}, \widetilde{Y}, \widetilde{t}), \widetilde{V}(\widetilde{X}, \widetilde{Y}, \widetilde{t}), 0]$$

$$\widetilde{A}_{n} = \frac{\partial \widetilde{A}_{n-1}}{\partial \widetilde{t}} + \widetilde{A}_{n-1} \left(\nabla \widetilde{V}\right) + \left(\nabla \widetilde{V}\right)^{\mathrm{T}} \widetilde{A}_{n-1}, n > 1$$
(8)

The thermodynamic analysis of the model [7] shows that if all the fluid motion is thermodynamically compatible, then these motions must satisfy the Clausius–Duhem inequality. Moreover, it is also supposed that the specific Helmholtz free energy be minimal in equilibrium, then

$$\mu \ge 0, \quad \alpha_1 \ge 0, \quad |\alpha_1 + \alpha_2| \le \sqrt{24\mu\beta_3}, \quad \beta_1 = \beta_2 = 0, \quad \beta_3 \ge 0.$$
 (9)

In the present investigation, we assume that fluid is thermodynamically compatible, so Equation (7) reduces to the form

$$\widetilde{S} = \left(\mu + \beta_3 tr(\widetilde{A}_1^2)\right)\widetilde{A}_1 + \alpha_1 \widetilde{A}_2 + \alpha_2 \widetilde{A}_1^2$$
(10)

2.3. Electrohydrodynamics (EHD)

The Poisson equation [11] in a microchannel is described as:

$$\nabla^2 \widetilde{\Phi} = -\frac{\rho_e}{\epsilon} \tag{11}$$

Here $\rho_{e} \in \text{and } \overline{\Phi}$ represents total charge density, dielectric permittivity, and electric potential.

2.4. Potential Distribution

The net charge density ρ_e follows the Boltzmann distribution [12], given as

$$\rho_e = -z_v e \left(\widetilde{n}^- - \widetilde{n}^+ \right), \tag{12}$$

here the anions (\tilde{n}^-) and cat-ions (\tilde{n}^+) are defined through ρ_e of the Boltzmann Equation:

$$\widetilde{n}^{\pm} = \widetilde{n}_0 e^{\left(\pm \frac{ez_0}{T_{av} K_B} \widetilde{\Phi}\right)},\tag{13}$$

where n_0 represents bulk concentration, z_v the charge balance, K_B the Boltzmann constant, e the electronic charge and T_{av} is the average temperature.

Applying Debye-Hückel linearization approxiamtion [12], Equation (11) becomes

$$\frac{d^2\Phi}{dy^2} = m_e^2\Phi. \tag{14}$$

Here m_e is the electroosmotic parameter. The analytical solution of above Equation (14) subject to boundary conditions $\frac{\partial \Phi}{\partial y} = 0$, at y = 0 and $\Phi = 1$, at y = h(x) is obtained as;

$$\Phi(y) = \frac{\cosh(m_e y)}{\cosh(m_e h)}.$$
(15)

2.5. Non-Dimensionlization, Lubrication Approach and Boundary Conditions

In the stationary frame (\tilde{X}, \tilde{Y}) , the motion is unsteady due to the moving boundary. But, if viewed in a moving frame (\tilde{x}, \tilde{y}) , it can be considered as steady due to static boundary. Therefore, the translational transformation between two frames is [7]:

$$\widetilde{x} = \widetilde{X} - c\widetilde{t}, \, \widetilde{y} = \widetilde{Y}, \, \widetilde{u}(\widetilde{x}, \widetilde{y}) = \widetilde{U}(\widetilde{X}, \widetilde{Y}, \widetilde{t}) - c, \, \widetilde{v}(\widetilde{x}, \widetilde{y}) = \widetilde{V}(\widetilde{X}, \widetilde{Y}, \widetilde{t}), \\
\widetilde{p}(\widetilde{x}, \widetilde{y}) = \widetilde{P}(\widetilde{X}, \widetilde{Y}, \widetilde{t}), \, T(\widetilde{x}, \widetilde{y}) = \widetilde{T}(\widetilde{X}, \widetilde{Y}, \widetilde{t})$$
(16)

The above transformations is used in Equations (2)–(6) and then introducing the non-dimensional variables:

