



# Local thermal non-equilibrium analysis of conjugate free convection within a porous enclosure occupied with Ag–MgO hybrid nanofluid

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## Abstract

Current investigation aims to analyze the conjugate free convection inside a porous square cavity occupied with Ag–MgO hybrid nanofluid using the local thermal non-equilibrium (LTNE) model. Hybrid nanofluids are a novel kind of enhanced working fluids, engineered with enhanced thermo-physical and chemical properties. Two solid walls located between the horizontal bounds in two sides of cavity play the role of a conductive interface between the hot and cold walls, and moreover, the top and bottom bounds have been insulated. The governing differential equations are obtained by Darcy model and then for better representation of the results, converted into a dimensionless form. The finite element method is utilized to solve the governing equations. To evaluate the correctness and accuracy of the results, comparisons have been performed between the outcomes of this work and the previously published results. The results indicate that using the hybrid nanoparticles decreases the flow strength and the heat transfer rate. The heat transfer rate augments when  $R_k$  rises and the flow strength augments when  $Ra$  grows. Enhancing the porosity increases strongly the size and strength of the vortex composed inside the porous medium. When  $K_r$  is low, the heat transfer rate is low and by increasing  $K_r$ , thermal fields become closer to each other. The effect of hybrid nanoparticles on thermal fields with the thinner solid walls is more than that the thicker ones. An increment in  $H$  eventuates the enhancement of heat transfer and hence, the thermal boundary layer thickness. By increasing the volume fraction of the hybrid nanoparticles,  $Nu_{hnf}$  and  $Nu_s$  decrease in constant  $Ra$ . Besides, increase in  $Ra$  enhances the  $Nu_{hnf}$  and  $Nu_s$ . For a certain  $d$ , the reduction of  $Nu_s$  due to using the hybrid nanoparticles is more than that for  $Nu_{hnf}$ . The increment of  $d$  lessens  $Nu_{hnf}$  for all values of  $K_r$  and has not specific trends for  $Nu_s$ . Utilizing hybrid nanoparticles decreases  $Nu_s$  (except  $d = 0.4$ ), rises  $Nu_s$  when  $K_r < 18$ , while it can increase  $Nu_s$  for  $K_r > 42$ . In constant  $d$ , increment of  $H$ , respectively, decreases and boosts  $Nu_{hnf}$  and  $Nu_s$ . For all values of  $d$ , increment of  $\varepsilon$  declines  $Nu_{hnf}$ . In low value of  $d$ , the increase in  $\varepsilon$  reduces  $Nu_s$ , whereas at higher values,  $Nu_s$  has continuously enhancing trend. For different values of  $d$ , the increase in  $\varepsilon$  scrimps  $Nu_{hnf}$ . The increment of  $d$  and also  $\varepsilon$ , and  $H$  are, respectively, decreases and increases the heat transfer rate.

**Keywords** Hybrid nanofluids · Ag–MgO/water · Porous medium · Conjugate natural convection · Local thermal non-equilibrium approach

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## Introduction

The conjugate heat exchange in porous media with applying the thermal non-equilibrium (TNE) models is extremely significant area in high-power heat transfer usages [1]. Many researchers have focused on conjugate free convective heat exchange in enclosures due to its significance for a lot of engineering and industrial systems, like heat exchangers, solar collector technology, cooling and storage of radioactive waste containers, flows in microchannels, material processing, electrical units, thermal exchange circuits, and the building energy components [1–3]. On the other hand, various engineering and practical usages of local thermal non-equilibrium (LTNE) such as fuel cells, heat pipes and sinks, dryers, computer chips, and catalytic reactors motivate a lot of authors to investigate this model with a goal of precise prediction in the thermal treatment of porous medium [4–7]. The design of using hybrid nanoparticles as nano-additives can be managed in such usages for multi-objective advantages containing heat transfer enhancement or inhibition. Accordingly, this investigation aims to theoretically analyze the influence of the presence of a hybrid nanofluid in cavity filled by porous media.

Nanofluids are new kind of emended (reinforced) fluids first presented by Choi [8] which contain well-dispersed solid nanometer-sized particles [2, 9–11]. Experiments proved that with addition of nanoparticles, physical characteristics of ordinary host fluid including density, viscosity, and thermal conductivity have increased [12, 13]. Thus, nanoparticles existence affects the heat transfer rate and is required in usages. Hybrid nanofluids are newfound category of nanofluids that consist of a small amount of metal nanoparticles (like Zn, Cu, Ag, and Al) and non-metallic nanoparticles (like  $\text{Al}_2\text{O}_3$ , CuO, MgO, and  $\text{Fe}_3\text{O}_4$ ). Metallic nanoparticles have high thermal conductivities, whereas they have limitations such as stability and reactivity. On the contrary, non-metallic nanoparticles give lower thermal conductivity compared to the metallic, while they have great number of beneficial properties like stability and chemical inertness. Hence, there is expectation that addition of metallic to non-metallic-based nanofluid able to raise the thermo-physical properties of the final mixture without lessening the stability [14, 15]. Newly, a lot of experimental investigations [16–19] and numerical studies [15, 20–22] have been carried out on hybrid nanofluid as a new conception in technology. Suresh et al. [14] examined the thermal conductivity and viscosity of the alumina–copper/water hybrid nanofluid. Esfe et al. [18, 19] measured and predicted the thermal conductivity of various hybrid nanofluids. Sarkar et al. [23] and Sundar et al. [24] published general reviews about the recent researches and

the future challenges in the area of hybrid nanofluids. Heat exchange in an annulus between two confocal elliptic cylinders filled by Cu– $\text{Al}_2\text{O}_3$ /water hybrid nanofluid conducted numerically by Tayebi and Chamkha [15]. A computational analysis of three various nanofluids ( $\text{Al}_2\text{O}_3$ ,  $\text{TiO}_2$  and  $\text{SiO}_2$ ) and their hybrids was performed by Minea [25]. Results indicated that thermal conductivity is rising by minimum 12% with adding the nanoparticles.

Convective heat transfer within enclosures occupied with nanofluid considering various conditions is a basic problem and conducted by a lot of authors as an attractive subject [16, 26–33]. Cavities are sometimes filled by porous environment that is saturated with a fluid or nanofluid. Kasaeian et al. [34] investigated recently the progresses in the area of nanofluid flow and heat transfer in a porous medium in format of a comprehensive review. Chamkha and Ismael [29] checked out the free convective heat transfer within a nanoliquid enclosure partially filled with a porous medium. Ghalambaz et al. [32] have considered the existence of viscous dissipation and radiation influences on free convective heat transfer within the porous nanofluid enclosure. Sheikholeslami et al. [35] investigated free convection heat exchange in a region having sinusoidal wall occupied by nanofluid affected by magnetic field. Free and mixed convective heat transfer in a 2-D nanofluid cavity considering multiple pairs of heat source–sinks with simulating the two-phase flow is performed by Garoosi et al. [36]. Most recently, investigation of utilizing a hybrid nanofluid in issues related to cavity has examined by some of the researchers [37–39]. Chamkha et al. [37] conducted the melting process of a nanoparticles-enhanced phase change material in a square cavity over a heated horizontal cylinder located in the middle of the enclosure under the influence of both single and hybrid nanoparticles.

Recently, a number of researchers have focused on different thermal boundary conditions in enclosure. One of the most challenging issues in this field is using the LTE (when the fluid and solid temperature are equal) [40] and LTNE (where the fluid and solid temperature are varied) [41, 42] as two well-known models for porous media. Many investigations have utilized the LTE model, whereas LTNE model has not attracted great consideration. Baytas and Pop [43], Kuznetsov and Nield [41], Agarwal and Bhadauria [42] examined an influence of LTNE approach on free convection in different case studies. Lately, Alsabery et al. [1, 44, 45] and Zargartalebi et al. [6] checked out an influence of the LTNE on natural convective heat transfer inside various enclosures filled by nanofluid. Other studies in this field can be found in the literature [46, 47].

