



IONIC TRANSITION OF ELECTROVISCOUS BOUNDARY LAYER FLOW WITH THE ANALYSIS OF MAXWELL AND NERNST-PLANCK EQUATION

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Abstract The aim of the present study is to explore a numerical model for an ionic transition of boundary layer electroviscous fluid flow over a spinning disc. For the purpose, the Nernst-Planck, Maxwell and Poisson's equations are coupled and simulated with the Navier Stokes equation for the ionic species conservation. The basic equations of the electroviscous boundary layer flow are diminished to an ordinary dimensionless system of differential equations. The Matlab built-in package bvp5c has been implemented for the numerical outcomes of the transform equations. The outputs are compared to the existing literature, which seemed to be in good agreement with each other, to ensure that the results are accurate. It has been presumed that the action of Debye length coefficient reduces the potential gradient and exposes an excess number of ions to the flow field, which causes an enhancement in the fluid velocity. The consequences of physical constraints on velocity as well as negative and positive charge profiles are drawn and briefly addressed.

MSC: 47H09

Keywords: Maxwell equation; Dielectric constant; bvp5c technique; Anion and cation ions; Surface charge density; Nernst-Planck equations

1. INTRODUCTION

An ionic medium is made up of cations and anions, which have opposite charges. High conductivity, tunable viscosity, low volatility, a broad electrochemical window, and low toxicity are some of the differences between the two [1]. Extractions, gas and liquid chromatography, mass spectrometer, capillary electrophoresis, spectroscopy, electrochemistry, and sensors are only a few of the uses for ionic liquids in chemical analyzing a fluid [2]. Endocrine-disrupting chemicals (EDCs) have recently advanced to the point that they are a threat to the global biological environment and humanity. During the last few decades, various methods such as fluid extracting, Soxhlet extraction and microwave-assisted extraction have been linked to invigilating EDC in soil or, on the other hand, residual conditions [3]. In a couple of these methods, unpredictable natural solvents were used as the extractants, and the bulk of these solvents have a high toxic content, posing a threat to biological communities and humanity. Ionic liquids are a comparatively recent group of solvent that are frequently thought to be less harmful than standard solvents and provide a substitute to chemicals. Human cations and anions make up ionic compounds, which are capable of dissolving a wide spectrum of organic and inorganic compounds [4]. At the micrometer size, physical values that are minor or irrelevant at the full scale may become daunting. The viscous and surface forces are accounted for in the cases. The proximity of surfaces, in particular, means that values such as surface tension and electrokinetic impacts have a greater significance [5]. Isolating and extricating natural contaminants from different matrices, for example, can be accomplished by varying the cation & anion or modifying their sequential, to meet the specifications of numerous technologies [6]. Deng et al. [7] explored the possibility of using imidazolium organic solvents as an energy-efficient green substrate for extracting deep dehydrogenation of ionic liquid. Fluid microextraction and solid stage solvent extraction both employ ionic fluids as a medium [8]. Ionic fluids have efficiently emerged as an excellent dissolvable medium for enzymatic reactions and other nutrient activities [9]. The ionic fluids are made up of ions that maintain their fluid form at temperatures below 1000C or even at room temperature. They've been termed "green solvents" due to their unrivaled qualities, which include a low melting and vapor pressure, thermal stability, minimizing the risk, and reduced negative impacts on the environment [10]. The prerequisite will modify the properties of ILs by changing their constituents (cation and anion) [11]. Protein thermal stabilization is based on the appropriate combination of ILs, that they deposited the protein throughout a high temperature. The spinning disc flow has a long history in the world of computing, with many uses. For example, in processors, heat convection affects the temperature of adjacent electronic components, causing a flow near a spinning hard disc. The invention of centrifugal spinners, external compressors, plate hydrometer, and crystalline processes

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employed in machine memory highlighted the importance of turning disc flows in the mechanic's field Polymer ejection, fluid film accumulation, emulsified coating on photographic films, liquefy swirling, and heat rollers are all examples of boundary extending operations in technology and industries. A thin polymer layer on a constantly changing surface creates a nonuniform pace. The rotating disc electrode device is mostly used to scrutinize the reaction kinetics that has a mass propagation impairment [12]. The MHD, iron ferrite and CNTs hybrid nanofluid flow over an infinite gyrating disc, was studied by Tassaddiq et al. [13]. They reported that increasing the disc spinning velocity significantly increased the thermal energy. The Bingham fluid model was used by Mustafa et al.[14] to investigate viscous dissipation and heat transfer for twirling viscoelastic fluid flow over a permeable rotating disc. A faster thermal efficiency can be achieved by using a higher suction effect at the surface of the disc. The mass and heat flux hypotheses in Oldroyd-B fluid flow over an expanding disc were investigated by Hafeez and Khan [15]. It's worth noting that the solutal relaxation time coefficient has a negative effect on the rate of mass distribution. Gul et al. [16] used many instances to model the hybrid nanofluid flow between disc and cone. From the study, it was concluded that if the surface temperature remained constant, a spinning disc with a fixed cone could achieve the results for different instruments. Fractional evaluations for Darcy flow over a rotating porous disc were recorded by Li et al. [17]. They used the slip conditions at the spinning disk's tip. Shah et al. [18] examined the performance of effectiveness of chemical catalytic reactors and copper selenide with magnetic field by using MATLAB package bvp4c and RK4 Scheme. The heat transference with the natural convection of hybrid nanofluid flow in a quadrilateral cavity attached with two animated fins is calculated by Sowmya et al.[19] and Xia et al. [20]. Zaydan et al. [21] thoroughly considered the thermostability of a conducting nanofluid with an outside applied magnetic field as well as vertical flow. During their time modeling any technical breakthroughs, the scientists run into extremely nonlinear BVPs that aren't easily dealt by a straightforward computational approach. To solve BVPs, MATLAB provides the solvers bvp4c and bvp5c. In certain cases, the solvers should be used interchangeably. The key distinction between the solvers is that bvp4c uses a fourth-order algorithm, while bvp5c uses a fifth-order formula. The bvp5c is based on the Lobatto IIIa formula. The algebraic equations are solved directly by bvp5c, and the scheme is supplemented with trivial differential equations to account for undefined parameters [22, 23]. Kumar et al.[24] Khan used a Runge-Kutta-based shooting approach to investigate the energy and mass transport behavior of chemical reactive Maxwell & MHD Casson fluids through a stretched sheet with an interior heat source. Nazir et al. [25] used the finite element technique to explore the effect of nanocrystals on the thermal efficiency of a partly ionized non-Newtonian liquid subjected to magnetic field and thermal radiations. Gopal et al. [26] computed the thermophysical parameters of elevated chemical operations and viscous dissipation on ferrofluid with a continually extending porous sheet using the numerical method RK4 and the shooting methodology. Kumaran et al. [27] used the RK premised shooting mechanism to calculate magnetohydrodynamic thermally reacting Maxwell and Casson fluids past a stretched sheet. Zubaidi et al. [28] and Cui et al. [29] computational reported the role of metallic nanomaterials in atomic fluid phase transitions.