$$\begin{aligned} x &= \frac{2\pi\tilde{x}}{\lambda}, \ y &= \frac{\tilde{y}}{a}, \ t &= \frac{2\pi c\tilde{t}}{\lambda}, \ u &= \frac{\tilde{u}}{c}, \ v &= \frac{\tilde{v}}{c}, \ p &= \frac{2\pi a^2}{\lambda c\mu} \tilde{p}, \alpha &= \frac{2\pi a}{\lambda}, \ h &= \frac{\tilde{h}}{a}, \Phi &= \frac{\Phi}{a}, \\ P_e &= \frac{c\lambda}{D}, m_e &= \frac{a}{\lambda D}, n &= \frac{\tilde{n}}{n_0}, \varepsilon &= \frac{b}{a}, \ \theta &= \frac{\tilde{T}-T_0}{T_0}, \ \Omega &= \frac{\tilde{C}-C_0}{C_0}, \ \lambda_D &= \frac{1}{ez_v} \sqrt{\frac{T_0 \in K_B}{2n_0}}, \\ E_c &= \frac{c^2}{c_p T_0}, \Phi &= \frac{ez_v}{T_{av} K_B} \tilde{\Phi}, \ U_{hs} &= -\frac{E_x \tilde{\epsilon}}{c\mu}, \ S &= \frac{a}{c\mu} \tilde{S}, \ \lambda_1 &= \frac{\alpha_1 c}{a\mu}, \ \lambda_2 &= \frac{\alpha_2 c}{a\mu}, \ \gamma_1 &= \frac{\beta_1 c^2}{a^2 \mu}, \\ S_c &= \frac{v}{D_m}, \ \gamma_2 &= \frac{\beta_2 c^2}{a^2 \mu}, \ \gamma_3 &= \frac{\beta_3 c^2}{a^2 \mu}, \ \Psi &= \frac{\tilde{\Psi}}{ca}, \ u &= \frac{\partial \Psi}{\partial y}, \ v &= -\alpha \frac{\partial \Psi}{\partial x}, \ R_e &= \frac{\rho ca}{\mu}, \\ P_r &= \frac{\mu c_p}{k}, \ B_r &= E_c P_r, \ \Gamma &= \gamma_3 + \gamma_2, \ f &= \frac{\tilde{q}}{ca}, \ \Theta &= \frac{\tilde{Q}}{ca}, \ S_r &= \frac{T_0 \ k_r \ D_m}{v \ T_m \ C_0}. \end{aligned}$$

Here, *x* and *y* are the axial and transverse coordinate, m_e is electroosmotic parameter, *p* is dimensionless pressure, λ_D is Debye length, α is peristaltic wave number, Θ is dimensionless volume flow rate, θ is dimensionless temperature, Ω is dimensionless concentration, U_{hs} is Helmholtz–Smoluchowski velocity, ψ is non-dimensional stream function, P_e is ionic Peclet number, B_r is Brinkman, R_e is Reynolds number, P_r is Prandtl, Γ is Deborah number, ε is amplitude ratio, *S* is non-dimensional shear stress, S_r is non-dimensional Soret number, S_c is Schmidt number and E_c is Eckert number.

The relation of velocity and stream function ψ is defined as

$$u = \frac{\partial \Psi}{\partial y}, v = -\alpha \frac{\partial \Psi}{\partial x}.$$
 (18)

After introducing Equations (17) and (18), Equation (2) is satisfied identically and Equations (3)–(6) becomes

$$R_e \alpha \left(\frac{\partial \psi}{\partial y}\frac{\partial}{\partial x} - \frac{\partial \psi}{\partial x}\frac{\partial}{\partial y}\right)\frac{\partial \psi}{\partial y} = -\frac{\partial p}{\partial x} + \left(\alpha \frac{\partial S_{xx}}{\partial x} + \frac{\partial S_{xy}}{\partial y}\right) + m_e^2 U_{hs}\Phi,$$
(19)

$$-R_e \alpha^3 \left(\frac{\partial \psi}{\partial y} \frac{\partial}{\partial x} - \frac{\partial \psi}{\partial x} \frac{\partial}{\partial y} \right) \frac{\partial \psi}{\partial x} = -\frac{\partial p}{\partial y} + \alpha \left(\alpha \frac{\partial S_{xy}}{\partial x} + \frac{\partial S_{yy}}{\partial y} \right), \tag{20}$$

$$R_{e}\alpha\left(\frac{\partial\psi}{\partial y}\frac{\partial}{\partial x}-\frac{\partial\psi}{\partial x}\frac{\partial}{\partial y}\right)\theta = \frac{1}{P_{r}}\left(\alpha^{2}\frac{\partial^{2}}{\partial x^{2}}+\frac{\partial^{2}}{\partial y^{2}}\right)\theta + E_{c}\left[\alpha^{2}\frac{\partial}{\partial x}\left(\frac{\partial\psi}{\partial y}\right)S_{xx}-\alpha\frac{\partial}{\partial y}\left(\frac{\partial\psi}{\partial x}\right)S_{yy}+\left(\frac{\partial^{2}\psi}{\partial y^{2}}-\alpha^{2}\frac{\partial^{2}\psi}{\partial x^{2}}\right)S_{xy}\right], \quad (21)$$

$$R_e \alpha \left(\frac{\partial \psi}{\partial y}\frac{\partial}{\partial x} - \frac{\partial \psi}{\partial x}\frac{\partial}{\partial y}\right) \Omega = \frac{1}{S_c} \left(\alpha^2 \frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2}\right) \Omega + S_r \left(\alpha^2 \frac{\partial^2}{\partial x^2} + \frac{\partial^2}{\partial y^2}\right) \theta$$
(22)