Some of researchers focused on conjugate natural convections in enclosures [2, 3, 48–52] for its significant usages in building energy components, electrical parts

cooling, collectors of solar energy, material production, etc. Kimura et al. [53] have explained in a review paper that the fundamental hypotheses about the mathematical approaches of conjugate convection–conduction heat exchange process in porous medium. Influence of natural convective–conductive heat transfer in a cavity bounded by a solid vertical wall of different thermal conductivity ratio is investigated numerically by Kaminski and Prakash [50]. Chamkha and Ismael [2] checked out conjugate heat exchange in a porous cavity saturated by nanoliquids and heated with triangular-shaped thick wall. Mahmoudi et al. [3] investigated the conjugate heat exchange in a cavity with solid walls having Cu–water nanoliquid. It was demonstrated that the divider’s placement affects the heat transfer increment with Rayleigh numbers and solid particles concentration. Sheremet and Pop [48] perused the conjugate free convection in porous nanoliquid cavity by the finite difference method. Free convective–conductive heat exchange in a nanoliquid enclosure having solid partition on various walls is evaluated by Selimefendigil and Öztop [51]. Good references on convective nanofluid flow and heat transfer were presented in the books by Das et al. [54], Nield and Bejan [55], and Shenoy et al. [56] and in the review papers by Buongiorno [57], Kakaç and Pramuanjaroenkij [12], Wong and Leon [58], Wen et al. [59], Mahian et al. [60], and many others.

Due to the presented literature survey and by our best of knowledge, there are no investigations which analyzed natural convective–conductive heat exchange within a porous enclosure saturated with Ag–MgO hybrid nanoliquid with LTNE approach. We believe that this work is noteworthy, and its usefulness for different engineering systems as discussed above motivated us. In hybrid nanofluid usage, it is the first effort to evaluate the conduction–convection effects of natural heat transfer considering the LTNE model within a cavity. The current investigation studies the effects of the Darcy–Rayleigh number, nanoparticles concentration, porosity, interface parameter, modified thermal conductivity ratio, and the width of the solid wall on the nanoliquid flow and thermal fields. The governing differential equations are written using the Darcy model and then transformed into a non-dimensional form. Solution of the boundary-value problem has been performed using the finite element method. Several examinations are performed to find out the accuracy of the present model. The numerical data are validated using the results available in the literature.

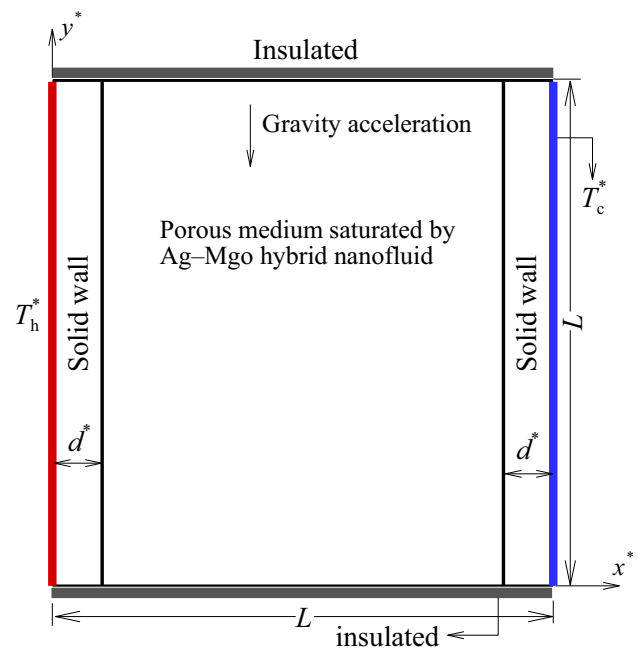


Fig. 1 A simple schematic of the problem

## Description and formulation of the model

A simple schematic view of the under study conjugate free convection problem is observed in Fig. 1. Two solid walls which have the finite thickness of  $d^*$  are located between the horizontal bounds in two sides of a square cavity with size  $L$ . Indeed, these solid walls play the role of a conductive interface between the hot and cold walls with constant temperatures of  $T_h$  and  $T_c$ , respectively, and the porous entity occupied by Ag–MgO hybrid nanofluid. The top and bottom bounds have been thermally insulated. It should be mentioned that all of the walls are fully impermeable. Here, the volumetric heat exchange between the hybrid nanofluid and porous matrix is finite and nonzero. This specific type of the heat transfer is considered using the LTNE model. The hybrid nanoparticles always remain suspended in the pores of the cavity. There is no contact resistance in the interface boundary between the solid walls and porous medium. In addition, the dynamic and thermal slips between nanoparticles and host liquid are negligible. Only body force applied is gravity force that acts in reverse direction of  $y$ -axis. During the process of the natural convection, all the characteristics of the host liquid and nanoparticles are unchangeable except for the density in the buoyancy term in momentum equation where its variation will be modeled using Boussinesq approximation.

Using the aforementioned assumptions to derive the equations leads to the following formulations

$$\frac{\partial u_{\text{hnf}}^*}{\partial x^*} + \frac{\partial v_{\text{hnf}}^*}{\partial y^*} = 0, \tag{1}$$

$$0 = -\frac{\partial p^*}{\partial x^*} - \frac{\mu_{\text{hnf}}}{K} u_{\text{hnf}}^*, \tag{2}$$

$$0 = -\frac{\partial p^*}{\partial y^*} - \frac{\mu_{\text{hnf}}}{K} v_{\text{hnf}}^* - (\rho\beta)_{\text{hnf}}(T_{\text{hnf}}^* - T_0)\mathbf{g}_y, \tag{3}$$

$$\begin{aligned} &(\rho C_p)_{\text{hnf}} \left( u_{\text{hnf}}^* \frac{\partial T_{\text{hnf}}^*}{\partial x^*} + v_{\text{hnf}}^* \frac{\partial T_{\text{hnf}}^*}{\partial y^*} \right) \\ &= \varepsilon k_{\text{hnf}} \left( \frac{\partial^2 T_{\text{hnf}}^*}{\partial x^{*2}} + \frac{\partial^2 T_{\text{hnf}}^*}{\partial y^{*2}} \right) + h(T_s^* - T_{\text{hnf}}^*), \end{aligned} \tag{4}$$

$$0 = (1 - \varepsilon)k_s \left( \frac{\partial^2 T_s^*}{\partial x^{*2}} + \frac{\partial^2 T_s^*}{\partial y^{*2}} \right) + h(T_{\text{hnf}}^* - T_s^*). \tag{5}$$

The energy equation without considering convection term and heat absorption or generation for the solid wall will be summarized as follows

$$0 = \frac{\partial^2 T_w^*}{\partial x^{*2}} + \frac{\partial^2 T_w^*}{\partial y^{*2}}. \tag{6}$$