We evaluated an incompressible, axially symmetric 3D ionic flow characteristic over a perforated gyrating disc in the present issue The current work intends to establish a numerical simulation for ionic transportation over a revolving disc. For ionic maintenance of species,



the Nernst-Planck, Maxwell and Poisson’s equations are related and simulated. Through the resemblance approach, the modelled equations of the electroviscous boundary layer flow are concentrated to a dimensionless system of ODEs. The Matlab built-in package bvp5c has been implemented for the numerical outcomes of the transform equations. In the coming segment, the problem has been expressed, resolved and discoursed.

2. FORMULATION OF THE PROBLEM

We assumed an incompressible ionic fluid flow over a porous surface of a rotating disc. The disk is spinning with an angular velocity as revealed in Figure 1. The fluid is assumed to be of an electrolyte nature, while considering fixed density and thickness Hence, in this sense, the cations (+) and anions (-) have parallel valences $z = 1$. It has been considered that the fluid is infinite and expands in the +ive z-direction. The modeled equations of the flow, electric and magnetic field, and ionic transition are supposed to be dimensionless by using speed \vec{V} , length R^* , time $\frac{R^*}{\vec{V}}$ thickness of cations and anions n_0 Here, T is the temperature, k is the Boltzmann coefficient and e are the elementary charge. The dimensionless equations are expressed as [30]:

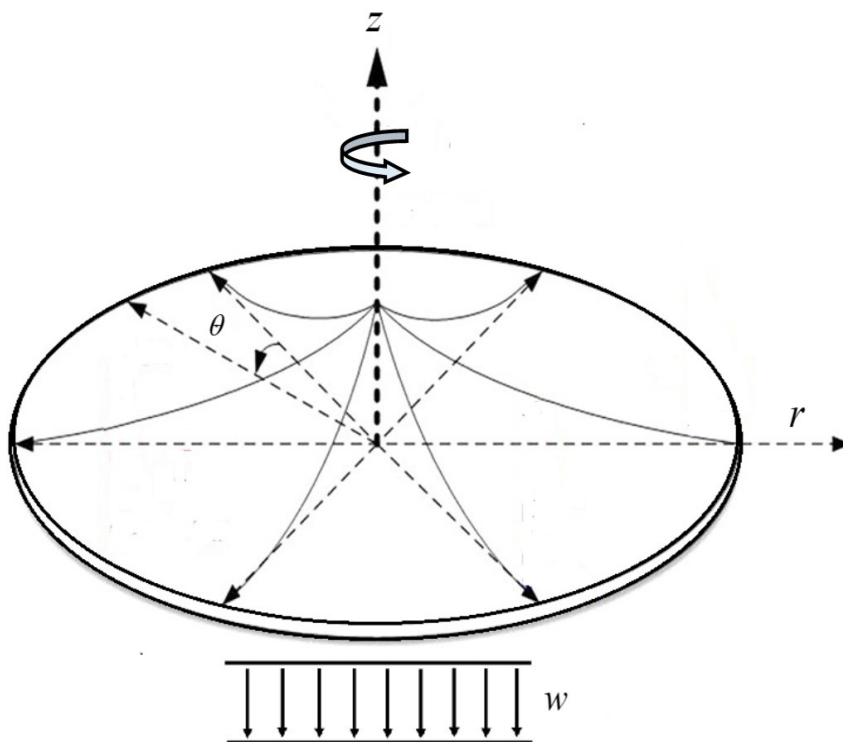


FIGURE 1. Physical sketch of the spinning cone



$$\nabla \cdot V = 0, \quad (2.1)$$

$$\frac{\partial V}{\partial T} + (V \cdot \nabla)V = -\nabla P + \frac{1}{Re} \nabla \cdot (\nabla V + (\nabla V)^T) - \frac{BK^2}{Re^2} (n_+ - n_-) \nabla U, \quad (2.2)$$

$$\nabla^2 U = -\frac{1}{2} K^2 (n_+ - n_-), \quad (2.3)$$

$$\frac{\partial n_+}{\partial t} + \nabla \cdot (V n_+) = \frac{1}{Re Sc} (\nabla^2 n_+ + \nabla \cdot (n_+ \nabla U)), \quad (2.4)$$

$$\frac{\partial n_-}{\partial t} + \nabla \cdot (V n_-) = \frac{1}{Re Sc} (\nabla^2 n_- + \nabla \cdot (n_- \nabla U)), \quad (2.5)$$

$$\rho \frac{\partial B}{\partial t} = \rho \nabla \times (V \times B) + \frac{\rho}{\sigma \mu_2} \nabla^2 B. \quad (2.6)$$