Now subject to long-wavelength assumption [10] on Equations (19)–(22), ignoring higher powers of α , we get;

$$\frac{\partial p}{\partial x} = \frac{\partial}{\partial y} \left[\frac{\partial^2 \psi}{\partial y^2} + 2\Gamma \left(\frac{\partial^2 \psi}{\partial y^2} \right)^3 \right] + m_e^2 U_{hs} \Phi, \qquad (23)$$

$$\frac{\partial p}{\partial y} = 0 \tag{24}$$

$$\frac{\partial^2 \theta}{\partial y^2} + B_r \left[\left(\frac{\partial^2 \psi}{\partial y^2} \right)^2 + 2\Gamma \left(\frac{\partial^2 \psi}{\partial y^2} \right)^4 \right] = 0$$
(25)

$$\frac{\partial^2 \Omega}{\partial y^2} + S_c S_r \left(\frac{\partial^2 \theta}{\partial y^2} \right) = 0$$
(26)

where $S_{xy} = \frac{\partial^2 \psi}{\partial y^2} + 2\Gamma \left(\frac{\partial^2 \psi}{\partial y^2}\right)^3$. Eliminating pressure from Equations (23) and (24), we get

$$\frac{\partial^2}{\partial y^2} \left[\frac{\partial^2 \psi}{\partial y^2} + 2\Gamma \left(\frac{\partial^2 \psi}{\partial y^2} \right)^3 \right] + m_e^2 U_{HS} \frac{\partial \Phi}{\partial y} = 0$$
(27)

The dimensionless boundary conditions are:

$$\frac{\partial^2 \psi}{\partial y^2} = 0, \quad \psi = 0, \quad \frac{\partial \theta}{\partial y} = 0, \quad \frac{\partial \Omega}{\partial y} = 0, \quad \text{at } y = 0, \quad (28)$$

$$\frac{\partial \Psi}{\partial y} = -1, \ \Psi = f, \quad \theta = 0, \ \Omega = 0, \quad \text{at } y = h(x) = 1 + \varepsilon \sin x$$
 (29)

here f is dimensionless mean flow rate in moving frame and Θ is dimensionless mean flow rate in fixed frame. Furthermore, f in the moving frame is connected to the Θ in the fixed frame as:

$$\Theta = f + 1. \tag{30}$$

here f = $\int_0^{h(x)} \frac{\partial \psi}{\partial y} dy = \psi(h(x)) - \psi(0).$

3. Solution Methodology

3.1. Series Solutions

The governing differential system given in Equations (23)–(27) subject to boundary conditions (28,29) consists of highly non-linear and coupled equations. Exact solution of the governing system of equations is impossible. Therefore, analytical solution of the system is presented through the perturbation technique by taking Deborah number Γ as a perturbation parameter. So, we expand ψ , θ , $\Omega_{,.} dp/dx$ and f as in [8]:

$$\psi = \psi_0 + \Gamma \psi_1 + O(\Gamma)^2 \tag{31}$$

$$\theta = \theta_0 + \Gamma \theta_1 + O(\Gamma)^2 \tag{32}$$

$$\Omega = \Omega_0 + \Gamma \Omega_1 + O(\Gamma)^2 \tag{33}$$

$$dp/dx = dp_0/dx + \Gamma dp_1/dx + O(\Gamma)^2$$
(34)

$$f = f_0 + \Gamma f_1 + (\Gamma)^2$$
 (35)

3.2. Zero Order System $(\Gamma)^0$

$$\frac{\partial^4 \psi_0}{\partial y^4} + m_e^2 U_{HS} \frac{\partial \Phi}{\partial y} = 0, \qquad (36)$$

$$\frac{\partial^2 \theta_0}{\partial y^2} + B_r \left(\frac{\partial^2 \psi_0}{\partial y^2}\right)^2 = 0 \tag{37}$$

$$\frac{\partial^2 \Omega_0}{\partial y^2} + S_c \cdot S_r \left(\frac{\partial^2 \theta_0}{\partial y^2} \right) = 0$$
(38)

6 of 20

Appl. Sci. 2019, 9, 2164

$$\frac{\partial p_0}{\partial x} = \frac{\partial^3 \Psi_0}{\partial y^3} + m_e^2 U_{HS} \Phi, \qquad (39)$$

subject to

$$\begin{split} \psi_0 &= 0, \quad \frac{\partial^2 \psi_0}{\partial y^2} = 0, \quad \frac{\partial \theta_0}{\partial y} = 0, \quad \frac{\partial \Omega_0}{\partial y} = 0, \text{ at } y = 0 \\ \psi_0 &= f_0, \quad \frac{\partial \psi_0}{\partial y} = -1, \quad \theta_0 = 0, \quad \Omega_0 = 0, \text{ at } y = h \end{split}$$
(40)

