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Comparative Numerical Analysis of Heat and Mass Transfer Characteristics in Sisko Al₂O₃-Eg and TiO₂-Eg Fluids on a Stretched Surface

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ABSTRACT

In the current research, a thorough examination unfolds concerning the attributes of magnetohydrodynamic (MHD) boundary layer flow and heat transfer inherent to nanoliquids derived from Sisko Al₂O₃-Eg and TiO₂-Eg compositions. Such nanoliquids are subjected to an extending surface. Consideration is duly given to slip boundary conditions, as well as the effects stemming from variable viscosity and variable thermal conductivity. The analytical approach applied involves the application of suitable similarity transformations. These conversions serve to transform the initial set of complex nonlinear partial differential equations into a more manageable assembly of ordinary differential equations. Through the utilization of the FEM, these reformulated equations are solved, considering the specified boundary conditions. The outcomes attained are graphically depicted by means of plots and tables. These visual aids facilitate a comprehensive exploration of how diverse parameters exert influence over the distributions of velocity, temperature, and concentration. Furthermore, detailed scrutiny is directed towards the fluctuations characterizing pivotal parameters, viz., Nusselt number, skin-friction coefficient, and Sherwood number. It is identified that the Nusselt number showcases a diminishing trend coinciding with increasing values of the volume fraction parameter (ϕ). This trend remains consistent regardless of whether the nanoliquid under consideration is Al₂O₃-Eg or TiO₂-Eg based. In contrast, both the skin-friction coefficient and Sherwood number assume lower values as the volume fraction parameter (ϕ) escalates. This pattern remains congruent across both classifications of nanoliquids. The findings of the study impart valuable insights into the complex interplay governing the characteristics of HMT pertaining to Sisko Al₂O₃-Eg and TiO₂-Eg nanoliquids along an extending surface.



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KEYWORDS

MHD boundary layer flow; heat transfer; Sisko Al₂O₃-Eg and TiO₂-Eg nanoliquids; slip boundary conditions; variable thermal conductivity; nusselt number

Nomenclature

\mathbf{B}_0	Magnetic field strength
В	Sisko fluid parameter
\mathbf{B}_2	Variable thermal viscosity
B ₃	Variable thermal conductivity
Sh_x	Sherwood number
$C_{\rm f}$	Skin-friction coefficient
Cr	Chemical reaction parameter
\mathbf{D}_{m}	Mean fluid concentration
$f(\eta)$	Dimensionless stream function
FEM	Finite element method
G	Vertical velocity
\mathbf{J}_{w}	Wall mass flux, Jw
Κ	Permeability parameter, H/m
Μ	Magnetic parameter
Nu _x	Nusselt number
ODE	Ordinary differential equations
PDE	Partial differential equations
Pr	Prandtl number
q _r	Radiative heat flux, W/m ²
Ŕ	Radiation parameter
Ra _a	Rayleigh number
\mathbf{V}_0	Suction parameter
σ	Electrical conductivity
ψ	Stream function
μ	Fluid viscosity, Pa.s

1 Introduction

At present, providing efficient thermal fluids for heat transfer and cooling purposes, particularly in engineering practice, has become increasingly challenging. Therefore, it remains a significant undertaking for researchers. Various approaches can be employed to enhance heat transfer. The thermal conductivity of any nanofluid is affected by the type, size, and shape of the nanoparticles involved. The primary purpose of nanoliquids is to enhance the thermal conductivity within heat transfer fluids. Generally, nanoliquids are created by suspending nanoparticles in base fluids such as water, oil, kerosene, and so on. Nanoliquids represent the cutting edge of technological advancement in this field. The extensive practical implications of nanotechnology have sparked enthusiasm across various technological and industrial domains. These span sectors like information technology, power generation, ecological studies, healthcare, national security, food integrity, and transportation. Several research studies have contributed to these advancements. Tasawar et al. [1] conducted a classical study on the rotating disk problem, while Ramesh et al. [2] explored the anomalous thermal behavior between two rotating disks. Choi et al. [3] investigated the enhancement of thermal conductivity in nanotube suspensions. Sheremet et al. [4] discussed the influence of a constant magnetic field on mixed convection in a trapezoidal cavity filled with Cu-water nanofluid. Hassan et al. [5] focused on the combined effects of heat and mass transfer (HMT), as well as the impact of thermal radiation, in the flow of sisko nanofluid with gold nanoparticles using blood as the base fluid. These studies contribute to our understanding and application of nanotechnology in improving heat transfer and fluid dynamics in various systems.

In the domain of computational investigation, Sheremet et al. [6] performed research to explore the impact of Brownian diffusion and thermophoresis on natural convection within a water-based nanofluid, constrained within two triangular cavities. The remaining walls of the cavities were maintained at a low constant temperature, while the top and bottom walls were kept at a high constant temperature. Furthermore, Sheremet et al. [7] conducted numerical scrutiny of heat transfer during steady-state free convection within a right-angled triangular porous enclosure that was occupied by a nanofluid, using the Buongiorno model. An evaluation of statistical patterns pertaining to free convection heat conveyance within a parallelogram-shaped enclosure with porous media was specifically undertaken, focusing on Cu-water nanoliquids [8]. Moreover, the utilization of the fifthorder Runge-Kutta-Fehlberg technique in conjunction with a shooting method is deployed to explore the impacts of a consistent magnetic field and nonlinear thermal radiation on the transfer of mass and heat within the flow of Sisko nanofluid over an extending surface [9,10]. These numerical studies contribute to the understanding of heat transfer phenomena in various complex systems and environments.

Convective boundary conditions and thermal radiation were harnessed by Jyothi et al. [11] to explore the characteristics of flow and heat transfer involving single-walled carbon nanotubes (SWCNTs) and multi-walled carbon nanotubes (MWCNTs) positioned amidst two rotating discs. The investigation encompassed diverse combinations of rotating and stretching velocities. Hashim et al. [12] developed a new model to examine the intricacies of HMT in the flow of Carreau nanoliquids. Hayat et al. [13] studied the three-dimensional Darcy-Forchheimer flow of Carreau nanofluid with a convectively heated surface, considering zero nanoparticles mass flux conditions. Raju et al. [14] investigated the flow of nanofluid over a bidirectional stretching surface in a porous medium, incorporating thermal radiation and the effects of non-uniform heat source/sink along with magnetohydrodynamics (MHD) and the Sisko ferro model. Reddy et al. [15] conducted a numerical analysis of HMT in single-phase nanofluid scenarios over rotating cylinders, considering the Cattaneo-Christov heat flux and slip effects. These studies contribute to our understanding of heat transfer phenomena and fluid dynamics in various nanofluid systems under different boundary conditions.