The dimensionless terms in eq (1-6) are:

$$Re = \frac{\rho \vec{V} R^*}{\mu_0}, Sc = \frac{\mu_0}{\rho D}, B = \frac{\rho k^2 T^2 \varepsilon_0 \varepsilon}{2 Z^2 e^2 (\mu_0)^2}, K^2 = \frac{2 Z^2 e^2 n_0 R^{*2}}{\varepsilon_0 \varepsilon k T}, R = \frac{d^2}{V}, R_2 = \frac{\alpha d^2}{2 \nu}. \quad (2.7)$$

The parameter K is the inverse Debye length, Sc is the Schmidt number, Re is the Reynold number, B is a fixed constant used for a given at certain temperature respectively. Where, ε_0 and ε is permeability of free space and dielectric constant. R is the rotating Reynolds number, R_2 is the magnetic based Reynolds number. In which, p_0 and q_0 are the dimensional quantities, which are used to make the B_r and B_z dimensionless. The boundary conditions for the fluid flow:

$$u = -\frac{\alpha r \gamma}{1 - \alpha t}, v = \frac{r \Omega_1}{1 - \alpha t}, w = 0, n_+ = n_- = \frac{\alpha}{1 - \alpha t}, \nabla \cdot V n_0 = S \quad \text{at } Z = 0, \\ u \rightarrow 0, U \rightarrow 0, v \rightarrow 0, n_+ = n_- \rightarrow 0 \quad \text{as } z \rightarrow \infty \quad (2.8)$$

Equation (1) & (2) are the Navier-Stokes equations with the addition of electrical charge as a body force, due to free charge in the momentum equation. The equation (3) is Poisson relation, which express the electric potential U . The aggregate potential at the uniform disk surface can be rebound as $U = \psi(r) - E\phi(z)$ in which, $\psi(r) = \frac{r^2}{R^2(1-\alpha t)}$ where, E and $\phi(z)$ are the uniform electric field and axial potential energy. Equation (4) & (5) are the Nernst-Plank equations, describing the conservation of particles (n_+, n_-). Equation (6) is the Maxwell equation. Equation (7) displays the boundary conditions, in which $S = \frac{ze\sigma R}{\varepsilon_0 \varepsilon k T}$ is the surface charge density and (n_0) is the external unit vector.



2.1. SIMILARITY TRANSFORMATION

The following similarity framework has been utilized to modify the system of PDEs to a system of ODEs:

$$\begin{aligned}
 u &= \frac{\alpha r f'(\eta)}{2(1-\alpha t)}, & v &= \frac{\Omega r g(\eta)}{1-\alpha t}, & w &= \frac{\alpha R f(\eta)}{\sqrt{1-\alpha t}}, & \eta &= \frac{z}{R\sqrt{1-\alpha t}}, \\
 B_r &= \frac{\alpha r p_0}{2R(1-\alpha t)} p'(\eta), & B_z &= \frac{q_0 q(\eta)}{\sqrt{1-\alpha t}}, & U &= -\frac{r^2}{R^2(1-\alpha t)} E H(\eta), \\
 n_+ &= \frac{\alpha}{(1-\alpha t)} m'(\eta), & n_- &= \frac{\alpha}{(1-\alpha t)} n(\eta).
 \end{aligned}
 \tag{2.9}$$

By implementing equation (9) to equations (1-8), we get:

$$\begin{aligned}
 f'''(\eta) &= R_2 \left(\frac{\eta}{4} f''(\eta) + \frac{1}{2} f'(\eta) + \frac{1}{4} f'^2(\eta) - \left(\frac{R}{R_2} \right)^2 g^2(\eta) - \frac{1}{2} f(\eta) f''(\eta) \right. \\
 &\quad \left. + 2 \left(\frac{R}{R_2} \right)^2 g^2(\eta) + \frac{2Bk^2}{\alpha R^2 Re^2} \right),
 \end{aligned}
 \tag{2.10}$$

$$g''(\eta) = R_2 \left(\frac{\eta}{2} g'(\eta) + g(\eta) + f'(\eta)g(\eta) - f'(\eta) - f(\eta)g'(\eta) \right),
 \tag{2.11}$$

$$f''(\eta) = R_2 \left(\frac{\eta}{2} f'(\eta) + \frac{1}{2} f(\eta) - f(\eta)f'(\eta) + \frac{Bk^2 E}{\alpha R^2 Re^2} (m'(\eta) - n(\eta)h'(\eta)) \right),
 \tag{2.12}$$

$$\begin{aligned}
 m''(\eta) &= R^2 \alpha Re Sc \left(\frac{\eta}{2} m''(\eta) - m'(\eta) - f(\eta)m''(\eta) + Em''(\eta)h'(\eta) \right. \\
 &\quad \left. + Em'(\eta)h''(\eta) - 4m'(\eta) \right),
 \end{aligned}
 \tag{2.13}$$

$$n''(\eta) = R^2 \alpha Re Sc \left(\frac{\eta}{2} n'(\eta) + n(\eta) - f(\eta)n'(\eta) + 4n(\eta) - Eh'(\eta) \right),
 \tag{2.14}$$

$$Eh''(\eta) = -\frac{1}{2} k^2 R^2 (m'(\eta) - n(\eta)) - 4,
 \tag{2.15}$$

$$\begin{aligned}
 p'''(\eta) &= \frac{2R_2}{R} \eta Bt p''(\eta) + \frac{4R_2}{R^2} Bt p'(\eta) - \frac{2Bt}{R} f'(\eta)h'(\eta) - \frac{2Bt}{R} f''(\eta)h(\eta) \\
 &\quad - \frac{4R_2 Bt}{R} f'(\eta)p''(\eta) - \frac{2(R_2)^2 Bt}{R} f'(\eta)p'(\eta),
 \end{aligned}
 \tag{2.16}$$

$$q'' = R_2 Bt q'(\eta)\eta + R_2 Bt q(\eta) + 2R_2 Bt q(\eta)f'(\eta) + \frac{2(R_2)^2 Bt}{R} f(\eta)p'(\eta).
 \tag{2.17}$$

the modify conditions are

$$\begin{aligned}
 f(0) &= 0, & f'(0) &= -2\gamma, & g(0) &= m(0) = n(0) = 1, & p'(0) &= q(0) = 1, & h'(0) &= -\frac{RS}{E}, \\
 g(\infty) &\rightarrow 0, & f'(\infty) &\rightarrow 0, & m(\infty) &\rightarrow 0 & n(\infty) &\rightarrow 0, & p'(\infty) &\rightarrow 0 & q(\infty) &\rightarrow 0, & h'(\infty) &\rightarrow 0.
 \end{aligned}
 \tag{2.18}$$