3.3. First Order System $(\Gamma)^1$

$$\frac{\partial^4 \psi_1}{\partial y^4} + 6 \left(\frac{\partial^2 \psi_0}{\partial y^2}\right)^2 \frac{\partial^4 \psi_0}{\partial y^4} + 12 \left(\frac{\partial^3 \psi_0}{\partial y^3}\right)^2 \frac{\partial^2 \psi_0}{\partial y^2} = 0, \tag{41}$$

$$\frac{\partial^2 \theta_1}{\partial y^2} + 2B_r \left[\left(\frac{\partial^2 \psi_0}{\partial y^2} \right) \left(\frac{\partial^2 \psi_1}{\partial y^2} \right) + \left(\frac{\partial^2 \psi_0}{\partial y^2} \right)^4 \right] = 0$$
(42)

$$\frac{\partial^2 \Omega_1}{\partial y^2} + S_c \cdot S_r \left(\frac{\partial^2 \theta_1}{\partial y^2} \right) = 0$$
(43)

$$\frac{\partial p_1}{\partial x} = \frac{\partial^3 \psi_1}{\partial y^3} + 6 \left(\frac{\partial^2 \psi_0}{\partial y^2}\right)^2 \frac{\partial^3 \psi_0}{\partial y^3},\tag{44}$$

subject to

$$\psi_1 = 0, \quad \frac{\partial^2 \psi_1}{\partial y^2} = 0, \quad \frac{\partial \theta_1}{\partial y} = 0, \quad \frac{\partial \Omega_1}{\partial y} = 0, \text{ at } y = 0, \\ \psi_1 = f_1, \quad \frac{\partial \psi_1}{\partial y} = 0, \quad \theta_1 = 0, \quad \Omega_1 = 0, \text{ at } y = h$$
(45)

The solution for zero and first order system is defined in the appendix. By substituting the zero and first order solution in the Equations (32)–(35) we get the final first order solution for ψ , θ , Ω and dp/dx.

The dimensionless pressure rises per wavelength ΔP_{λ} is as follows:

$$\Delta P_{\lambda} = \int_{0}^{1} \frac{\partial p}{\partial x} dx.$$
(46)

The heat transfer coefficient *Z* is defined as:

$$Z = h_x \frac{\partial \theta}{\partial y} \Big|_{y = h(x)}.$$
(47)

The shear stress S_{xy} at the wall y = h(x) is defined as

$$S_{xy} = \left(\frac{\partial^2 \psi}{\partial y^2} + 2\Gamma \left(\frac{\partial^2 \psi}{\partial y^2}\right)^3\right) \bigg|_{y = h(x)}.$$
(48)

4. Computational Discussion of Results

This section deals with the impacts of different parameters (i.e., Deborah number Γ , electroosmotic parameter m_e , Helmholtz–Smoluchowski/maximum electroosmotic velocity U_{hs} , Brinkman number B_r , volume flow rate Θ , Soret number S_r and Schmidt number S_c) on the pressure rise ΔP_{λ} , pressure gradient dp/dx, velocity distribution u, temperature distribution θ , concentration distribution Ω , heat transfer coefficient Z and streamlines ψ . Figures 2–11 are plotted to serve the purpose.

7 of 20

Figure 2 reveals the overlapping comparison between the analytical and numerical solution of velocity. Note that the analytical solution is obtained through perturbation technique and numerical solution is from built-in scheme NDsolve of working software. It is concluded that our results through perturbation (i.e., truncated up to first order of third-order fluid parameter) are in good agreement with the numerical scheme.



Figure 2. Comparative analysis between Perturbation solution and Numerical solution for axial velocity, while other parameters are $\Gamma = 0.01$, $x = \pi$, $\varepsilon = 0.2$, $\Theta = 1$, $m_e = 2$, $U_{hs} = -1$, $B_r = 1$, $S_c = 1$, $S_r = 1$.

4.2. Flow Characteristics

The axial velocity deserves to provide the main features of flow behavior in the microchannel for microfluidic applications. Figure 3a–c is portrayed to inspect the evolutions in the velocity profile across the microchannel for various values of Deborah number Γ , electroosmotic parameter m_e and maximum electroosmotic velocity U_{hs} . Figure 3a shows the variation of axial velocity for various values of Γ . It is demonstrated that the magnitude of velocity u decreases as Γ increases at y = 0. The reason behind this trend is EDL (electric double layer). It means flow of fluid resists in the presence of EDL. But opposite behavior is witnessed near the channel wall y = h(x). Figure 3b predicts that m_e boosts the magnitude of u i.e., magnitude of u increases as m_e increases at the central region of the channel and reduces near the walls of the channel. Since m_e is the ratio of the channel height to the Debye length λ_D , it specifies that λ_D is inversely proportional to EDL. Hence, more fluid flows in the central region y = 0. Figure 3c depicts that u decent as U_{hs} ascents in y = 0, whereas opposite movement is noticed at y = h(x). As U_{hs} physically concludes that velocity of fluid decreases when thickness of EDL increases. Therefore, fluid flow decreases in EDL presence.