An experimental exploration into the heat transfer and flow attributes of nanoliquids containing multi-walled carbon nanotubes (MWCNTs) within a double-pipe heat exchanger was carried out by Moradi et al. [16]. Hashim et al. [17] engaged in numerical investigation of the HMT characteristics of Williamson nanofluid over a wedge geometry, accounting for temporal variations. Sreedevi et al. [18] delved into the analysis of HMT within a vertical cone, incorporating both single- and multi-wall carbon nanotubes, and incorporating convective boundary conditions. Li et al. [19] analyzed HMT in magnetohydrodynamic (MHD) Williamson nanofluid over an exponentially porous stretching surface, noting the effects of thermophoresis and Brownian motion parameters on heat transmission rate. Veera et al. [20] investigated MHD fluid flow of nanoliquids through a vertically traveling absorbent plate in the existence of radiation absorption. Azmi et al. [21] examined heat transmission and friction concerns in the turbulent flow of titanium oxide (TiO₂) and aluminum oxide (Al₂O₃)

nanoliquids within a tube. An artificial neural network (ANN) framework was introduced by Ahmadloo et al. [22] to anticipate the thermal conductivity ratio of diverse nanoliquids concerning the base fluid. This prediction model takes input variables like the nature of nanoparticles and the base fluid into consideration. Investigation into the interplay of heat and mass transfer within an unsteady chemically reactive boundary layer flow was undertaken by Prashar et al. [23]. This study focused on a ZnO-MWCNTs/ethylene glycol hybrid nanoliquid and encompassed the incorporation of a non-Newtonian flow model alongside Arrhenius activation energy effects. The impact of nanoparticle migration on the convective heat transfer coefficient within nanoliquids based on Al₂O₃-EG/water was subject to experimental inquiry by Choi et al. [24]. Nabil et al. [25] conducted empirical research into the heat transfer and friction factor of nanoliquids containing TiO₂-SiO₂, in conjunction with an H₂O:EG mixture. Maddah et al. [26] experimentally examined the heat exchanger performance using Al₂O₃-TiO₂ hybrid nanofluid in the presence of thermal radiation. Urmi et al. [27] discussed the thermophysical properties of 40% EG-based TiO_2 -Al₂O₃ hybrid nanoliquids and highlighted the improved thermal conductivity and low viscosity of hybrid nanoliquids for realistic heat transfer applications. Yıldız et al. [28] evaluated the efficacy of heat transfer within an Al_2O_3 -SiO₂/H₂O hybrid nanofluid, additionally contrasting theoretical and experimental models for thermal conductivity. Several researchers [29–31] have undertaken investigations into the heat transfer attributes of diverse fluids, while considering a range of geometries.

This paper focuses on the analysis of HMT characteristics of MHD Sisko nanoliquid over a stretching sheet, with a specific emphasis on comparing Al_2O_3 -Eg and TiO_2 -Eg nanoliquids. The study takes into account slip conditions and radiation effects. It is noteworthy that this topic has not been previously addressed, making this analysis novel and unique. The outcomes of this study bear immediate relevance in numerous domains, notably in the blending of lubricants, food products, and the fusion of plasma with lubricants and moisturizers.

2 Problem Formulation

Under consideration is a stretching sheet characterized by velocities, viz., $u = \frac{c_1 x}{1 - \beta t}$, $v = \frac{d_1 y}{1 - \beta t}$ within the context of a 3D MHD Sisko nanoliquid. This nanoliquid is formulated utilizing Al₂O₃ and TiO₂ as nanoparticles, while Eg serves as the base fluid. These cases are visually represented in Fig. 1. Observation reveals that u represents the velocity along the x-axis, where $c_1 > 0$ signifies the stretching rate, while v denotes the velocity along the y-axis, where $d_1 > 0$ signifies the stretching rate. Fluid movement is constrained to the z-axis and maintains parallel alignment with the surface. A logical assumption can be made that the temperature of the nanoliquid is higher near the sheet's surface compared to temperatures observed at substantial distances. Moreover, an external magnetic field denoted as B-0 is exerted along the direction of the y-axis. In Table 1, comprehensive information pertaining to the thermophysical characteristics of the nanoliquids is provided. The governing equations that encompass momentum, energy, and concentration factors, accommodating fluctuations in both thermal conductivity and viscosity, can be formulated as in Eqs. (1)–(5), in accordance with the assumptions [3–5].

$$\frac{\partial v}{\partial y} + \frac{\partial v}{\partial y} + \frac{\partial w}{\partial z} = 0 \tag{1}$$

$$u\frac{\partial u}{\partial x} + v\frac{\partial u}{\partial y} + w\frac{\partial u}{\partial z} = \frac{\mu_{nf}}{\rho_{nf}}\frac{\partial}{\partial z}\left(\mu\left(T\right)\frac{\partial u}{\partial z}\right) - \frac{b}{\rho_{nf}}\frac{\partial}{\partial z}\left(-\frac{\partial u}{\partial z}\right)^{n} - \frac{\sigma_{nf}}{\rho_{nf}}B_{0}^{2}u$$
(2)

$$u\frac{\partial v}{\partial x} + v\frac{\partial v}{\partial y} + w\frac{\partial v}{\partial z} = \frac{\mu_{nf}}{\rho_{nf}} \left(\frac{\partial}{\partial z} \left(\mu \left(T \right) \frac{\partial v}{\partial z} \right) \right) - \frac{b}{\rho_{nf}} \frac{\partial}{\partial z} \left(-\frac{\partial u}{\partial z} \right)^{n-1} \frac{\partial v}{\partial z} - \frac{\sigma_{nf}}{\rho_{nf}} B_0^2 v \tag{3}$$

$$u\frac{\partial T}{\partial x} + v\frac{\partial T}{\partial y} + w\frac{\partial T}{\partial z} = \frac{K_{nf}}{\left(\rho_{cp}\right)_{nf}} \left(\frac{\partial}{\partial z}\left(K\left(T\right)\frac{\partial T}{\partial z}\right)\right) - \frac{1}{\left(\rho_{cp}\right)_{nf}}\frac{\partial q_r}{\partial z}$$
(4)

$$u\frac{\partial C}{\partial x} + v\frac{\partial C}{\partial y} + w\frac{\partial c}{\partial w} = D_B \frac{\partial^2 C}{\partial z^2} - K_r(c - c_\infty)$$
(5)



Figure 1: Physical explanation and cartesian geometry of the problem

Kg Fluid C_p k ρ $\overline{m^3}$ gKтK 46 Al_2O_3 3970 765 TiO_2 4250 686 8.90 Eg 1100 2415 0.26