3. PROBLEM SOLUTION

The following variables are introduced in equations (10)-(17) and (18):

$$\begin{aligned} f = l_1, f' = l_2, f'' = l_3, g = l_4, g' = l_5, m = l_6, m' = l_7, m'' = l_8, n = l_9, \\ n' = l_{10}, h = l_{11}, h' = l_{12}, p = l_{13}, p' = l_{14}, p'' = l_{15}, q = l_{16}, q' = l_{17}. \end{aligned} \quad (3.1)$$

The obtained system of nonlinear 1st order ODEs is:

$$\begin{aligned} l'_1 &= l_2, l'_2 = l_3, \\ l'_3 &= R_2 \left(\frac{\eta}{2} l_3 + \frac{1}{2} l_2 + \frac{1}{4} l_2^2 - \left(\frac{R}{R_2} \right)^2 l_4^2 - \frac{1}{2} l_1 l_3 + 2 \left(\frac{R}{R_2} \right) l_4^2 + \left(\frac{2Bk^2}{\alpha R^2 Re^2} \right) \right), \\ l'_5 &= R_2 \left(\frac{\eta}{2} l_5 + l_4 + l_2 l_4 - l_2 - l_1 l_5 \right), \\ l'_8 &= R^2 \alpha Re Sc \left(\frac{\eta}{2} l_8 - l_7 - l_1 l_8 + El_8 l_{11} + El_7 l_{12} - 4l_7 \right), \\ l'_9 &= R^2 \alpha Re Sc \left(\frac{\eta}{2} l_9 + l_8 - l_1 l_9 + 4l_8 - El_4 \right), \\ l'_{11} &= -\frac{1}{2} k^2 R^2 (l_7 - l_8) - 4, \\ l'_{14} &= \frac{2R_2}{R} \eta B t l_{14} + 4 \frac{R_2}{R^2} B t l_{13} - 2 \frac{Bt}{R} l_3 l_{11} - 2 \frac{Bt}{R} l_3 l_{10} - 4 \frac{R_2 Bt}{R} l_2 l_{14} - 2 \frac{R_2^2 Bt}{R} l_2 l_{13}. \end{aligned} \quad (3.2)$$

The boundary conditions are:

$$\begin{aligned} l_2(0) = -2\gamma, l_4(0) = 1, l_6(0) = l_8(0) = 1, l_{11} I(0) = \frac{-RS}{E}, l_{13}(0) = l_{15}(0) = 1, \\ l_2(\infty) \rightarrow 0, l_4(\infty) \rightarrow 0, l_6(\infty) = l_8(\infty) \rightarrow 0, l_{11}(\infty) \rightarrow 0, l_{13}(\infty) = l_{15}(\infty) \rightarrow 0. \end{aligned} \quad (3.3)$$

4. RESULT AND DISCUSSION

4.1. ANALYSIS OF RESULT

In the present mathematical model, we have formulated the Nernst-Planck equation, Maxwell equation and Poisson's equations along with the well-known Navier Stokes equation for the ionic fluid transition. The ion-ion interactions have a huge implementation in numerous mathematical models specially in the electro-chemistry. The current mathematical model is a tiny effort towards ionic liquid models, which have a broad range of applications not only in fluid dynamics, but in the fields of electro chemistry as well. For the solution of ionic fluid model, the Matlab built-in package bvp5c is used. Then any other numerical scheme, bvp5c have fast convergence ability.

4.2. DISCUSSION OF RESULT

Spinning disk flow of an ionic species has been calculated under the consequences of uniform electric and magnetic field at and ions charged surfaces via Figures (2-7). Figure 1 elaborate an ionic flow mechanism over a spinning disk. Figure 2(a)-(d) revealed the behavior of positive charge profile $m'(\eta)$ against parameter S (surface charge density),



rotational Reynold number R , Reynolds number Re and Schmidth number Sc respectively. It can be perceived that the positive charge profile is reduces with the increment of parameter S . At the disk surface ($\eta = 0$) *ive* charge profile remain same, because disk circulate all the charge ions with itself and eventually charge density relics unchanged. We can notice that the formulation of $S = \frac{ze\sigma R}{\epsilon\epsilon_0 kT}$ is in inverse relation with $\frac{kT}{ze}$ (electrical potential), which indicates that the positive charge profile decreases as the electric potential drops as shown in Figure 2(a). Increasing Reynolds number leads to a reinforcement of the inertial forces over the frictional viscous forces. The positive charge profile $m'(\eta)$ reduces with the action of rotational Reynold number R , local Reynold number Re , Schmidth number Sc as revealed through Figure 2(c)-(d).

Figure 3(a)-(d) revealed the behavior of negative charge profilen(η) against surface charge density S , rotational Reynold number R , Reynolds number Re and Schmidth number Sc respectively. In contrast to positive charge density, negative charge density develops at first, then declines until it approaches zero under the influence of above-mentioned parameters. $S = \frac{ze\sigma R}{\epsilon\epsilon_0 kT}$ Here, the rising effect of surface charge density encourage the electric conductivity of the fluid, as a result the negative charge strength reduces, as shown in Figure 3(a). The negative charge profile $n(\eta)$ reduces with the action of rotational Reynold number R , local Reynold number Re , Schmidth number Sc as displayed via Figure 3(c)-(d). The Reynold number Re effect the density of fluid flow, which decreases the negative charge profile as illustrated through Figure 3(b)-(c). Furthermore, the molecular diffusion rate decreases, while kinetic viscosity of fluid enhances with rising effect of Schmidth number, as a result negative charge profile $n(\eta)$ declines Figure 3(d).