4.3. Pumping Characteristics

Peristalsis is characterized by pushing the fluid forward against the pressure rise ΔP_{λ} . The characteristic of pumping can be analyzed in the form of ΔP_{λ} versus Θ , shown in Figure 4a–c Pumping action divides the whole region into the four segments: pumping region (adverse pressure gradient $\Delta P_{\lambda} > 0$, Positive pumping $\Theta > 0$, backward/retrograde pumping ($\Delta P_{\lambda} > 0$, $\Theta < 0$), augmented pumping (favorable pressure gradient $\Delta P_{\lambda} < 0$, Positive pumping $\Theta > 0$) and free pumping ($\Delta P_{\lambda} = 0$). Figure 4a portrays the ΔP_{λ} against Θ for different values of Γ . It is perceived that pumping increases as Γ increases in pumping region ($\Delta P_{\lambda} > 0$). But in co-pumping ($\Delta P_{\lambda} < 0$), pumping decreases as Γ increases. It means that pumping rate is high for third order fluid under parameter Γ in comparison to viscous fluid $\Gamma = 0$. For free pumping ($\Delta P_{\lambda} = 0$), curves coincide i.e., there is no difference between third-grade fluid and Newtonian fluid within the domain of -0.4 to 0.4. Figure 4b describes that ΔP_{λ} decreases as m_e increases. This means additional pressure is needed to drive the third-grade fluid for maximum thickness of EDL. It physically declares that formation of EDL on charged surface opposes the third-grade fluid flow. Furthermore, Figure 4c illustrates ΔP_{λ} decreases as U_{hs} increases. Here pumping is controlled through an electric field. Note that there is a linear relationship between ΔP_{λ} against Θ , because of U_{hs} .



Figure 3. Axial velocity u profile for (a). Γ (b) m_e (c) U_{hs} , while other parameters are $\Gamma = 0.01, x = \pi, \varepsilon = 0.358, \Theta = 0.8, m_e = 5, U_{hs} = -1.$

The distribution of pressure gradient dp/dx is plotted through Figure 5a–d for Deborah number Γ , electroosmotic parameter m_e , maximum electroosmotic velocity U_{hs} and volume flow rate Θ . Figure 5a portrays that by increasing the Γ , magnitude of pressure gradient dp/dx increase. Γ being a physical parameter showing the Non-Newtonian nature, one can observe pressure gradients are higher for

third order in comparison to Newtonian fluids. Figure 5b illustrates that by increasing the values of $\Theta_{,.}$ dp/dx decreases. Because there is an inverse relationship between dp/dx and Θ . Figure 5c highlights that by raising the values of $m_{e,.}$ dp/dx decreases. It means that EDL presence in charged surface resists the flow of third grade fluid. Similarly, it is important that the characteristics of pumping can be modified by EDL mechanism and process of pumping can be systematized by thinning and thickening EDL width. Also, Figure 5d presents that the magnitude of dp/dx increases as U_{hs} increases.



Figure 4. Pressure rise ΔP_{λ} profile for (**a**). Γ (**b**). m_e (**c**). U_{hs} , while other parameters are $\Gamma = 0.1, \varepsilon = 0.258, m_e = 2, U_{hs} = -1.$



Figure 5. Pressure gradient profile for (a). Γ (b). Θ (c). m_e (d). U_{hs} , while other parameters are $\Gamma = 0.01, \varepsilon = 0.4, \Theta = 0.1, m_e = 2, U_{hs} = -1, B_r = 1.$

4.4. Trapping Characteristics

Trapping for different values of Deborah number Γ , electroosmotic parameter m_e and maximum electroosmotic velocity U_{hs} are shown in Figures 6–8. Figure 6a–d reveal a streamline structure for various values of Γ . The size of the enclosed bolus decreases as Γ increases. It means that size of trapped bolus is strongly affected by changing the fluid. Similarly, Figure 7a–d visualize that the number of enclosed bolus increases with rise in electroosmotic parameter m_e . It scrutinizes that as the m_e increases, the enclosed bolus strongly appears in EDL. It means that more fluid can be trapped in the presence of more electric field. Figure 8a–d illustrate streamline makeup for U_{hs} . It demonstrates that the accumulation of streamline reduces with rise in U_{hs} . Thus, stronger U_{hs} means stronger the external electric field, the number of enclosed significantly decreases for large values of bolus U_{hs} .

4.5. Temperature Characteristics

In the present subsection, we examined the effect of Deborah number Γ , electroosmotic parameter m_e , maximum electroosmotic velocity U_{hs} and Brinkman number B_r on heat transfer characteristics. Figure 9a reveals the various values of Γ , for temperature distribution θ . It examines that θ enhances for more values of Γ . Also, Figure 9b displays that θ increases by increasing m_e . Therefore, the decrease in EDL produces a rise in θ remarkably. Figure 9c, disclose the effects of maximum electroosmotic velocity U_{hs} on θ . It signifies that for more values of U_{hs} , θ declines. Figure 9d exemplifies the variation of B_r on θ i.e., θ increases for more values of B_r . The main fact in B_r is the viscosity, which creates resistance. This resistance is responsible for the collision of fluid particles and the resulting collision causes an increase in temperature.

1.0

0.8

0.0

0.4

0.2

0.0

1.0

0.8

0.6

>

0 1 2 3 4

x

(a)

1

0.4 0.4 0.2 0.2 0.0 0.0 0 2 3 0 2 3 4 1 1 4 1 1 x х (c) (**d**)

1.0

0.8

0.6

0.4

0.2

0.0

1.(

0.8

0.0

0 1 2 3 4

x

(b)

Figure 6. Streamline distribution for (**a**). $\Gamma = 0.00$ (**b**). $\Gamma = 0.02$ (**c**). $\Gamma = 0.04$ (**d**). $\Gamma = 0.06$, while other parameters are $\varepsilon = 0.258$, $\Theta = 0.8$, $m_e = 2$, $U_{hs} = -1$.