The boundary conditions subjected in dimensional  $x^*$  and  $y^*$  coordinates are

$$\begin{aligned} T_w^* &= T_h^* \quad \text{on} \quad x^* = 0, 0 \leq y^* \leq L, \\ T_w^* &= T_c^* \quad \text{on} \quad x^* = L, 0 \leq y^* \leq L, \end{aligned} \tag{7-a}$$

$$\begin{aligned} u_{\text{hnf}}^* &= v_{\text{hnf}}^* = 0, \quad \frac{\partial T_s^*}{\partial y^*} = \frac{\partial T_{\text{hnf}}^*}{\partial y^*} = 0 \quad \text{on} \\ &y^* = 0, d^* \leq x^* \leq L - d^*, \end{aligned} \tag{7-b}$$

$$\begin{aligned} u_{\text{hnf}}^* &= v_{\text{hnf}}^* = 0, \quad \frac{\partial T_s^*}{\partial y^*} = \frac{\partial T_{\text{hnf}}^*}{\partial y^*} = 0 \quad \text{on} \\ &y^* = L, d^* \leq x^* \leq L - d^*, \end{aligned}$$

$$\begin{aligned} \frac{\partial T_w^*}{\partial y^*} &= 0 \quad \text{on} \\ &0 \leq x^* \leq d^*, L - d^* \leq x^* \leq L, y^* = 0, y^* = L. \end{aligned} \tag{7-c}$$

At the solid wall–porous medium interface, we have the following conditions:

$$\left. \begin{aligned} &u_{\text{hnf}}^* = v_{\text{hnf}}^* = 0, T_{\text{hnf}}^* = T_s^* = T_w^* \\ &k_w \frac{\partial T_w^*}{\partial x^*} = \varepsilon k_{\text{hnf}} \frac{\partial T_{\text{hnf}}^*}{\partial x^*} + (1 - \varepsilon)k_s \frac{\partial T_s^*}{\partial x^*} \end{aligned} \right\} \text{on} \tag{7-d}$$

$$x^* = d^*, x^* = L - d^* \text{ and } 0 \leq y^* \leq L.$$

At the present time, the model which can accurately give the thermal conductivity and dynamic viscosity of the hybrid nanoliquids has not been provided. Hence, this study applies the experimental values of these two thermo-physical properties in [18] to do a realistic analysis. Esfe et al. [18] experimentally determined the thermal conductivity and viscosity of the MgO–Ag hybrid nanoliquid in which the diameters of Ag and MgO particles were 25 and 40 nm, respectively. The volume fraction of

each nanoparticle was evenly 50% of the total volume fraction. They represented the new relations to calculate the thermal conductivity (Eq. 8) and viscosity (Eq. 9) of the MgO–Ag hybrid nanoliquid by curve fitting on the empirical results. The empirical data of Table 1 are the values that the curve fitting have been performed based on them and are utilized to represent the results. Additionally, in Table 1,  $M$  and  $\alpha_r$  are two dimensionless functions of the volume fractions and will be introduced in the following.

There are excellent correspondences between empirical data and models proposed by Esfe et al. [18]. The other thermo-physical properties employed in this study (shown in Table 2) are obtained using the classical models that are mostly in good coincidence with the empirical results. Here  $\varphi_{\text{hnf}}$  is the total volume fraction which equals the summation of the volume fractions of Ag and MgO.

$$k_{\text{hnf}} = \left( \frac{0.1747 \times 10^5 + \varphi_{\text{hnf}}}{0.1747 \times 10^5 - 0.1498 \times 10^6 \varphi_{\text{hnf}} + 0.1117 \times 10^7 \varphi_{\text{hnf}}^2 + 0.1997 \times 10^8 \varphi_{\text{hnf}}^3} \right) k_{\text{bf}}, \tag{8}$$

$0 \leq \varphi_{\text{hnf}} \leq 0.03$

$$\begin{aligned} \mu_{\text{hnf}} &= \left( 1 + 32.795 \varphi_{\text{hnf}} - 7214 \varphi_{\text{hnf}}^2 + 714600 \varphi_{\text{hnf}}^3 \right. \\ &\quad \left. - 0.1941 \times 10^8 \varphi_{\text{hnf}}^4 \right) \mu_{\text{bf}}. \quad 0 \leq \varphi_{\text{hnf}} \leq 0.02 \end{aligned} \tag{9}$$

To eliminate the pressure terms from momentum equations, the cross differencing can be applied for the momentum equations along  $x$  and  $y$  directions:

$$\frac{\mu_{\text{hnf}}}{K} \left( \frac{\partial u_{\text{hnf}}^*}{\partial y^*} - \frac{\partial v_{\text{hnf}}^*}{\partial x^*} \right) = -g(\rho\beta)_{\text{hnf}} \frac{\partial T_{\text{hnf}}^*}{\partial x^*}, \tag{10}$$

Now, a function should be defined that corresponds to the continuity equation and the velocity components  $u_{\text{hnf}}^*$  and  $v_{\text{hnf}}^*$  can be obtained by partial differentials of that function. In fact, this function is same as the stream function that is written as:

$$u_{\text{hnf}}^* = \frac{\partial \psi^*}{\partial y}, \quad v_{\text{hnf}}^* = -\frac{\partial \psi^*}{\partial x}, \tag{11}$$

To have a dimensional parametric analysis, the non-dimensional parameters are employed to transfer the

**Table 1** Experimental data for thermal conductivity and viscosity of hybrid MgO–Ag/water nanoliquid. Reproduced with permission from [18]

$\varphi_{\text{hnf}}/\%$	$k_{\text{hnf}}/k_{\text{bf}}$	$\mu_{\text{hnf}}/\mu_{\text{bf}}$	$M$	$\alpha_r$
0	1	1	1	1
0.56	1.05376	1.063619	0.93789	1.0555
1.12	1.08296	1.160534	0.85746	1.0866
1.5	1.13208	1.300207	0.76407	1.1371
2	1.1573	1.3815	0.71753	1.1642

**Table 2** Applied relations for hybrid nanofluid properties. Reproduced with permission from [61]

Hybrid nanofluid properties	Applied relation
Density	$\rho_{\text{hnf}} = \rho_f(1 - \varphi_{\text{hnp}}) + \rho_{\text{Ag}}\varphi_{\text{Ag}} + \rho_{\text{MgO}}\varphi_{\text{MgO}}$
Buoyancy coefficient	$(\rho\beta)_{\text{hnf}} = (1 - \varphi_{\text{hnp}})(\rho\beta)_f + \varphi_{\text{Ag}}(\rho\beta)_{\text{Ag}} + \varphi_{\text{MgO}}(\rho\beta)_{\text{MgO}}$
Heat capacity	$(\rho c_p)_{\text{hnf}} = (1 - \varphi_{\text{hnp}})(\rho c_p)_f + \varphi_{\text{Ag}}(\rho c_p)_{\text{Ag}} + \varphi_{\text{MgO}}(\rho c_p)_{\text{MgO}}$
Thermal diffusivity	$\alpha_{\text{hnf}} = k_{\text{hnf}}/(\rho c_p)_{\text{hnf}}$

equations and boundary conditions from dimensional  $x^*$ - $y^*$  coordinates to dimensionless  $x$ - $y$  coordinates:

$$x = x^*/L, \quad y = y^*/L, \quad \psi = \psi^*/\alpha_{\text{bf}}, \quad T_{\text{hnf}} = (T_{\text{hnf}}^* - T_c^*)/(T_h^* - T_c^*),$$

$$T_s = (T_s^* - T_c^*)/(T_h^* - T_c^*), \quad H = \frac{hL^2}{\varepsilon k_{\text{bf}}}, \quad K_r = \frac{\varepsilon k_{\text{bf}}}{(1 - \varepsilon)k_s}. \tag{12}$$