Figure 4(a)-(d) illustrated the consequences of several physical entities on radial $f'(\eta)$ velocity. Figure 4(a) elaborate the dominance of parameter S on the velocity distribution. The system is consisting of anion (negative charge) and cation (positive charge) in fluid flow over spinning disc. It has been observed that at less charge density the fluid flows occurred only by the action of electro osmosis. At normal surface density, fluid flow manifest maximum. While, on high charge surface causes the absorption and immobilization of ions near the disc surface. The rising effect of Schmidth number Sc , reduces the thermal diffusivity of fluid flow, which improves the radial velocity profile $f'(\eta)$ as presented in Figure 4(b). Figure 4(c) expressed parameter B impact on velocity profile $f'(\eta)$ as presented in Figure 4(b). Figure 4(c) expressed parameter B impact on velocity profile $f'(\eta)$ B is mostly kept constant for a proposed fluid at a specific temperature. The increment in B reduces the fluid velocity propagation. The behavior of radial velocity profile versus the magnetic Reynolds number is depicted via Figure 4(d). Physically, the positive effect of R_2 , enhances the induction and advection upshot on the fluid flow, which become the cause of reduction in radial velocity profile.

Figure 5(a)-(d) elaborated the nature of axial velocity versus many physical constraints. The same impact of charge density S has been perceived as Fig. 4(a) against axial velocity in Fig. 5(a). The electro-osmosis effect combines with the surface density enhances the axial velocity $f(\eta)$ Figure 5(b) displays the B parameter consequences on the velocity profile. The reverse flow effect has been noticed with an increases of B term. The retarding behavior of velocity versus magnetic force E is communicated in Fig. 5(c). Physically, the magnetic Lorentz forces increase due to the interaction between the electrically conducting fluid and the velocity of the flow which tends to oppose the upward convective motion of the fluid. Consequently, increase the thermal stability of the fluid, that is why, such phenomena have been perceived. Fig. 5(d) is drawn to exhibit the axial



velocity against Debye length coefficient K . Because the potential gradient is not zero near the disk surface, and almost equals to the S . That's the reason, that an increment K enhances the potential gradient, Because EDLs account up a much smaller fraction of the disc surface and expose more counter-ions to the flow, that improves the axial velocity component.

The influence of parameter S, R_2, E & R are displayed respectively via Fig. 6(a)-(d). Magnetic field strength is always proportional to the surface charge density. The hydration layer is more stable to thermal agitation when the surface charge density is higher. Because the viscous fluid around the ions is polarized to some extent by high charge density, minor ions encounter more resistible forces than bigger ions with lower charge density as observed in Fig. 6(a). Figure 6(b) depicts the reduction in tangential velocity as the magnetic Reynolds number R_2 rises. Physically, increasing the magnetic Reynolds number minimizes the fluid's kinematic viscosity and improves the system's linear dimension, which causes in lowering the fluid's velocity. Figure 6(c) shows how tangential velocity decreases as magnetic strength E grows. The E field provides resistance to the flow profile, causing velocity to drop. Obviously, when the Reynolds number rises, the tangential velocity of the fluid rises, and if the R rises without bounds, the laminar flow will shift to the turbulent flow seen in Figure 6(d).

Figure 7(a)-(d) demonstrated the magnetic strength profile behavior along axial $p'(\eta)$ and radial $q(\eta)$ direction for Batchlor number Bt , rotational Reynold number R , Batchlor number Bt and magnetic Reynold number R_2 respectively. The effect of Batchlor number Bt enhances the axial magnetic strength $p'(\eta)$ profile as illustrated in Fig. (a) & (c). Batchlor number is expressed as $Bt = \sigma\mu_0 2$ it measures the fluid slips across the magnetic field. A low current creates a big induced magnetic field when Bt is large. When Bt is weak, however, a moderate-sized current will only create induced fields, which have a minor effect on the applied field. While the Reynold number declines the magnetic strength profile, in both axial and radial magnetic strength profile as shown in Fig. 7(d).