Figure 7. Streamline distribution for (a). $m_e \rightarrow 0$ (b). $m_e = 2$ (c). $m_e = 5$ (d). $m_e = 10$, while other parameters are $\varepsilon = 0.358$, $\Theta = 0.7$, $\Gamma = 0.01$, $U_{hs} = -1$.



Figure 8. Streamline distribution for (**a**). $U_{hs} = -1.0$ (**b**). $U_{hs} = 0.0$ (**c**). $U_{hs} = 1.0$, while other parameters are $\varepsilon = 0.758$, $\Theta = 0.7$, $\Gamma = 0.01$, $m_e = 2$.



Figure 9. Temperature profile for (**a**). Γ (**b**). m_e (**c**). U_{hs} (**d**). B_r , while other parameters are $\Gamma = 0.01, x = 1, \varepsilon = 0.2, \Theta = 1, m_e = 2, U_{hs} = -1, B_r = 1.$



Figure 10. Cont.



Figure 10. Concentration profile for (**a**). Γ (**b**). m_e (**c**). U_{hs} (**d**). B_r . (**e**). S_c (**f**). S_r , while other parameters are $\Gamma = 0.01$, $x = \pi$, $\varepsilon = 0.2$, $\Theta = 1$, $m_e = 2$, $U_{hs} = -1$, $B_r = 1$, $S_c = 1$, $S_r = 1$.



Figure 11. Heat transfer coefficient for (**a**). Γ (**b**). m_e (**c**). U_{hs} (**d**). B_r , while other parameters are $\Gamma = 0.01, \varepsilon = 0.2, \Theta = 1.2, m_e = 10, U_{hs} = -1, B_r = 1.$

4.6. Concentration Characteristics

Here, we observed the impact of Deborah number Γ , electroosmotic parameter m_e , maximum electroosmotic velocity U_{hs} , Brinkman number B_r , Schmidt number S_c and Soret number S_r on mass transfer characteristics. Figure 10a reveals the various values of Γ , for temperature distribution θ . It examines that Ω decreases for more values of Γ . Figure 10b displays that Ω decreases by increasing m_e . Figure 10c discloses the effects of U_{hs} on Ω . It signifies that for more values of U_{hs} , θ rises. Figure 10d exemplifies the variation of B_r on Ω i.e., Ω decreases for more values of B_r . Figure 10e discloses the effects of S_c on Ω . It implies that Ω decreases for more values of S_c . Also, Figure 10f shows that Ω decreases for more values of S_r .

4.7. Heat Transfer Coefficient

Figure 11a–d demonstrates the changes in heat transfer coefficient *Z* for various values of Deborah number Γ , electroosmotic parameter m_e , maximum electroosmotic velocity U_{hs} , and Brinkman number B_r . This reveals that the various values of Γ , B_r and m_e , the heat transfer coefficient increases. On contrary *Z* is decreased for different values of U_{hs} .

4.8. Shear Stress Distribution

Distribution of shear stress provides very useful information about the nature of dissipation at walls. Table 1a–c show the distribution of axial shear stress for different values of Deborah number Γ , maximum electroosmotic velocity U_{hs} and electroosmotic parameter m_e . Table 1a explains that the magnitude of shear stress decreases in the axial direction while it decreases (near the channel wall) then increases (away from channel wall) for different values of Γ . Here, it is obvious that the third-grade fluid act as a shear thinning and shear thickening fluid. Similarly, Table 1b displays the variation of shear stress for U_{hs} . It is found that the axial distribution of shear stress enhances for increasing values of U_{hs} . Table 1c presents the shear stress distribution for different values of m_e . It is concluded that the same behavior is observed as of Γ . Therefore, we say that the presence of electroosmotic parameter strongly infers the characteristics of third order fluid parameter.

Table 1. Numerical values of shear stress at the wall $y = 1 + \varepsilon \sin x$ for different values of Deborah number Γ , Helmholtz–Smoluchowski velocity U_{hs} and electroosmotic parameter m_e .

(a) $\varepsilon = 0.385, \Theta = 1.0, m_e = 2, U_{hs} = -1.0.$					
x	Г				
	0.0	0.2	0.4		
0.1	-2.4917	-3.0742	-0.8223		
0.2	-2.3669	-3.8790	-2.2711		
0.3	-2.6497	-4.5689	-4.3309		
(b) $\varepsilon = 0.385, \Theta = 1.0, m_e = 2, \Gamma = 0.1.$					
x	U_{hs}				
	-1.0	0.0	1.0		
0.1	-3.6869	-1.1370	0.5727		
0.2	-3.7974	-1.2903	0.4311		
0.3	-3.8395	-1.4188	0.3188		
(c) $\varepsilon = 0.385, \Theta = 1.0, \Gamma = 0.1, U_{hs} = -1.0$					
x	m _e				
	2	4	6		
0.1	-3.6869	-10.778	-6.3530		
0.2	-3.7974	-8.7149	-4.7301		
0.3	-3.8395	-7.0505	-3.6590		

5. Concluding Remarks

From current analysis, we conclude that:

- The axial velocity of third-grade fluid in the microchannel is enhanced due to an increase in electroosmotic parameter.
- For higher values of Schmidt or Soret number, concentration profile decreases.
- The magnitude of pressure rise decreases in the pumping region with the increase of electroosmotic parameter.
- Pressure gradient is more for EOF of third order fluid in comparison to peristaltically flowing fluid.