 Table 1: Thermo-physical properties of nanoparticles

The related boundary situations are

at
$$z \to 0$$
,
 $u = u_w = \frac{c_1 x}{1 - \beta t} \frac{\partial u}{\partial z}; v = v_w = \frac{d_1 y}{1 - \beta t}; T = T_w = T_\infty + \frac{c_1 x}{1 - \beta t} \frac{\partial T}{\partial z}; C = C_w = C_\infty + \frac{c_1 x}{1 - \beta t} \frac{\partial C}{\partial z}$ (6)
at $z \to \infty$,

$$u \to 0, v \to 0, T \to T_{\infty}, C \to C_{\infty} \tag{7}$$

To reorganize the mathematical analysis of the problem, we introduce the following similarity transformations:

$$\begin{split} \mathbf{u} &= \frac{c_{1}x}{1-\beta t} \mathbf{f}(\eta), \mathbf{v} = \frac{d_{1}y}{1-\beta t} \mathbf{g}'(\eta), \\ \mathbf{W} &= -c_{1} \left(\frac{c^{n-2}}{\frac{b}{\rho_{f}}} \right)^{\frac{1}{n+1}} \left[\frac{2n}{n+1} \mathbf{f}(\eta) + \eta \left(\frac{1-n}{1+n} \right) \mathbf{f}'(\eta) + \mathbf{g}(\eta) \right] \mathbf{x}^{\frac{n-1}{1+n}} \left(1 - \beta t \right)^{\frac{1-2n}{n+1}}, \\ \eta &= \mathbf{Z} \left(\frac{c^{n-2}}{\frac{b}{\rho_{f}}} \right)^{\frac{1}{n+1}} \mathbf{x}^{\frac{1-n}{1+n}} (1-\beta t)^{\frac{n-2}{n+1}}, \theta(\eta) = \frac{\mathbf{T} - \mathbf{T}_{\infty}}{\mathbf{T}_{w} - \mathbf{T}_{\infty}}; \quad \mathbf{S}(\eta) = \frac{\mathbf{C} - \mathbf{C}_{\infty}}{\mathbf{C}_{w} - \mathbf{C}_{\infty}} \end{split}$$
(8)

The nanoliquid's dynamic viscosity μ_{nf} , density ρ_{nf} , thermal diffusivity α_{nf} , heat capacitance $(\rho c_p)_{nf}$, thermal conductivity k_{nf} , electric conductivity σ_{nf} along with base fluid's kinematic viscosity ν_f are categorized as follows:

$$\begin{split} \mu_{nf} &= \frac{\mu_{f}}{(1-\phi)^{2.5}} (\rho c_{p})_{nf} = (1-\phi) \left(\rho c_{p}\right)_{f} + \phi (\rho c_{p})_{nf}, \rho_{nf} = (1-\phi) \rho_{f} + \rho_{nf} \phi \\ k_{nf} &= k_{f} \left(\frac{(1-\phi) + 2\phi \left(\frac{k_{nf}}{k_{nf} - k_{f}}\right) \ln \left(\frac{k_{nf} + k_{f}}{2k_{f}}\right)}{(1-\phi) + 2\phi \left(\frac{k_{f}}{k_{nf} - k_{f}}\right) \ln \left(\frac{k_{nf} + k_{f}}{2k_{f}}\right)} \right), \frac{\sigma_{nf}}{\sigma_{f}} = 1 + \frac{3 \left(\frac{\sigma_{nf}}{\sigma_{f}} - 1\right) \phi}{\left(\frac{\sigma_{nf}}{\sigma_{f}} - 1\right) \phi}, \alpha_{nf} = \frac{k_{nf}}{(\rho c_{p})_{nf}} \end{split}$$
(9)

Through the utilization of the Rosseland approximation to model radiation effects, the radiative heat flux, q_r can be characterized as follows:

$$q_r = -\frac{4\sigma^*}{3K^*} \frac{\partial T^4}{\partial z} \tag{10}$$

Assuming that the temperature variations within the flow result in an approximation of the T^4 term as a linear function of temperature, this is achieved by expanding T^4 using a Taylor series centered around the free stream temperature T_{∞} which can be characterized as follows:

$$T^{4} = T_{\infty}^{4} + 4T_{\infty}^{3} (T - T_{\infty}) + 6T_{\infty}^{2} (T - T_{\infty})^{2} + \dots$$
(11)

Disregarding terms of higher order in above Eq. (11) past the linear degree in $(T - T_{\infty})$, we get $T^4 \cong 4T_{\infty}^{\ 3}T - 3T_{\infty}^{\ 4}$. (12)

Thus, substituting Eq. (12) into Eq. (10), we get

$$q_r = -\frac{16T_{\infty}^{3}\sigma^*}{3K^*}\frac{\partial T}{\partial z}.$$
(13)

Utilizing Eqs. (8), (9), and (13), the controlling non-linear partial differential Eqs. (1)–(5), in conjunction with the boundary constraints (6) and (7), are simplified to

$$\Pr \mathbf{B}[(1+\epsilon\theta)f'''+\epsilon\theta'f'] - A_1A_2\mathbf{B}n(-f'')^{n-1}f''' + \mathbf{M}A_1f' + A_1A_2\left[\frac{2n}{n+1}f + g\right]f'' - A_1A_2f'^2 = 0$$
(14)

$$\Pr B[(1+\epsilon\theta)g'''+\epsilon\theta'g''] - A_1B(n-1)(-f'')^{n-2}f'''g'' + MA_1g' + A_1A_2\left[\frac{2n}{n+1}f + g\right]g'' - A_1A_2g'^2 = 0$$
(15)

$$\left[(1+\epsilon\theta)\theta''+\epsilon\theta'\right] + \mathbf{R}\mathbf{A}_{4}\mathbf{A}\theta'' + \mathbf{A}_{3}\mathbf{A}_{4}\left[\frac{2n}{n+1}\mathbf{f} + \mathbf{g}\right]\theta' - \mathbf{A}_{3}\mathbf{A}_{4}\mathbf{f}'\theta = 0$$
(16)

$$S'' - Sc Cr S - Sc \phi f' + Sc \left[\frac{2n}{n+1}f + g\right]S' = 0$$
(17)

The renovated boundary conditions are

$$f(0) = V_0, \quad f'(0) = 1 + \lambda f', \quad g(0) = 0, \quad \theta(0) = 1 + \xi \theta', \quad S(0) = 1 + \beta S',$$

$$f'(\infty) \to 1, \quad g'(0) = 0, \quad \theta(\infty) \to 0, \quad S(\infty) \to 0$$
(18)

where, main denotes ordinary differentiation with respect to η . In conventional notation,

$$\begin{aligned} \Pr &= \frac{\mu_{f}}{\rho_{f}}, M = \frac{\sigma B_{0}^{2}}{\rho_{f}} \left(\frac{1 - \beta t}{c_{1}} \right), k_{1} = \frac{\nu_{f}}{2K\omega_{1}}, R = \frac{16T_{\infty}^{3}\sigma^{*}}{3K^{*}k_{f}}, A_{1} = (1 - \varphi)^{2.5}, \\ \lambda &= \left(\frac{c_{1}}{1 - \beta t} \right) Re_{b}^{\frac{1}{n+1}}, \xi = \left(\frac{c_{1}}{1 - \beta t} \right)^{2} \frac{Re_{b}^{\frac{1}{n+1}}}{(T_{w} - T_{\infty})}, A_{2} = (1 - \varphi) + \varphi \left(\frac{\rho_{s}}{\rho_{f}} \right), A = \frac{Re_{b}^{\frac{2}{n+1}}}{Re_{a}}, \tau = \frac{\beta}{c_{1}}, \\ Sc &= \frac{1}{D_{B}}, C_{r} = \frac{k_{r}(1 - \beta t)}{c_{1}}, \beta = \left(\frac{c_{1}}{1 - \beta t} \right)^{2} \frac{Re_{b}^{\frac{1}{n+1}}}{(C_{w} - C_{\infty})}, A_{3} = (1 - \varphi) + \varphi \left(\frac{(\rho C_{p})_{s}}{(\rho C_{p})_{f}} \right), A_{4} = \frac{k_{f}}{k_{nf}}, \\ q &= \frac{16T_{\infty}^{3}\sigma^{*}}{3K^{*}k_{f}}, Re_{a} = \frac{U_{w}x\rho_{f}}{a}, Re_{b} = \frac{U_{w}^{2-n}x^{n}\rho_{f}}{b} \end{aligned}$$