So, the final non-dimensional equations can be rewritten:

$$\frac{\partial^2 \psi}{\partial x^2} + \frac{\partial^2 \psi}{\partial y^2} = -Ra \cdot M \cdot \frac{\partial T_{\text{hnf}}}{\partial x}, \tag{13}$$

$$\frac{1}{\varepsilon} \left( \frac{\partial \psi}{\partial y} \frac{\partial T_{\text{hnf}}}{\partial x} - \frac{\partial \psi}{\partial x} \frac{\partial T_{\text{hnf}}}{\partial y} \right) = \alpha_r \left( \frac{\partial^2 T_{\text{hnf}}}{\partial x^2} + \frac{\partial^2 T_{\text{hnf}}}{\partial y^2} \right) + \frac{(\rho C_p)_{\text{bf}}}{(\rho C_p)_{\text{hnf}}} H(T_s - T_{\text{hnf}}), \tag{14}$$

$$0 = \left( \frac{\partial^2 T_s}{\partial x^2} + \frac{\partial^2 T_s}{\partial y^2} \right) + K_r H(T_{\text{hnf}} - T_s), \tag{15}$$

$$0 = \left( \frac{\partial^2 T_w}{\partial x^2} + \frac{\partial^2 T_w}{\partial y^2} \right). \tag{16}$$

In Eq. (13),  $Ra$  is the abbreviation of the Darcy–Rayleigh number which is defined as

$$Ra = gK(\rho\beta)_{\text{bf}}(T_h - T_c)L/(\alpha_{\text{bf}}\mu_{\text{bf}}).$$

Two dimensionless numbers  $M$  and  $\alpha_r$  are also introduced as

$$M = ((\rho\beta)_{\text{hnf}}/(\rho\beta)_{\text{bf}})(\mu_{\text{bf}}/\mu_{\text{hnf}})$$

and  $\alpha_r = \alpha_{\text{hnf}}/\alpha_{\text{bf}}$ . The thermal and dynamic boundary conditions of the model have been listed below:

$$T_w = 1 \quad \text{on} \quad x = 0, 0 \leq y \leq 1,$$

$$T_w = 0 \quad \text{on} \quad x = 1, 0 \leq y \leq 1, \tag{17 - a)}$$

$$\psi = 0, \quad \frac{\partial T_s}{\partial y} = \frac{\partial T_{\text{hnf}}}{\partial y} = 0 \quad \text{on} \quad y = 0, d \leq x \leq 1 - d,$$

$$\psi = 0, \quad \frac{\partial T_s}{\partial y} = \frac{\partial T_{\text{hnf}}}{\partial y} = 0 \quad \text{on} \quad y = 1, d \leq x \leq 1 - d, \tag{17 - b)}$$

$$\frac{\partial T_w}{\partial y} = 0 \quad \text{on} \quad 0 \leq x \leq d, 1 - d \leq x \leq 1, y = 0, y = 1, \tag{17 - c)}$$

$$\psi = 0, T_{\text{hnf}} = T_s = T_w$$

$$\left. \begin{aligned} \frac{\partial T_{\text{hnf}}}{\partial x} = R_k \frac{k_{\text{bf}}}{k_{\text{hnf}}} \frac{\partial T_w}{\partial x} - K_r^{-1} \frac{k_{\text{bf}}}{k_{\text{hnf}}} \frac{\partial T_s}{\partial x} \end{aligned} \right\} \quad \text{on} \tag{17 - d)}$$

$$x = d, x = 1 - d \text{ and } 0 \leq y \leq 1.$$

Parameter appeared in the last boundary conditions,  $R_k$ , is named as the thermal conductivity ratio and introduced as:

$$R_k = \frac{k_w}{\varepsilon k_{\text{bf}}}, \tag{18}$$

To evaluate the heat transfer rate through the porous cavity, the average Nusselt number of the solid and liquid phases of the porous medium at the porous-wall interface bounds is defined as follows:

$$Nu_{\text{hnf}} = \int_0^1 \left( \frac{\partial T_{\text{hnf}}}{\partial x} \right)_{x=d} dy, \tag{19}$$

$$Nu_s = \int_0^1 \left( \frac{\partial T_s}{\partial x} \right)_{x=d} dy, \tag{20}$$

Further, the rate of heat exchange across the walls can be obtained using the weighted sum of the aforesaid average Nusselt numbers such as the following:

$$Q_w = R_k^{-1} Nu_{\text{hnf}} + R_k^{-1} K_r^{-1} Nu_s, \tag{21}$$

or

$$Q_{\text{wall}} = \frac{q_w L}{k_w} = \int_0^1 \left( \frac{\partial T_w}{\partial x} \right)_{x=0} dy, \tag{22}$$



### Numerical approach, grid sensitivity test and validation

The formulated Eqs. (12)–(15) are nonlinear and coupled; therefore, the finite element method can be more effective for the present study. The details of the used finite element method can be found in [62]. To have an accurate solution which is independent of the number of the elements, it needs to do the grid independency test. The grid independency examination is performed based on assessing the variations of  $Nu_{hnf}$ ,  $Nu_s$  and  $|\psi|_{max}$  with the grid size. From Table 3, the maximum error because of the variations of grid size, belonged to  $|\psi|_{max}$ , is 0.3% when grid size enhances from  $50 \times 50$  to  $100 \times 100$ . This means that a  $50 \times 50$  mesh can give very satisfactory outcomes. However, to receive the grid independence results at all cases, the grid of  $100 \times 100$  nodes is confidently employed to discretize the computational domain. The accuracy of the outcomes of the present work is evaluated via comparing these results and those reported by Sun and Pop [30], Sheremet et al. [28] and Saeid [63] in Tables 4 and 5. The excellent agreements found between the data of the present work and reported in the literature certify the accuracy of present modeling and simulating.

### Results and discussion

This part has focused on influences of parameters appeared in the equations and boundary conditions such as Darcy–Rayleigh number  $Ra = 10$ –1000, porosity  $\varepsilon = 0.1$ –0.9, interface parameter  $H = 1$ –1000,  $K_r = 0.1$ –10, volume fraction of the hybrid nanofluid  $\varphi_{hnf} = 0$ –0.02 and the width of the solid wall  $d = 0.1$ –0.4 on the nanoliquid flow and thermal fields as well as the rates of heat exchange through the solid walls, the solid and liquid phases of the porous medium.

The set of images shown in Fig. 2 illustrates the dependency of streamlines, isotherms of the liquid and solid phases of the porous medium on  $R_k$  at constant parameters  $Ra = 10^3$ ,  $\varepsilon = 0.5$ ,  $d = 0.1$ ,  $K_r = H=1$ . In the

**Table 3** Grid independency test for  $Ra = 10^3$ ,  $H = 100$ ,  $K_r = 10$ ,  $R_k = 1$ ,  $d = 0.1$ ,  $\varepsilon = 0.6$ ,  $\varphi_{Ag} = \varphi_{MgO} = 0.0$

Grid size	$Nu_{hnf}$	Error/%	$Nu_s$	Error/%	$ \psi _{max}$	Error/%
$50 \times 50$	12.662	–	2.689	–	12.518	–
$100 \times 100$	12.641	0.17	2.694	0.18	12.478	0.3
$150 \times 150$	12.637	0.03	2.695	0.03	12.471	0.05
$200 \times 200$	12.635	0.01	2.695	0.0	12.468	0.02

**Table 4** Average Nusselt number in a porous triangular cavity for different Rayleigh numbers and nanoparticles volume fraction