- Volume of trapped bolus is strongly dependent upon the electroosmotic parameter whereas it is suppressed for Deborah number and maximum electroosmotic velocity.
- Temperature distribution in microchannel is crucial for electroosmotic parameter, Deborah number and Brinkman number.

Electro-osmotic exchange of energy and mass has a role in reservoir engineering. The outcomes of the present model may be used by engineers in chemical industry and in micro-fabrication technologies. In the future, this model accompanying porosity will help in understanding the hydrodynamics of rheological fluids in a typical channel employed in a LOC system. This course requires promotion, being of significance in factual life applications.

Author Contributions: Formal analysis, S.N.; Funding acquisition, A.H.; Investigation, S.W.; Methodology, S.W.; Resources, S.N.; Supervision, S.N.; Writing—original draft, S.W.; Writing—review & editing, A.H.

Funding: This research received no external funding. The APC was funded by Ton Duc Thang University, Ho Chi Minh City, Vietnam. However, no grant number is available from source.

Acknowledgments: The third author would like to thank Ton Duc Thang University, Ho Chi Minh City, Vietnam for the financial support.

Conflicts of Interest: The authors declare no conflict of interest.

Nomenclature

Symbols	Meaning	Dimensions
a	Channel's half width	[L]
\widetilde{A}_n	Revilin Erickson tensor	$[M/LT^2]$
b	Wave amplitude	[L]
B_r	Brinkman number	[-]
С	Wave speed	[L/T]
\widetilde{C}	Dimensional concentration field	[N/L ³]
<i>C</i> ₀ , <i>C</i> ₁	Concentration field at upper and lower wall	[N/L ³]
D	Diffusivity of chemical species	[-]
D_m	Mass diffusivity coefficient	$[L^2/T]$
E _c	Eckert number	[-]
E_x	Axial electric field	[ML/AT ³]
е	Electron charge	[AT]
f	Non-dimensional mean flow rate in moving frame	[-]
\widetilde{h}	Dimensional upper wall	[L]
h	Non-dimensional upper wall	[-]
k	Thermal conductivity	[ML/KT ³]
k_t	Ratio of thermal diffusion	[N/L ³]
K _B	Boltzmaan constant	$[ML^2/T^2K]$
m _e	Electroosmotic parameter	[-]
\widetilde{n}^{\pm}	Positive and negative ions	[AT]
<i>n</i> ₀	Bulk concentration	[AT]
\widetilde{P}	Dimensional pressure field	$[ML/T^2]$
р	Non-dimensional pressure field	[-]
P_e	Peclet number	[-]
P_r	Prandtl number	[-]
Q	Dimensional volume flow rate in fixed frame	[L ³ /T]
R_e	Reynolds number	[-]
ŝ	Dimensional shear stress	$[M/LT^2]$
S	Non-dimensional shear stress	[-]
Sc	Schmidt number	[-]
S_r	Soret number	[-]

\overline{t}	Dimensional time	[T]
t	Non-dimensional time	[-]
\widetilde{T}	Dimensional temperature field	[K]
T_0, T_1	Temperature field at upper and lower wall	[K]
T_m	Mean temperature	[K]
\widetilde{u}	Dimensional velocity along \widetilde{X} direction	[L/T]
и	Non-dimensional velocity along <i>x</i> direction	[-]
U_{hs}	Helmholtz-Smoluchowski velocity	[L/T]
\widetilde{V}	Dimensional velocity along \widetilde{Y} direction	[L/T]
υ	Non-dimensional velocity along <i>y</i> direction	[-]
\widetilde{X}	Dimensional axial coordinate in fixed frame	[L]
x	Non-dimensional axial coordinate in moving frame	[-]
\widetilde{Y}	Dimensional transverse coordinate in fixed frame	[L]
у	Non-dimensional transverse coordinate in moving frame	[-]
Z	heat transfer coefficient	[-]
Z_v	Valence of ions	[-]
α	Peristaltic wave number	[-]
α_1, α_2	Material constants	[-]
$\beta_1, \beta_2, \beta_3$	Material constants	[-]
$\gamma_1, \gamma_2, \gamma_3$	Material coefficients	[-]
λ	Wavelength	[L]
λ_1, λ_2	Material coefficients	[-]
λ_D	Debye length	[L]
∈	Dielectric permittivity	[-]
$ ho_e$	Toatl charge density	$[AT/L^3]$
ρ	Density of fluid	$[M/L^3]$
$\widetilde{\Phi}$	Dimensional electric potential	$[ML^2/T^2]$
Φ	Non-dimensional electric potential	[-]
Г	Non-dimensional Deborah number	[-]
ε	Amplitude ratio	[-]
Θ	Non-dimensional volume flow rate	[-]
θ	Non-dimensional temperature field	[-]
Ω	Non-dimensional concentration field	[-]
$\widetilde{\psi}$	Dimensional stream function	$[L^2/T]$
ψ	Non-dmensional stream function	[-]