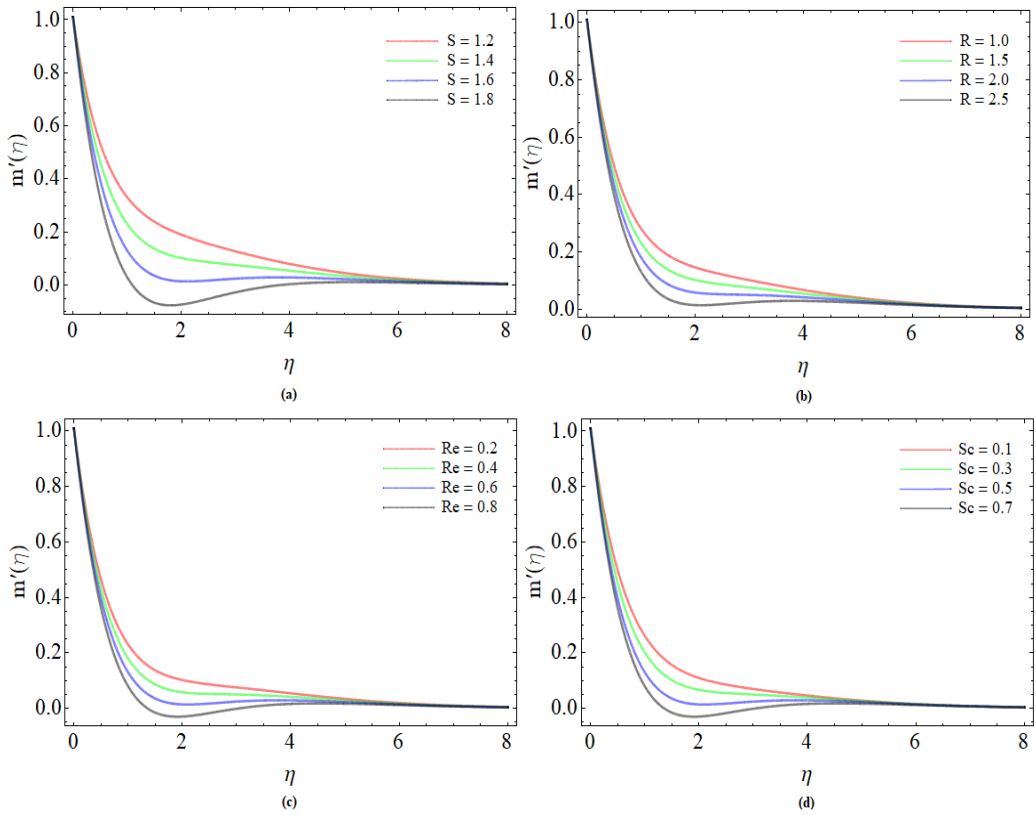


FIGURE 2. Positive charge profile $m'(\eta)$ for (a) surface charge density (b) rotational Reynolds number (c) Reynolds number (d) Schmidt number.



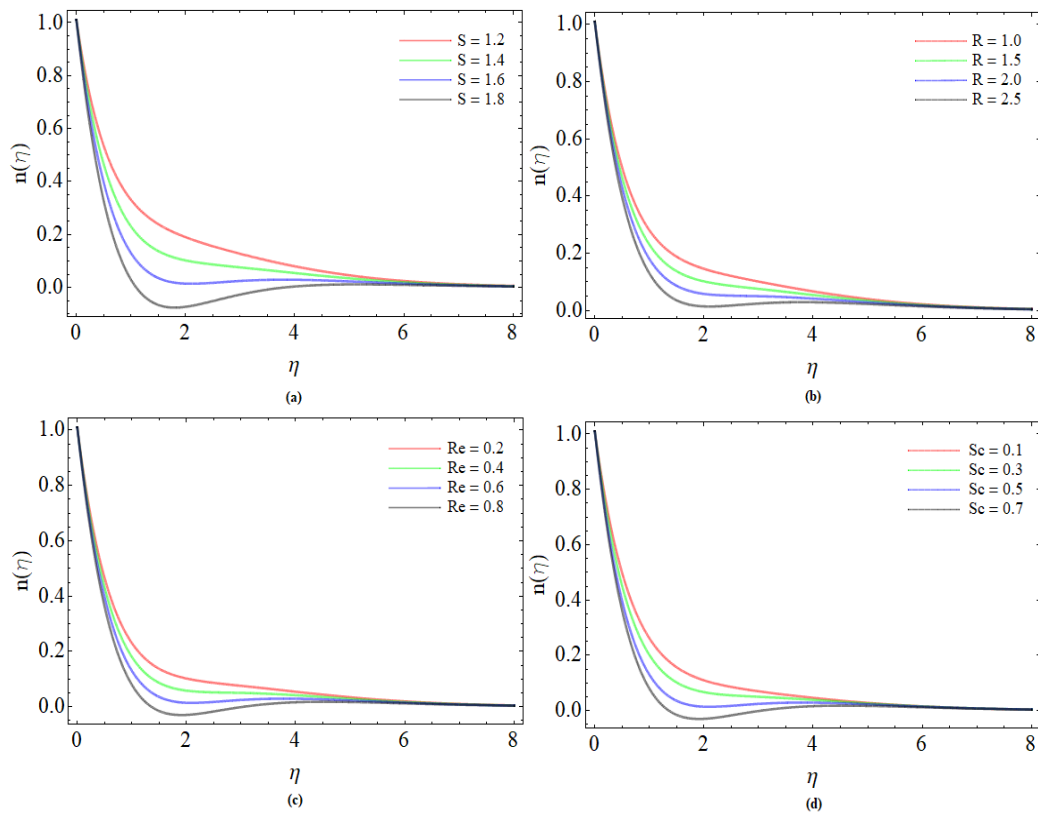


FIGURE 3. Negative charge profile for $n(\eta)$ (a) surface charge density (b) rotational Reynolds number (c) Reynolds number (d) Schmidt number.