The skin-friction coefficients in the x and y directions, as well as the local Nusselt number (Nu_x) , and the local Sherwood number (Sh_x) are all important quantities in this field and can be defined as follows:

$$C_{fx} = \frac{\tau_{xz}}{\frac{1}{2}\rho U_w^2}, C_{fy} = \frac{\tau_{yz}}{\frac{1}{2}\rho U_w^2}, Nu_x = -\frac{xq_w}{(T_f - T_\infty)} \left(\frac{\partial T}{\partial z}\right)|_{z=0}, Sh_x = -\frac{xJ_w}{(C_f - C_\infty)} \left(\frac{\partial C}{\partial z}\right)|_{z=0}$$

The afore mentioned in the form of non-dimensional variables are

$$\frac{1}{2}Re_{b}^{\frac{1}{n+1}}C_{fx} = Af''(0) - \left[(-f''(0))\right]^{n},$$

$$\frac{1}{2}Re_{a}^{\frac{1}{n+1}}C_{fy} = \frac{V_{w}}{U_{w}}\left[Ag''(0) + \left[(-f''(0))^{(n-1)}g''(0)\right]\right]$$

$$Re^{-\frac{1}{n+1}}Nu_{x} = -\theta'(0), Re^{-\frac{1}{n+1}}Sh_{x} = -S'(0)$$

3 Numerical Method of Solution

1

The numerical technique known as the finite element method (FEM) [28-30] is utilized for the computation of Eqs. (14)–(17) in conjunction with boundary conditions (18). In contrast to alternative numerical approaches, FEM is recognized as a sophisticated method for precisely and effectively solving both partial and ordinary differential equations.

3.1 FEM

FEM proves highly adept at resolving both regular and partial differential equations. This technique functions by partitioning the complete domain into smaller units referred to as "finite elements," each possessing bounded dimensions as depicted in Fig. 2. This approach is especially fitting for tackling integral equations spanning diverse domains, including heat conduction, fluid dynamics, chemical engineering, electrical networks, and other branches of engineering science.



Figure 2: Stages in FEM

Choosing appropriate form functions to approximate real functions is a pivotal consideration. When tackling the solution of a system of non-linear ordinary differential Eqs. (14)–(17) coupled with boundary conditions (18), the first assumption is that.

$$\frac{\mathrm{df}}{\mathrm{d}\eta} = \mathrm{h} \tag{19}$$

The Eqs. (14) to (17) then reduces to

$$\Pr B[h''(1+\epsilon\theta)+\epsilon\theta'h'] - A_1A_2Bn^{(-h')^{n-1}}h'' + MA_1h + A_1A_2\left[\left(\frac{2n}{1+n}\right)f + g\right]h' - A_1A_2h^2 = 0$$
(20)

$$\frac{\mathrm{d}g}{\mathrm{d}\eta} = \mathbf{k} \tag{21}$$

$$\Pr B[k''(1+\epsilon\theta)+\epsilon\theta'k'] - A_1B(n-1)^{(-h')^{n-2}}h''k' + MA_1k + A_1A_2\left[\left(\frac{2n}{1+n}\right)f + g\right]k' - A_1A_2k^2 = 0 \quad (22)$$

$$\left[(1+\epsilon\theta)\,\theta''+\epsilon\theta \right] + \mathbf{R}A_4A\theta'' + A_3A_4 \left[\left(\frac{2n}{1+n}\right)\mathbf{f} + \mathbf{g} \right] \theta' - A_3A_4h\theta = 0 \tag{23}$$

$$S'' - ScCrS - ScSh + Sc\left[\left(\frac{2n}{1+n}\right)f + g\right]S' = 0$$
(24)

The boundary conditions take the form

$$\begin{split} f(0) &= V0, \quad h(0) = 1 + \lambda h', \quad g(0) = 0, \quad \theta(0) = 1 + \xi \theta', \quad S(0) = 1 + \beta S', \\ h(\infty) &\to 1, \quad k(0) \to 0, \quad \theta(\infty) \to 0, \quad S(\infty) \to 0 \end{split}$$

3.2 Variational Formulation

The variational formulation corresponding to Eqs. (19) to (24) across a standard linear element (η_e, η_{e+1}) is expressed as

$$\int_{\eta_e}^{\eta_{e+1}} w_1 \left(\frac{\mathrm{df}}{\mathrm{d\eta}} - \mathbf{h}\right) \mathrm{d\eta} = 0 \tag{26}$$

$$\int_{\eta_e}^{\eta_{e+1}} w_2 \left(\Pr B[h''(1+\epsilon\theta) + \epsilon\theta'h'] - A_1 A_2 B n^{(-h')^{n-1}} h'' + M A_1 h + A_1 A_2 \left[\left(\frac{2n}{1+n} \right) f + g \right] h' - A_1 A_2 h^2 \right) d\eta = 0$$
(27)

$$\int_{\eta_{e}}^{\eta_{e+1}} w_{3} \left(\frac{\mathrm{d}g}{\mathrm{d}\eta} - \mathbf{k} \right) \mathrm{d}\eta = 0$$

$$\int_{\eta_{e}}^{\eta_{e+1}} w_{4} \left(\Pr B[k''(1 + \epsilon\theta) + \epsilon\theta'k'] - A_{1}B(n-1)^{(-h')^{n-2}}h''k' + MA_{1}k \right)$$
(28)

$$+A_1A_2\left[\left(\frac{2n}{1+n}\right)\mathbf{f} + \mathbf{g}\right]k' - A_1A_2k^2\right)\mathrm{d}\eta = 0$$
(29)

$$\int_{\eta_e}^{\eta_{e+1}} w_5 \left(\left[(1+\epsilon\theta) \,\theta'' + \epsilon\theta \right] + \mathbf{R}AA_4\theta'' + A_3A_4 \left[\left(\frac{2n}{1+n} \right) \mathbf{f} + \mathbf{g} \right] \theta' - A_3A_4h\theta \right) \mathrm{d}\eta = 0 \tag{30}$$

$$\int_{\eta_e}^{\eta_{e+1}} w_6 \left(S'' - ScCrS - ScSh + Sc \left[\left(\frac{2n}{1+n} \right) f + g \right] S' \right) d\eta = 0$$
(31)

here w_1, w_2, w_3, w_4, w_5 and w_6 represent arbitrary test functions and can be seen as variations in f, h, g, k, θ and S correspondingly.