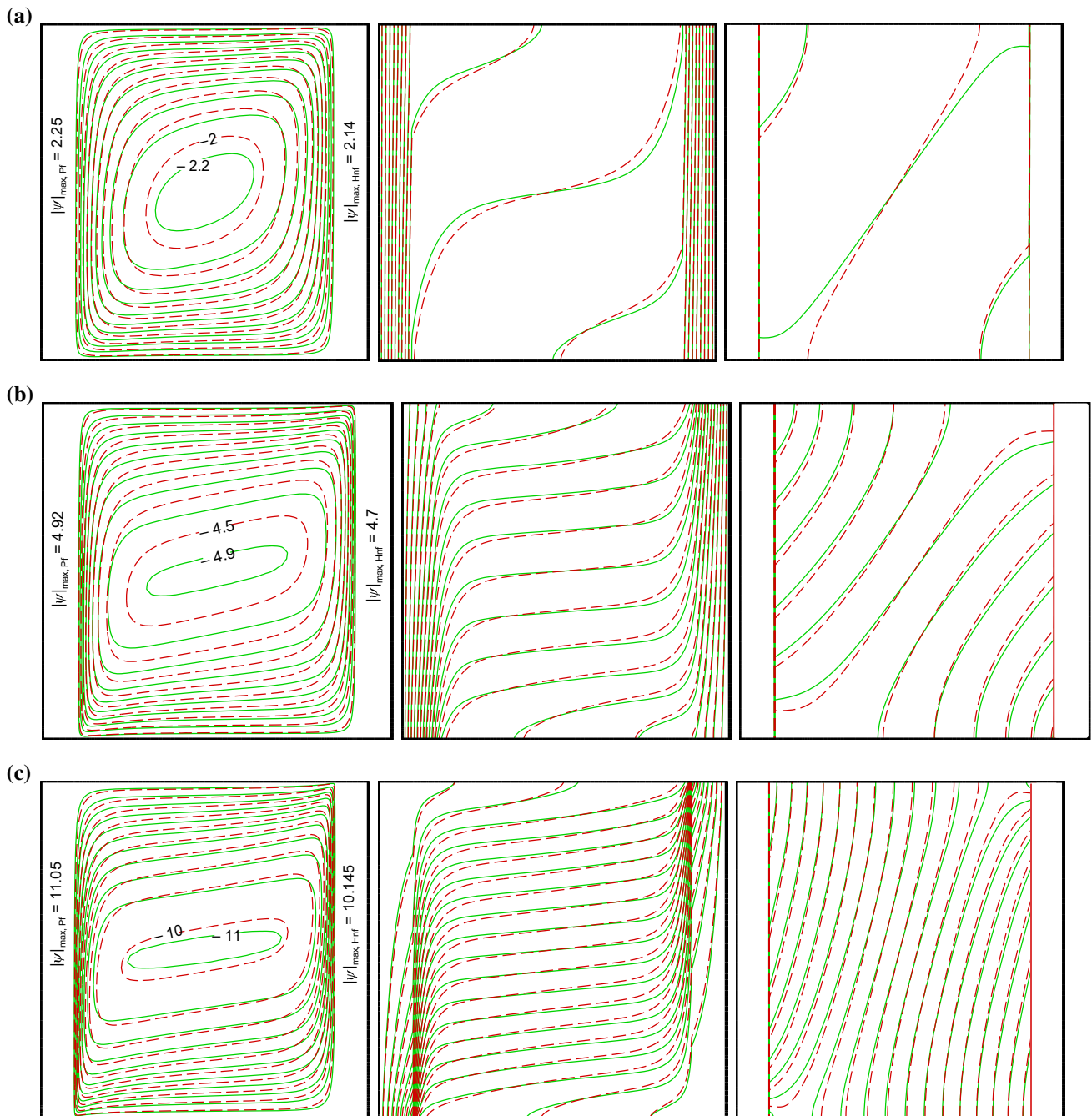
$Ra$	$\varphi$	Sun and Pop [30]	Sheremet et al. [28]	Present work
500	0	9.66	9.65	9.64
1000	0.1	9.42	9.41	9.42
500	0	13.9	14.05	13.96
1000	0.2	12.85	12.84	12.85

**Table 5** Data of the present numerical solution and those performed by Saeid [63] at  $Ra = 10^3$ ,  $d = 0.1$ ,  $K_r = H=1$

$ \psi _{max}$	$\bar{Q}_w$	$\bar{Nu}_s$	$\bar{Nu}_f$	$R_k$	
3.614	4.559	0.112	0.344	0.1	Present results
8.053	3.348	0.425	2.923	1	
16.409	1.105	1.007	10.04	10	
3.536	4.357	0.110	0.326	0.1	Saeid [63]
7.898	3.232	0.418	2.814	1	
16.219	1.090	1.010	9.887	10	

figures, the solid and dash lines are indicants of the pure fluids and Ag–MgO hybrid nanofluids, respectively. It is visible that the use of the Ag–MgO hybrid nanoparticles decreases the strength of the fluid flow. Indeed, using the Ag–MgO hybrid nanoparticles amplifies the dynamic viscosity of the base fluid which acts as a resistant force against the buoyancy effects. The increment of the density of the isotherms attributed to each of the phases by increasing  $R_k$  bodes that the heat transfer rate through the porous region raises when  $R_k$  increases. Comparing the isotherms of the fluid and solid phases at low and high values of  $R_k$  indicates that when  $R_k$  is low, the influences of the presence of the hybrid nanoparticles on the temperatures fields are more than those compared to the high values of  $R_k$ , while the relative decrease in the strength of the flow for the high value of  $R_k$  is more than that for the low value of  $R_k$ . These relative decreases are 4.8 and 8.0% for  $R_k = 0.1$  and 10, respectively.

From Fig. 3, it is apperceived that a weak circular-shaped vortex is appeared in the center of the porous cavity when  $Ra = 10$ . In this case, the parallelism of the isotherms attributed to the fluid phase of the porous medium with vertical bounds reflects the heat conduction dominance on the thermal convection mechanism during the heat transfer process. As  $Ra$  enhances, the strength and the size of the formed vortices becomes more as a result of increasing the buoyancy effects. In addition, it is visible that the hybrid