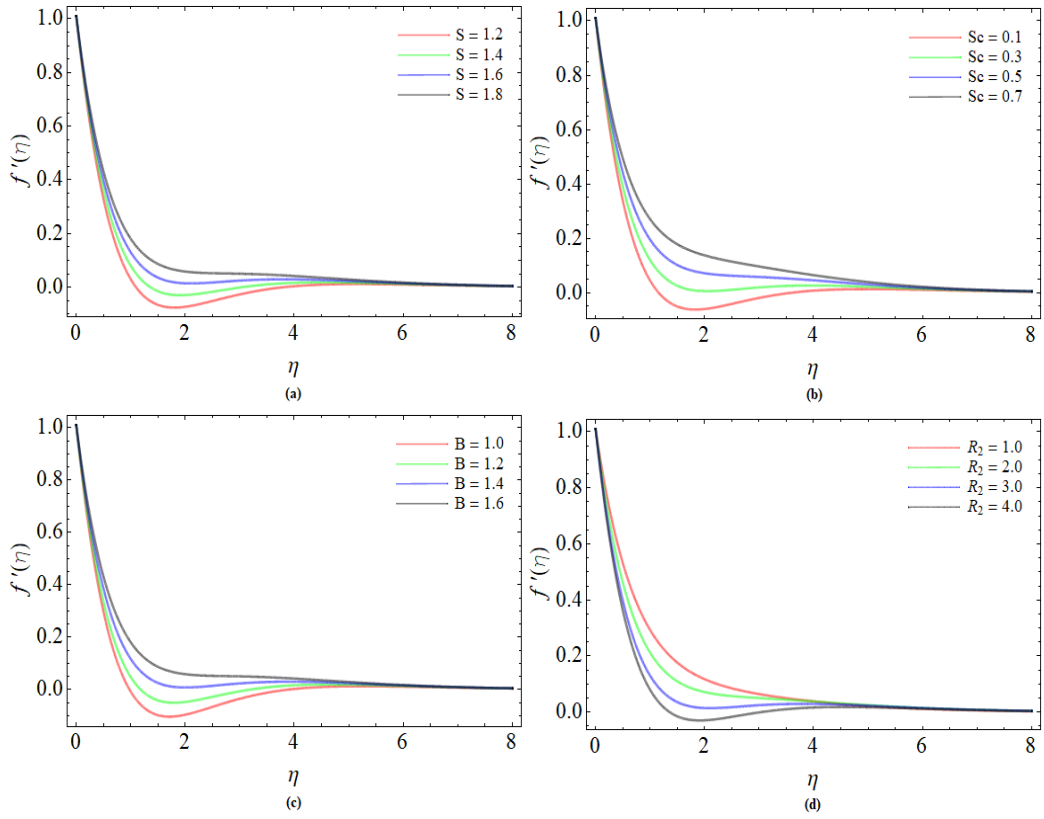


FIGURE 4. Radial velocity profile for $f'(\eta)$ (a) surface charge density (b) Schmidt number (c) Fixed constant (d) magnetic Reynolds number.



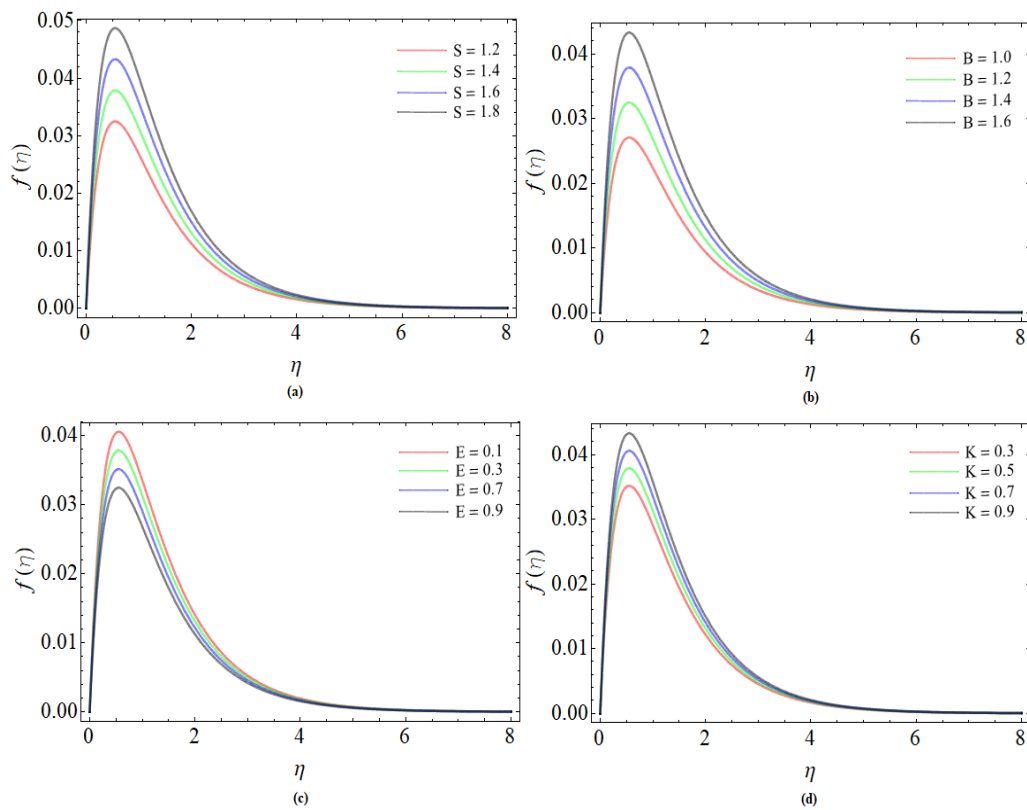


FIGURE 5. Axial velocity profile for $f(\eta)$ (a) surface charge density (b) Fixed constant (c) magnetic strength (d) Debye length.