3.3 Finite Element Formulation

Upon incorporating finite element approximations of the specified kind into the above-mentioned equations, the finite element model can be generated.

$$f = \sum_{j=1}^{2} f_j \psi_j, h = \sum_{j=1}^{2} h_j \psi_j, g = \sum_{j=1}^{2} g_j \psi_j, k = \sum_{j=1}^{2} k_j \psi_j, \theta = \sum_{j=1}^{2} \theta_j \psi_j, S = \sum_{j=1}^{2} S_j \psi_j.$$

with, $w_1 = w_2 = w_3 = w_4 = w_5 = w_6 = \psi_i, (i = 1, 2, 3, 4, 5, 6).$

where ψ_i are the shape functions for a typical element (η_e, η_{e+1}) and are indicated as

$$\psi_{1}^{e} = \frac{(\eta_{e+1} + \eta_{e} - 2\eta)(\eta_{e+1} - \eta)}{(\eta_{e+1} - \eta_{e})^{2}}, \quad \psi_{2}^{e} = \frac{4(\eta - \eta_{e})(\eta_{e+1} - \eta)}{(\eta_{e+1} - \eta_{e})^{2}},$$
$$\psi_{3}^{e} = \frac{(\eta_{e+1} + \eta_{e} - 2\eta)(\eta - \eta_{e})}{(\eta_{e+1} - \eta_{e})^{2}}, \quad \eta_{e} \le \eta \le \eta_{e+1}$$
(32)

The resulting finite element representation of the formulated equations is as follows:

$$\begin{bmatrix} K^{11} & [K^{12}] & [K^{13}] & [K^{14}] & [K^{15}] & [K^{16}] \\ [K^{21}] & [K^{22}] & [K^{23}] & [K^{24}] & [K^{25}] & [K^{26}] \\ [K^{31}] & [K^{32}] & [K^{33}] & [K^{34}] & [K^{35}] & [K^{36}] \\ [K^{41}] & [K^{42}] & [K^{43}] & [K^{44}] & [K^{45}] & [K^{46}] \\ [K^{51}] & [K^{52}] & [K^{53}] & [K^{54}] & [K^{55}] & [K^{56}] \\ [K^{61}] & [K^{62}] & [K^{63}] & [K^{64}] & [K^{65}] & [K^{66}] \end{bmatrix} \begin{bmatrix} f \\ h \\ g \\ k \\ \theta \\ S \end{bmatrix} = \begin{bmatrix} \{r^1 \\ r^2 \\ r^3 \\ r^4 \\ r^5 \\ r^6 \end{bmatrix}$$

where $[K^{nm}]$ and $[r^{m}]$ (m, n = 1, 2, 3, 4, 5, 6) are defined as

$$\begin{split} & \mathcal{K}_{\eta}^{(1)} = \int_{w}^{u_{q+1}} \psi_{i} \frac{\partial \psi_{j}}{\partial \eta} d\eta, \mathcal{K}_{\eta}^{(2)} = -\int_{w}^{v_{q+1}} \psi_{i} \psi_{j} d\eta, \mathcal{K}_{\eta}^{(3)} = \mathcal{K}_{\eta}^{(3)} = \mathcal{K}_{\eta}^{(3)} = \mathcal{K}_{\eta}^{(3)} = \mathcal{K}_{\eta}^{(3)} = \mathcal{K}_{\eta}^{(3)} = 0 \\ & \mathcal{K}_{\eta}^{(2)} = \Pr B \int_{w}^{u_{q+1}} (1 + \epsilon \psi_{i}) \frac{\partial \psi_{i}}{\partial \eta} \frac{\partial \psi_{j}}{\partial \eta} d\eta + \Pr B \epsilon \overline{\theta_{i}} \int_{w}^{u_{q+1}} \psi_{i} \psi_{i} \frac{\partial \psi_{j}}{\partial \eta} d\eta + \Pr B \epsilon \overline{\theta_{i}} \int_{w}^{u_{q+1}} \psi_{i} \psi_{j} \frac{\partial \psi_{j}}{\partial \eta} d\eta + \Pr B \epsilon \overline{\theta_{i}} \int_{w}^{u_{q+1}} \psi_{i} \psi_{j} \frac{\partial \psi_{j}}{\partial \eta} d\eta + \mathcal{A}_{1} A_{2} B \int_{w}^{u_{q+1}} \psi_{i} \psi_{j} \frac{\partial \psi_{j}}{\partial \eta} d\eta + \mathcal{A}_{1} A_{2} \left(\frac{2n}{1+n} \right) \overline{f_{i}} \int_{w}^{u_{q+1}} \psi_{i} \psi_{i} \frac{\partial \psi_{j}}{\partial \eta} d\eta + \mathcal{A}_{1} A_{2} \left(\frac{2n}{1+n} \right) \overline{f_{2}} \int_{w}^{u_{q+1}} \psi_{i} \psi_{j} \frac{\partial \psi_{j}}{\partial \eta} d\eta + \mathcal{A}_{1} A_{2} \left(\frac{2n}{1+n} \right) \overline{f_{2}} \int_{w}^{u_{q+1}} \psi_{i} \psi_{j} \frac{\partial \psi_{j}}{\partial \eta} d\eta + \mathcal{A}_{1} A_{2} \overline{g_{2}} \int_{w}^{u_{q+1}} \psi_{i} \psi_{j} \frac{\partial \psi_{j}}{\partial \eta} d\eta + \mathcal{A}_{1} A_{2} \int_{w}^{u_{q+1}} \psi_{i} \psi_{j} \frac{\partial \psi_{j}}{\partial \eta} d\eta \\ + \mathcal{A}_{1} A_{2} \overline{g_{1}} \int_{w}^{u_{q+1}} \psi_{i} \psi_{j} \frac{\partial \psi_{j}}{\partial \eta} d\eta + \mathcal{A}_{1} A_{2} \overline{g_{2}} \int_{w}^{u_{q+1}} \psi_{i} \psi_{j} \frac{\partial \psi_{j}}{\partial \eta} d\eta + \mathcal{A}_{1} A_{2} \int_{w}^{u_{q+1}} \psi_{i} \psi_{j} \frac{\partial \psi_{j}}{\partial \eta} d\eta \\ \mathcal{K}_{i}^{(3)} = 0, \mathcal{K}_{i}^{(3)} = 0, \mathcal{K}_{i}^{(3)} = 0, \mathcal{K}_{i}^{(3)} = 0, \mathcal{K}_{i}^{(3)} = 0 \\ \mathcal{K}_{i}^{(3)} = 0, \mathcal{K}_{i}^{(3)} = 0, \mathcal{K}_{i}^{(3)} = 0, \mathcal{K}_{i}^{(3)} = 0, \mathcal{K}_{i}^{(3)} = 0 \\ \mathcal{K}_{i}^{(4)} = 0, \mathcal{K}_{i}^{(4)} = 0, \mathcal{K}_{i}^{(3)} = 0, \mathcal{K}_{i}^{(3)} = 0, \mathcal{K}_{i}^{(3)} = 0 \\ \mathcal{K}_{i}^{(4)} = 0, \mathcal{K}_{i}^{(4)} = 0, \mathcal{K}_{i}^{(4)} = 0, \mathcal{K}_{i}^{(5)} = 0, \mathcal{K}_{i}^{(5)} = 0 \\ \mathcal{K}_{i}^{(4)} = 0, \mathcal{K}_{i}^{(4)} = 0, \mathcal{K}_{i}^{(5)} = 0, \mathcal{K}_{i}^{(5)} = 0, \mathcal{K}_{i}^{(5)} = 0, \mathcal{K}_{i}^{(5)} = 0 \\ \mathcal{K}_{i}^{(4)} = 0, \mathcal{K}_{i}^{(4)} = 0, \mathcal{K}_{i}^{(5)} = 0 \\ \mathcal{K}_{i}^{(5)} = 0, \mathcal{K}_{i}^{(5)} = 0, \mathcal{K}_{i}^{(5)} = 0, \mathcal{K}_{i}^{(5)} = 0, \mathcal{K}_{i}^{(5)} = 0 \\ \mathcal{K}_{i}^{(5)} = \int_{w}^{u_{i+1}} \psi_{i} \psi_{i} \frac{\partial \psi_{i}}{\partial \eta} d\eta + \mathcal{L}_$$