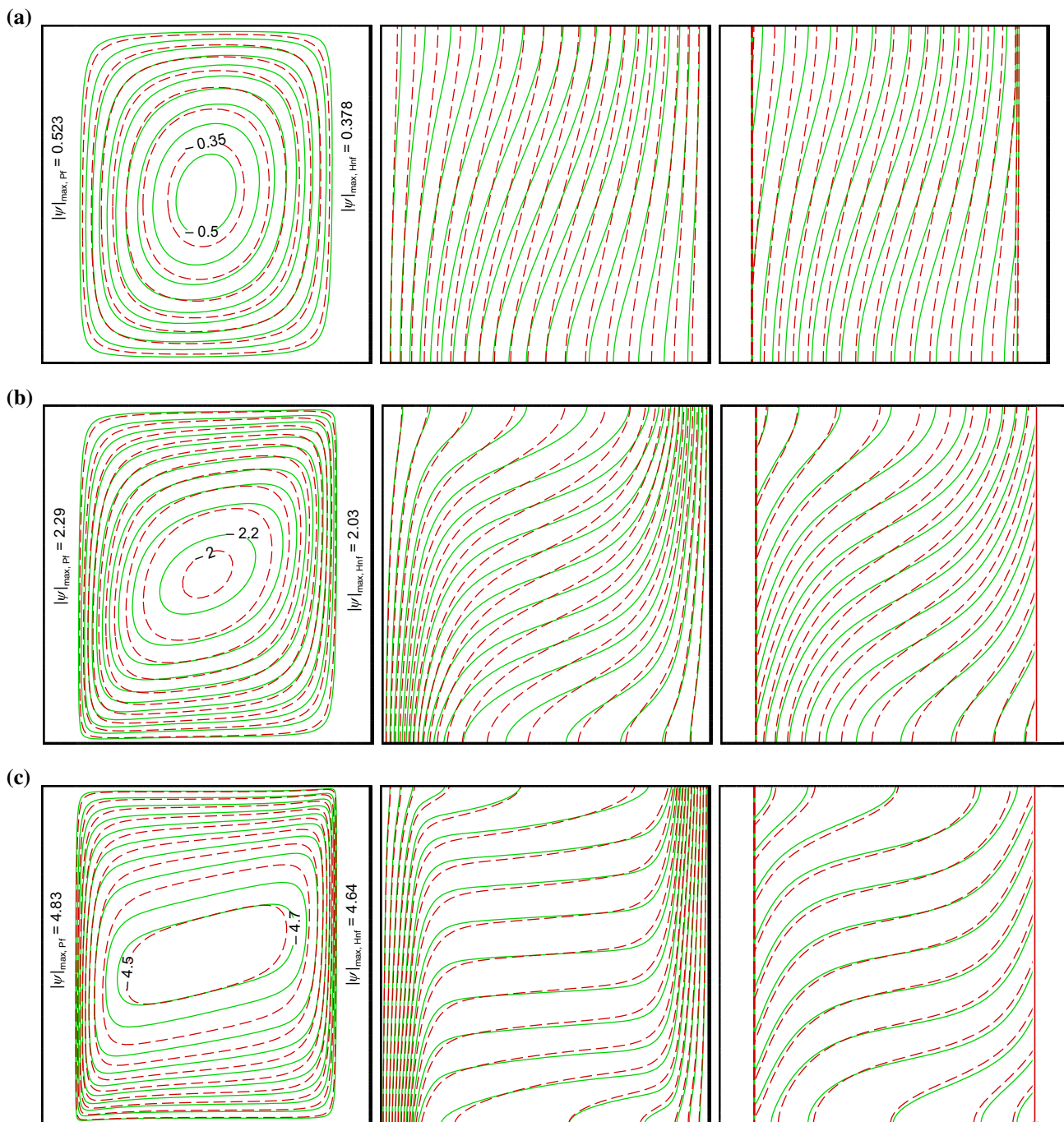
**Fig. 2** Effects of  $R_k$  on the flow, fluid temperature  $T_{hnf}$  and solid temperature  $T_s$  fields at  $Ra = 10^3$ ,  $\varepsilon = 0.5$ ,  $d = 0.1$ ,  $K_r = H = 1$ , **a**  $R_k = 0.1$ , **b**  $R_k = 1$  and **c**  $R_k = 10$

nanoparticles more impress the thermal fields, when  $Ra$  is low.

From Fig. 4, the increment of porosity  $\varepsilon$  increases strongly the size and the strength of the vortex created within the porous media. Indeed, when the porosity  $\varepsilon$  increases, the dynamic resistance resulting from the solid matrix, which was modeled by Darcy term in the momentum equation, is reduced. It is found that the elongation of isotherms along horizontal bounds decreases as  $\varepsilon$

increases; however, the effect of  $\varepsilon$  on the solid matrix temperature field is not so perceptible.

Efficacies of modified thermal conductivity ratio  $K_r$  on the velocity and temperatures patterns have been illustrated in Fig. 5. For this analysis, other characteristics are kept constant so that  $Ra = 10^3$ ,  $d = 0.1$ ,  $R_k = H = 10$  and  $\varepsilon = 0.5$ . At first, when  $K_r$  augments from 0.1 to 1, the intensity of the recirculating flow formed in the porous enclosure slightly increases. Then, the augmentation of  $K_r$

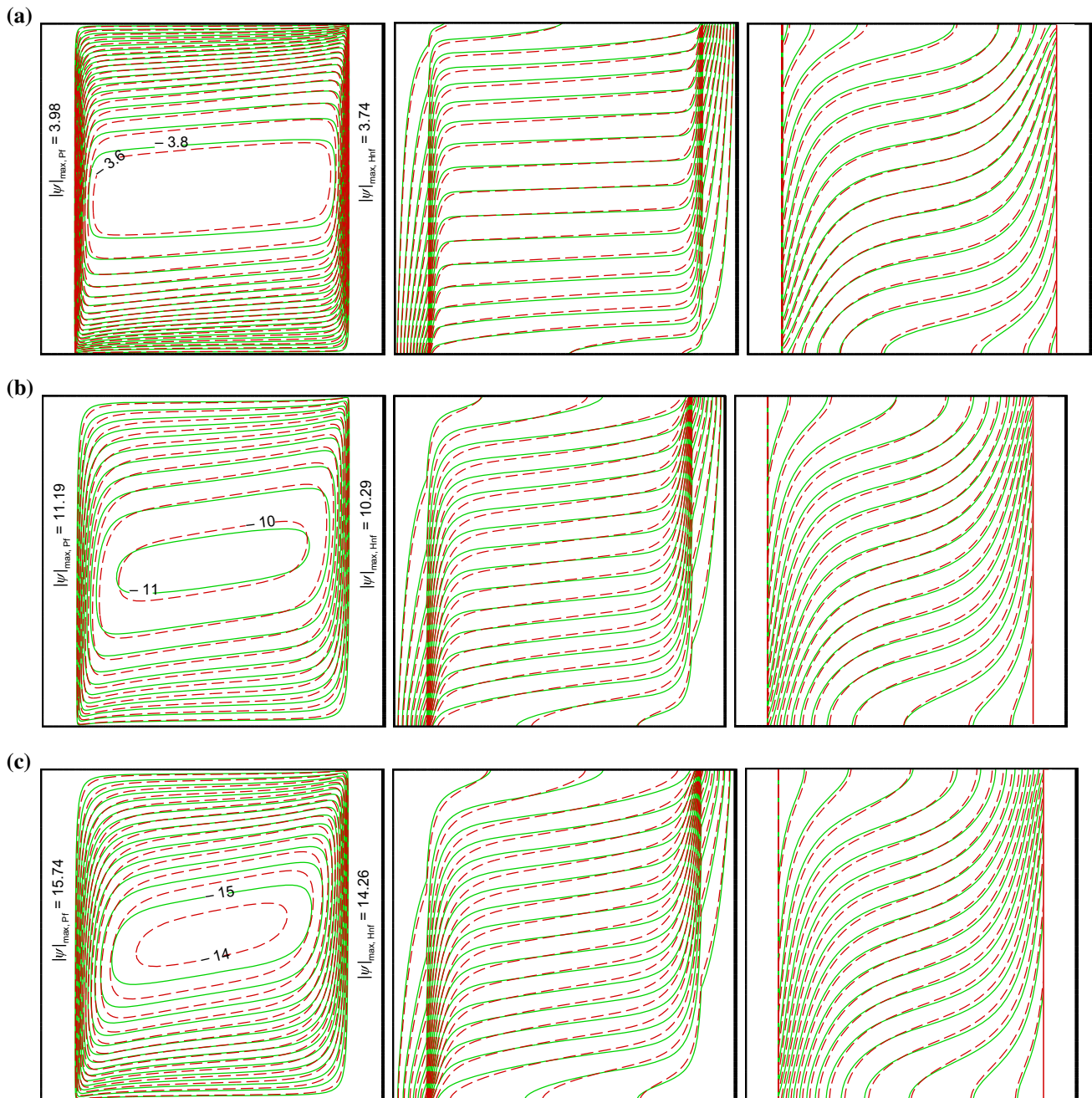


**Fig. 3** Effects of  $Ra$  on the flow, fluid temperature  $T_{hnf}$  and solid temperature  $T_s$  fields at  $R_k = 1$ ,  $\varepsilon = 0.5$ ,  $d = 0.1$ ,  $K_r = H = 10$ , **a**  $Ra = 10$ , **b**  $Ra = 100$  and **c**  $Ra = 1000$

from 1 to 10 declines this characteristic of the flow. This trend can literally be seen in Saied’s work [63]. According to the definition of the  $K_r$ , the ability of the fluid phase for heat transfer enhances as  $K_r$  increases. When  $K_r$  is low, the thermal resistance of the fluid occupying the pores of the porous medium is high; hence, the heat exchange rate between the fluid and solid phases of the porous matrix is low. This fact eventuates a drastic TNE condition between

the fluid and solid phases of the porous matrix. The difference of the configurations of the isotherms ascribed to two phases at the low value of  $K_r$  vouches the above statements. Then, it is observed that these thermal fields become closer to each other by increasing  $K_r$ . According to the above description, it is clear that this trend is a result of decreasing thermal resistance of the fluid phase when  $K_r$  enhances.