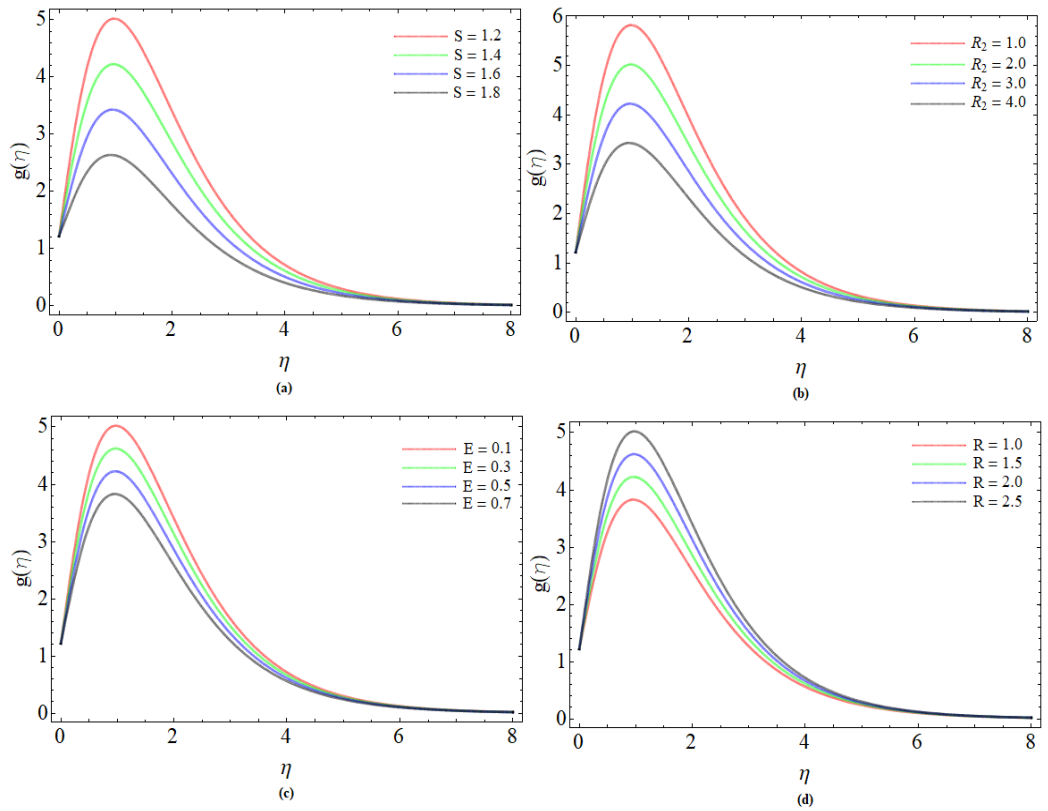


FIGURE 6. Tangential velocity profile for $g(\eta)$ (a) surface charge density (b) magnetic Reynolds number (c) magnetic strength (d) Reynolds number.



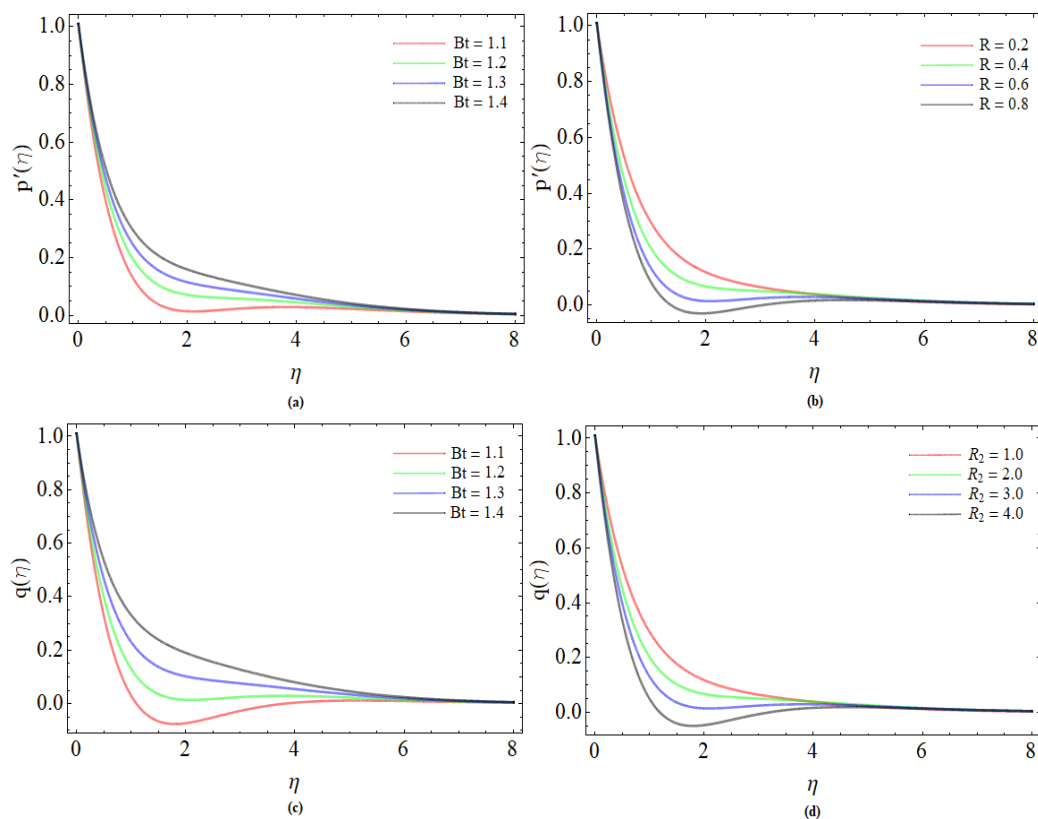


FIGURE 7. The magnetic strength profile behavior along axial $p'(\eta)$ and radial $q(\eta)$ direction for (a) Batchlor number (b) rotational Reynold number (c) Batchlor number (d) magnetic Reynold number.

5. CONCLUSION

We evaluated an unsteady, axisymmetric 3D ionic fluid flow over a permeable spinning disc in the present dilemma. The current effort aspires to establish a numerical analysis for ionic transport in an electroviscous boundary layer flow across a spinning disc. For ionic study the Maxwell equation and Poisson's equation are linked and simulated with the well-known Navier Stokes equations. The following conclusions were reached as a consequence of the aforementioned study and calculation:

- The electrical potential and the surface charge density S are inversely related. As a result, when the electric potential drops, the positive charge per unit volume improves. Work done against an electric field produces electric potential energy.
- The absorption and immobilization of counter ions at the disc surface due to a high charged surface diminishes the radial velocity.
- The Lorentz effect is generated inside the boundary layer by raising the magnetic intensity E , which causes some flow field deceleration, resulting in axial velocity reductions.



- Fluid flows mainly by electro osmotic pressure mechanism at lower charge densities. However, flow is optimum when the surface density is normal.
- The magnetic strength profiles both in axial $p'(\eta)$ and radial $q(\eta)$ direction enhances with the effect of Batchlor number Bt , while the increment in Reynold number declines the magnetic strength profiles.

Competing Interest Statement: No competing interest exists.

Data Availability: The relevant data exist in the article.

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