 $K_{ij}^{56} = 0, K_{ij}^{61} = 0, K_{ij}^{62} = 0, K_{ij}^{63} = 0, K_{ij}^{64} = 0,$

$$\begin{split} K_{ij}^{66} &= \int_{\eta_e}^{\eta_{e+1}} \frac{\partial \psi_i}{\partial \eta} \frac{\partial \psi_j}{\partial \eta} d\eta - ScCr \int_{\eta_e}^{\eta_{e+1}} \psi_i \psi_j d\eta + Sc \left(\frac{2n}{1+n}\right) \overline{f_1} \int_{\eta_e}^{\eta_{e+1}} \psi_i \psi_1 \frac{\partial \psi_j}{\partial \eta} d\eta \\ &+ Sc \left(\frac{2n}{1+n}\right) \overline{f_2} \int_{\eta_e}^{\eta_{e+1}} \psi_i \psi_2 \frac{\partial \psi_j}{\partial \eta} d\eta + Sc \overline{g_1} \int_{\eta_e}^{\eta_{e+1}} \psi_i \psi_1 \frac{\partial \psi_j}{\partial \eta} d\eta + Sc \overline{g_2} \int_{\eta_e}^{\eta_{e+1}} \psi_i \psi_2 \frac{\partial \psi_j}{\partial \eta} d\eta \\ &- Sc \overline{h_1} \int_{\eta_e}^{\eta_{e+1}} \psi_i \psi_1 \psi_j d\eta - Sc \overline{h_2} \int_{\eta_e}^{\eta_{e+1}} \psi_i \psi_2 \psi_j d\eta = 0 \\ r_i^2 &= 0, \ r_i^2 &= -\left(\psi_i \frac{d\psi_i}{d\eta}\right)_{\eta_e}^{\eta_{e+1}}, \ r_i^3 &= -\left(\psi_i \frac{d\psi_i}{d\eta}\right)_{\eta_e}^{\eta_{e+1}}, \ r_i^4 &= -\left(\psi_i \frac{d\psi_i}{d\eta}\right)_{\eta_e}^{\eta_{e+1}}, \ r_i^5 &= -\left(\psi_i \frac{d\psi_i}{d\eta}\right)_{\eta_e}^{\eta_{e+1}}, \end{split}$$

4 Results and Discussions

The comparison of sisko Al_2O_3 -Eg and TiO₂-Eg numerical resolutions of nanoliquids flow over a stretching sheet in the existence of MHD, were exemplified for various values of the related parameters and results were exposed graphically as well as in tabular form. The influence of the volume fraction parameter (ϕ) on the thickness of the hydrodynamic boundary layer, thermal boundary layer, and concentration boundary layer is depicted in the study of both Sisko Al_2O_3 -Eg and TiO₂-Eg nanoliquids using Figs. 3–5. It is identified that a rise in the value of (ϕ) tends to a fall in the radial velocity and temperature profiles. This reduction is more pronounced in the case of TiO₂-Eg nanofluid compared to Al_2O_3 -Eg nanofluid. However, the concentration profiles exhibit an inverted behavior.

The impact of the magnetic parameter (M) on the radial velocity, temperature, and concentration profiles of Sisko Al_2O_3 -Eg and TiO₂-Eg nanoliquids is illustrated in Figs. 6–8. The radial velocity profile exhibits a rise with higher values of (M) for both nanoliquids, with TiO₂-Eg nanofluid showing a more significant enhancement compared to Al_2O_3 -Eg nanofluid. On the other hand, the temperature profiles deteriorate for both Al_2O_3 -Eg and TiO₂-Eg nanofluid sas (M) increases. This reduction is more pronounced in TiO₂-Eg nanofluid compared to Al_2O_3 -Eg nanofluid. A similar decline is observed in the concentration profiles, with Al_2O_3 -Eg nanofluid exhibiting a higher decrease than TiO₂-Eg nanofluid.



Figure 3: Illustration of f' against ϕ



Figure 4: Representation of θ against ϕ



Figure 5: Variation of S against ϕ



Figure 6: Illustration of f' against M



Figure 7: Presentation of θ against M



Figure 8: Variation of S against M

The effects of the Siskofluid parameter (B) on the radial velocity, temperature, and concentration profiles of Sisko Al_2O_3 -Eg and TiO₂-Eg nanoliquids are depicted in Figs. 9–11. In both nanoliquids, an increase in (B) leads to an increase in the radial velocity profiles. The temperature profiles, on the other hand, decrease with higher values of (B) for both Al_2O_3 -Eg and TiO₂-Eg nanoliquids. This phenomenon is more pronounced in TiO₂-Eg-based nanofluid. Conversely, the concentration profiles increase with escalating values of (B). This accumulation trend is more significant in Al_2O_3 -Eg nanofluid compared to TiO₂-Eg-based nanofluid.



Figure 9: Variation of *f* ' against B



Figure 10: Variation of θ against **B**



Figure 11: Variation of S against B

Figs. 12 and 13 illustrate the effects of the Prandtl number (Pr) on the enhancements of radial velocity and temperature profiles in Sisko Al_2O_3 -Eg and TiO₂-Eg based nanoliquids. The radial velocity exhibits an increase with increasing values of Pr, and it is observed that this increase is more pronounced in TiO₂-Eg nanofluid compared to Al_2O_3 -Eg nanofluid. The same trend is observed in the temperature profiles as well.