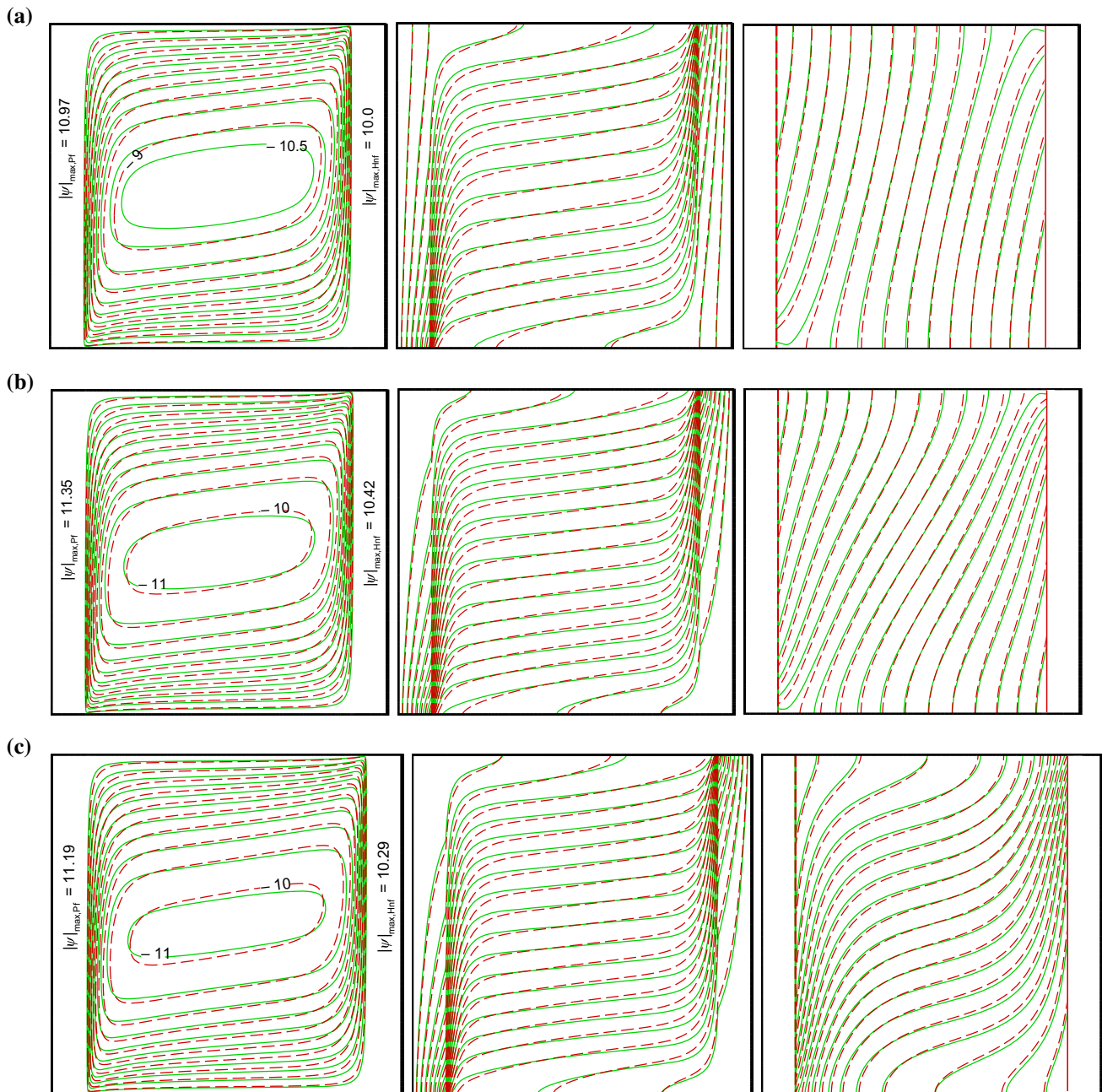




**Fig. 4** Effects of  $\varepsilon$  on the flow, fluid temperature  $T_{\text{hmf}}$  and solid temperature  $T_s$  fields at  $Ra = 10^3$ ,  $d = 0.1$ ,  $R_k = K_r = H = 10$ , **a**  $\varepsilon = 0.1$ , **b**  $\varepsilon = 0.5$  and **c**  $\varepsilon = 0.9$

The efficacies of the solid wall thickness on the flow and temperatures patterns are shown in Fig. 6. With the constant parameters,  $Ra = 100$ ,  $R_k = K_r = 10$ ,  $H = 100$ ,  $\varepsilon = 0.5$ , the strength and the size of the recirculation formed in the porous cavity decreases as the solid wall thickness rises. One argument says that since the wall thermal conductivity is limited, the thermal resistance of the wall is completely related to the thickness of the wall and increases with a growth of the wall thickness.

Consequently, when the wall is thick, less heat can reach the porous media from the hot bound. Decreasing trend of the fluid convection with being thicker solid wall reduces the share of the convection mode in the heat transfer mode and mutually amplifies the share of the conduction mode in the heat transfer process by the fluid phase. Comparing the isotherms of the pure fluid and hybrid nanofluid in a certain case to each other clarifies that the influence of the presence of the hybrid nanoparticles on the temperature fields