Appendix A

$$\psi_0 = A_1 y + A_2 y^3 + A_3 \sinh(m_e y),$$

$$u_0 = A_1 + 3A_2 y^2 + A_3 m_e \cosh(m_e y)$$

$$\frac{\partial p_0}{\partial x} = 3A_4 + A_5 \cosh(m_e y),$$

 $\theta_{0} = B_{1} + B_{2}y^{2} + B_{3}y^{4} + B_{4}y\cosh(m_{e}y) + B_{5}y\cosh(m_{e}y)^{3} + B_{6}y\sinh(m_{e}y) + B_{7}y\sinh(m_{e}y)^{3} + B_{8}y\cosh(m_{e}y)^{2}\sinh(m_{e}y) + B_{9}y\cosh(m_{e}y)\sinh(m_{e}y)^{2} + B_{10}\sinh(m_{e}y) + B_{11}\sinh(2m_{e}y) + B_{12}\sinh(m_{e}y)^{3} + B_{13}\sinh(2m_{e}y)^{3} + B_{14}\cosh(2m_{e}y)^{2}\sinh(2m_{e}y) + B_{15}\cosh(m_{e}y)\sinh(m_{e}y)^{2}$

$$\begin{aligned} \Omega_0 &= C_1 + C_2 y^2 + C_3 y^4 + C_4 y \cosh(m_e y) + C_5 y \cosh(m_e y)^3 + C_6 y \sinh(m_e y) + \\ C_7 y \sinh(m_e y)^3 + C_8 y \cosh(m_e y)^2 \sinh(m_e y) + C_9 y \cosh(m_e y) \sinh(m_e y)^2 + \\ C_{10} \cosh(m_e y) + C_{11} \cosh(2m_e y) + C_{12} \sinh(m_e y) + C_{13} \sinh(2m_e y) + C_{14} \cosh(m_e y)^3 + \\ C_{15} \cosh(2m_e y)^3 + C_{16} \sinh(m_e y)^3 + C_{17} \sinh(2m_e y)^3 + C_{18} \cosh(m_e y)^2 \sinh(m_e y) + \\ C_{19} \cosh(m_e y) \sinh(m_e y)^2 + C_{20} \cosh(2m_e y)^2 \sinh(2m_e y) + C_{21} \sinh(2m_e y)^2 \cosh(2m_e y). \end{aligned}$$

 $u_{1} = 3D_{2}y^{2} + 5D_{3}y^{4} + m_{e}D_{4}y^{2}\cosh(m_{e}y) + 2D_{4}y\sinh(m_{e}y) + m_{e}D_{5}y\sinh(m_{e}y)$ $+ D_{5}\cosh(m_{e}y) + 2m_{e}D_{6}y^{1}\sinh(2m_{e}y) + D_{6}\cosh(2m_{e}y)$ $+ m_{e}D_{7}\cosh(m_{e}y) + 2D_{8}m_{e}\cosh(2m_{e}y) + 3D_{9}m_{e}\cosh(3m_{e}y)$

$$\frac{\frac{\partial p_1}{\partial x}}{\frac{\partial p_1}{\partial x}} = D_{11} + D_{12}y^2 + D_{13}y^2 \cosh(m_e y) + D_{14}y \sinh(m_e y) + D_{15}y \sinh(2m_e y) \\ + D_{16}y \cosh(m_e y) \sinh(m_e y) + D_{17} \cosh(m_e y) + D_{18} \cosh(2m_e y) \\ + D_{19} \cosh(3m_e y) + D_{20} \cosh(m_e y) \sinh(m_e y)^2 + D_{21} \sinh(m_e y)^2$$

$$\begin{aligned} \theta_1 &= E_1 + E_2 y^2 + E_3 y^4 + E_4 y^6 + E_5 y^3 \sinh(m_e y) + E_6 y^2 \cosh(m_e y) \\ &+ E_7 y^2 \cosh(2m_e y) + E_8 y \sinh(m_e y) + E_9 y \sinh(2m_e y) \\ &+ E_{10} y \sinh(3m_e y) + E_{11} \cosh(m_e y) + E_{12} \cosh(2m_e y) \\ &+ E_{13} \cosh(3m_e y) + E_{14} \cosh(4m_e y) \end{aligned}$$

$$\begin{aligned} \Omega_1 = & F_1 + F_2 y^2 + F_3 y^4 + F_4 y^6 + F_5 y^3 \sinh(m_e y) + F_6 y^2 \cosh(m_e y) \\ & + F_7 y^2 \cosh(2m_e y) + F_8 + F_8 y \sinh(m_e y) + F_9 y \sinh(2m_e y) \\ & + F_{10} y \sinh(3m_e y) + F_{11} \cosh(m_e y) + F_{12} \cosh(2m_e y) \\ & + F_{13} \cosh(3m_e y) + F_{14} \cosh(4m_e y) \end{aligned}$$

where $A_1 - A_5, B_1 - B_{15}, C_1 - C_{21}, D_1 - D_{21}, E_1 - E_{14}$ and $F_1 - F_{14}$ are evaluated through Mathematica software package.

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