Figure 12: Variation of *f* ' against Pr



Figure 13: Illustration of θ against Pr

From the observations in Figs. 14 and 15, it is evident that the profiles of radial velocity and temperature intensify in both Sisko Al_2O_3 -Eg and TiO₂-Eg based nanoliquids as the Schmidt number (Sc) increases. Moreover, this enhancement is more significant in TiO₂-Eg nanofluid compared to Al_2O_3 -Eg nanofluid. As for the chemical reaction parameter (Cr). Figs. 16 and 17 showcase the influence of the chemical reaction parameter (Cr) on the radial velocity and concentration distributions in Sisko Al_2O_3 -Eg and TiO₂-Eg based nanoliquids. It is observed that the reduction in both nanoliquids is slightly more pronounced in TiO₂-Eg nanofluid compared to Al_2O_3 -Eg nanofluid in terms of the concentration distributions.



Figure 14: Variation of *f*['] against Sc



Figure 15: Variation of θ against Sc



Figure 16: Variation of f' against Cr



Figure 17: Illustration of S against Cr

Figs. 18 and 19 depict the variations in radial velocity and temperature profiles for different values of the Reynolds number (R) in both Sisko Al_2O_3 -Eg and TiO₂-Eg based nanoliquids. It is observed that as the values of R increase, there is a decrease in the radial distributions for both



nanoliquids. Furthermore, this decrease is more pronounced in TiO₂-Eg nanofluid compared to Al₂O₃-Eg nanofluid.

Figure 18: Variation of f' against R



Figure 19: Variation of θ against *R*

In Fig. 19, it is observed that the temperature variations are significant in both nanoliquids, with TiO₂-Eg nanofluid exhibiting higher scatterings compared to Al_2O_3 -Eg nanofluid. Figs. 20–22 present the radial velocity, temperature, and concentration variations for different values of the nanoparticle volume fraction (Nv) in Sisko Al_2O_3 -Eg and TiO₂-Eg based nanoliquids. As the values of Nv increase, the radial velocity intensifies in both nanoliquids. This intensification is more pronounced in TiO₂-Eg based nanofluid compared to Al_2O_3 -Eg based nanofluid. The temperature variations also increase with higher values of Nv, and this increment is more significant in TiO₂-Eg based nanofluid compared to Al_2O_3 -Eg based nanofluid. Additionally, there is a reversed pattern in the concentration distributions, where Al_2O_3 -Eg based nanofluid shows more significant changes compared to TiO₂-Eg based nanofluid.







Figure 21: Variation of θ against η



Figure 22: Variation of S against η

The radial velocity distributions for different values of Nc are examined in Fig. 23 for both nano liquids, revealing an observed dropout phenomenon. Interestingly, it is observed that the dropout is more pronounced in the TiO_2 -Eg based nanofluid compared to the Al_2O_3 -Eg based nanofluid.



Figure 23: Variation of θ against η

Figs. 24–26 present the disparities in radial velocity, temperature, and concentration patterns for different suction parameter values (V_0) in both Sisko nanoliquids. The radial velocity distributions of Sisko Al_2O_3 -Eg and TiO_2 -Eg based nanoliquids decrease as the values of (V_0) increase. Notably, this trend is more pronounced in the TiO_2 -Eg based nanofluid compared to the Al_2O_3 -Eg-based nanofluid. Conversely, the temperature and concentration distributions diminish as the values of (V_0) increase. The reason behind this behavior is that the extraction of heated fluid from the fluid region effectively reduces the thickness of the thermal and concentration boundary layers.



Figure 24: Variation of f' against V_0



Figure 25: Variation of θ against V_{o}



Figure 26: Variation of S against V_o

The influence of the viscosity ratio parameter (α) on radial velocity patterns and concentration amplifications is examined for both Sisko Al_2O_3 -Eg and TiO_2 -Eg based nanoliquids. As the values of (α) increase, the radial velocity decreases in both Sisko Al_2O_3 -Eg and TiO_2 -Eg based nanoliquids. This effect is more pronounced in the TiO_2 -Eg-based nanofluid compared to the Al_2O_3 -Eg-based nanofluid. However, with increasing values of (α), the concentration profiles of the Sisko Al_2O_3 -Eg and TiO_2 -Eg based nanoliquids exhibit an increasing trend. This increasing behavior is more prominent in the Sisko Al_2O_3 -Eg nanoliquid compared to the TiO_2 -Eg nanoliquid. These findings are depicted in Figs. 27 and 28.



Figure 27: Variation of f' against α



Figure 28: Variation of S against α

Fig. 29 demonstrates that the radial velocity of Sisko Al_2O_3 -Eg and TiO_2 -Eg-based nanoliquids increases as the slip parameter (λ) values escalate. This behavior is less pronounced in the radial velocity of Al_2O_3 -Eg nanoliquids and more prominent in TiO_2 -Eg nanofluid. Regarding temperature and concentration distributions, as depicted in Figs. 30 and 31, both exhibit a decreasing trend. This decrease is more significant in the temperature distributions of TiO_2 -Eg based nanofluid compared to Al_2O_3 -Eg based nanofluid. Conversely, the decrease in concentration distributions is more noticeable in Sisko Al_2O_3 -Eg nanoliquids than TiO_2 -Eg nanoliquids.



Figure 29: Variation of f' against λ



Figure 30: Change of θ against λ



Figure 31: Presentation of S against λ

The impact of the temperature slip parameter (β) on temperature distributions is illustrated in Fig. 32, revealing a decrease in fluid temperature as the values of (β) increase. This temperature reduction is more pronounced in TiO_2 -Eg nanofluid compared to Al_2O_3 -Eg nanoliquids. Furthermore, Fig. 33 displays concentration profiles, demonstrating a decrease in values for both Sisko Al_2O_3 -Eg and TiO_2 -Eg-based nanoliquids as the concentration slip parameter (ξ) values increase. This decrease is more prominent in Al_2O_3 -Eg nanoliquids than TiO_2 -Eg-based nanoliquids.



Figure 32: Presentation of θ against β



Figure 33: Variation of S against ξ

Additionally, discussions and graphs have been developed for all non-dimensional parameters found in the physical phenomena of Sisko Al₂ O₃-Eg and TiO₂-Eg-based nanoliquids. Efforts were made to present exclusively novel findings obtained from Mathematica 10.4. The non-dimensional parameters include volume fraction (ϕ), magnitude, V0, Nc, Nv, Sc, and others

The statistical data pertaining to the Skin friction, local Nusselt number, and local Sherwood number for both Sisko Al_2O_3 -Eg and TiO₂-Eg nanoliquids has been calculated and presented in Tables 2 and 3. Analysis of Table 2 reveals a consistent decrease in the numerical values of the skin-friction coefficient, Nusselt number, and Sherwood number for both Sisko Al_2O_3 -Eg and TiO₂-Eg cases as the parameter () increases. The impact of (M) on the dimensionless rates of velocity shows an increase along the x-direction and an increase along the y-direction, while rates of temperature and mass exhibit a slight depreciation, as indicated in Table 2. Moreover, as the values of (Pr) increase, there is a notable decline in the skin-friction coefficient, Nusselt number, and Sherwood number for both Sisko Al_2O_3 -Eg and TiO₂-Eg nanoliquids.

Table 2: The influence of different parameters in the problem on skin-friction coefficient (-f''(0)), local Nusselt number $(-\theta'(0))$ and local Sherwood number $(-\phi'(0))$ for fixed $\phi = 0.01$, M = 0.5, Pr = 28.0, B = 1.0

Parameters				-f''(0)		heta'(0)		φ' (0)	
φ	М	Pr	В	Al ₂ O ₃ -Eg	TiO ₂ -Eg	Al ₂ O ₃ -Eg	TiO ₂ -Eg	Al ₂ O ₃ -Eg	TiO ₂ -Eg
0.01	0.5	2.58	1.0	0.64368	0.64028	0.29778	0.35412	0.16719	0.16971
0.05	0.5	2.58	1.0	0.46756	0.45167	0.30472	0.36588	0.16598	0.16755
0.1	0.5	2.58	1.0	0.25719	0.26368	0.31245	0.39950	0.16406	0.16330
0.01	0.5	2.58	1.0	1.46218	1.46272	0.30337	0.36986	0.71393	0.70122
0.01	0.8	2.58	1.0	2.85498	2.83536	0.31474	0.40385	0.76536	0.74523
0.01	1.2	2.58	1.0	4.52842	4.52557	0.32530	0.43330	0.80889	0.79910
0.01	0.5	2.58	1.0	1.86653	1.87193	0.30489	0.37490	0.72655	0.73772
0.01	0.5	2.58	1.0	0.58383	0.58117	0.29516	0.34331	0.67283	0.68312
0.01	0.5	2.58	1.0	0.45100	0.44855	0.28248	0.33443	0.66387	0.65220
0.01	0.5	2.58	1.0	2.92436	2.92545	0.30110	0.35421	0.73391	0.72020
0.01	0.5	2.58	2.0	1.92671	1.92509	0.28933	0.37777	0.75752	0.74411
0.01	0.5	2.58	3.0	1.51711	1.51535	0.27248	0.39101	0.79889	0.78743

Table 3: The effect of different parameters in the problem on skin-friction coefficient (-f''(0)), local Nusselt number $(-\theta'(0))$ and local Sherwood number $(-\phi'(0))$ for fixed, R = 0.5, Cr = 0.1, $N_v = 0.2$, $N_c = 0.2$

Parameters				-f''(0)		heta'(0)		$\phi'(0)$	
R	Cr	Nv	Nc	Al_2O_3 -Eg	TiO ₂ -Eg	Al_2O_3-Eg	TiO ₂ -Eg	Al ₂ O ₃ -Eg	TiO ₂ -Eg
0.5	0.1	0.2	0.2	1.46218	1.46269	0.32296	0.33458	0.73939	0.71995
0.8	0.1	0.2	0.2	1.44124	1.42310	0.31209	0.32146	0.77688	0.77088
1.2	0.1	0.2	0.2	1.42462	1.42237	0.30654	0.30981	0.84537	0.84404

(Continued)

Tabl	Fable 3 (continued)										
Parameters				-f''(0)		$\theta'(0)$		$\phi'(0)$			
R	Cr	Nv	Nc	Al ₂ O ₃ -Eg	TiO ₂ -Eg	Al_2O_3 -Eg	TiO ₂ -Eg	Al ₂ O ₃ -Eg	TiO ₂ -Eg		
0.5	0.1	0.2	0.2	1.49102	1.48272	0.30337	0.31452	0.71379	0.71365		
0.5	0.3	0.2	0.2	1.45613	1.46892	0.29899	0.29989	0.71119	0.71120		
0.5	0.5	0.2	0.2	1.43976	1.44120	0.29576	0.28942	0.71002	0.71011		
0.5	0.1	0.2	0.2	1.26281	1.19452	0.38793	0.38742	0.68726	0.68712		
0.5	0.1	0.5	0.2	0.97692	0.89845	0.46138	0.46157	0.67337	0.67312		
0.5	0.1	0.7	0.2	0.79628	0.72164	0.60145	0.60178	0.65122	0.61034		
0.5	0.1	0.2	0.2	1.21822	1.20012	0.36899	0.36231	0.36232	0.36211		
0.5	0.1	0.2	0.4	1.24420	1.21962	0.35595	0.35487	0.35475	0.35462		
0.5	0.1	0.2	0.6	1.28389	1.23491	0.34548	0.33212	0.33200	0.33109		

Furthermore, in Table 2, it is evident that the values of the skin-friction coefficient decrease with an increase in (B) for both Sisko Al_2O_3 -Eg and TiO_2-Eg nanoliquids. The Nusselt number decreases for the Sisko Al_2O_3 -Eg nanofluid but increases for the TiO_2-Eg nanofluid. However, the Sherwood number shows an escalation for both Sisko Al_2O_3 -Eg and TiO_2-Eg nanoliquids.

Table 3 illustrates the dimensionless rates of velocity upsurges along the x-direction and their deterioration along the y-direction for both Sisko Al_2O_3 -Eg and TiO_2 -Eg nanoliquids. In these cases, the Nusselt number and skin-friction coefficient values decrease, while the Sherwood number values increase across various values of (R). Notably, there is a decreasing trend observed in the rates of velocity and mass transfer for higher values of (Nv), while the rate of heat transfer increases with (Nv) for both Sisko Al_2O_3 -Eg and TiO_2 -Eg nanoliquids. Furthermore, the skin-friction coefficient values rise with increasing values of (Nc) for both nanoliquids. Conversely, the Nusselt number and Sherwood number values decrease with increasing (Nc) for both cases of Sisko Al_2O_3 -Eg and TiO_2 -Eg nanoliquids. These findings align closely with previously published results [30–32].

5 Conclusions

The present study employs FEM to computationally solve nonlinear PDEs, transformed into ODEs using appropriate similarity transformations. Investigating variable thermal conductivity and viscosity effects on MHD boundary layer flow, HMT in Sisko Al_2O_3 -Eg and TiO₂-Eg nanoliquids over a stretching surface with slip boundary conditions, several key conclusions emerge. An increase in the volume fraction of nanoparticles (ϕ) corresponds to elevated non-dimensional heat transfer rates, suggesting an enhancement in heat transfer efficiency due to nanoparticle attributes. Additionally, higher Sisko parameter (B) values correlate with reduced temperature profiles within nanoliquids. Moreover, amplified magnitudes of the parameter characterizing variable thermal conductivity (Nc) induce temperature reductions in the fluid region, showcasing the influence of variable thermal conductivity on temperature distributions. Furthermore, augmenting the parameter for variable viscosity (Nv) leads to decreased velocities within Sisko nanoliquids. These findings significantly contribute to understanding nanoliquid behavior and potentially find applications in various engineering and industrial processes.

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