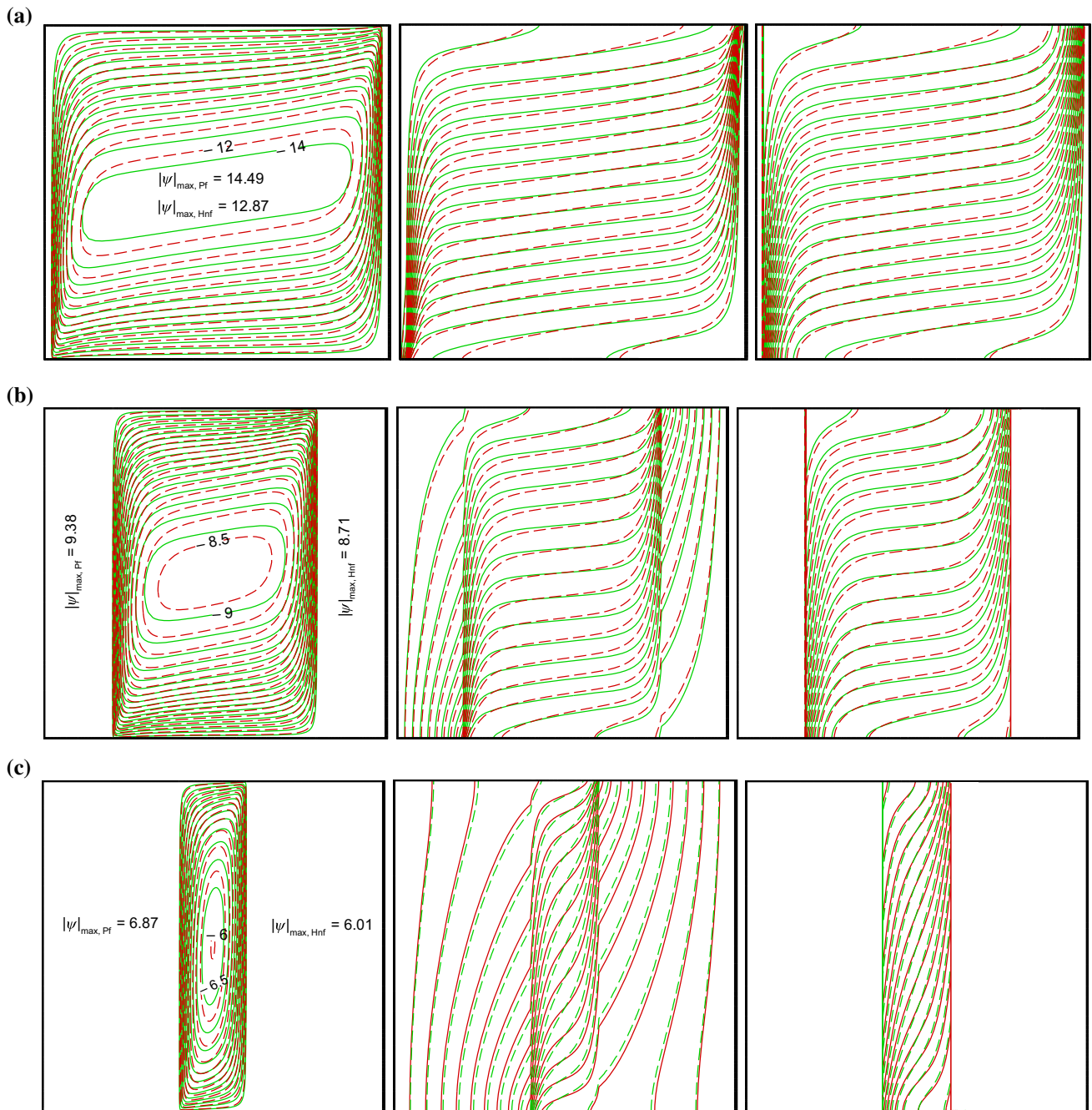


**Fig. 5** Effects of  $K_r$  on the flow, fluid temperature  $T_{\text{hnf}}$  and solid temperature  $T_s$  fields at  $Ra = 10^3$ ,  $d = 0.1$ ,  $R_k = H = 10$ ,  $\varepsilon = 0.5$ , **a**  $K_r = 0.1$ , **b**  $K_r = 1$  and **c**  $K_r = 10$

at the cavity with the thinner solid walls is more than that for the cavities with the thicker solid walls.

The LTNE conditions between the fluid and solid phases of the porous matrix, which are deputized by the Nield number  $H$ , make the different temperature fields for these two phases as can be observed in Fig. 7. In fact,  $H$  is a criterion of the heat exchange rate microscopically between the fluid and solid phases of the porous matrix. An increment in  $H$  augments the heat transfer between the fluid and solid phases of the porous medium. Accordingly, it is said

that the thickness of the thermal boundary layer rises with  $H$ . A detailed look at the fluid temperature field affirms this claim. Additionally, the increment of the interface parameter  $H$  causes a more elongation for the isotherms of the solid phase. Indeed, the growth of the heat exchange between these two phases arising from enhancing  $H$  directs both the thermal fields to a unique thermal field. It is visible that the influences of using hybrid nanoparticles on the isothermal fields of the solid phase are significant at high values of  $H$  ( $H = 100$  and  $1000$ ). It should be noted that the



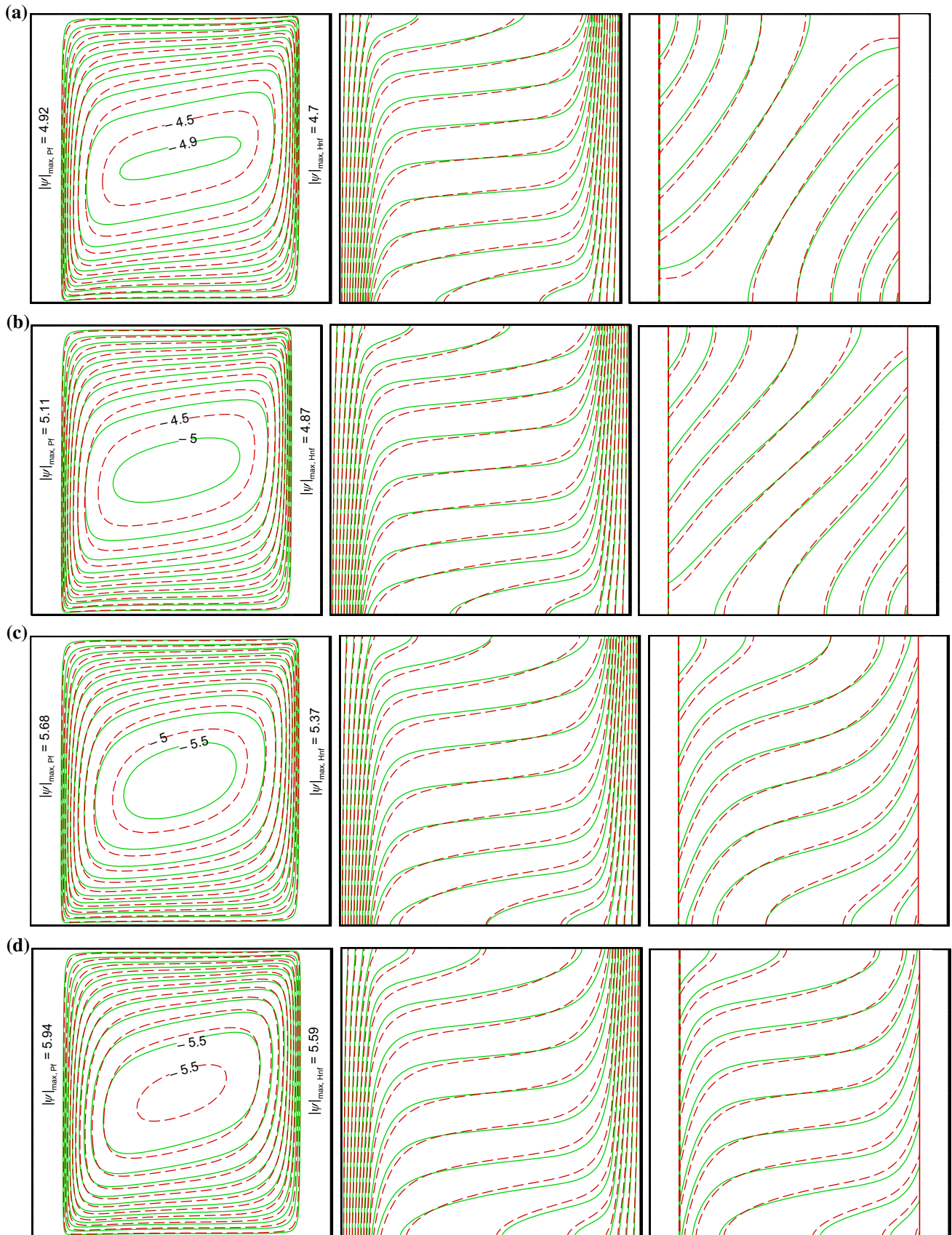
**Fig. 6** Effects of  $d$  on the flow, fluid temperature  $T_{\text{hnf}}$  and solid temperature  $T_s$  fields at  $Ra = 100$ ,  $R_k = K_r = 10$ ,  $H = 100$ ,  $\varepsilon = 0.5$ , **a**  $d = 0.02$ , **b**  $d = 0.2$  and **c**  $d = 0.4$

influence of the hybrid nanoparticles on the reduction of the local temperatures difference of two phases reduces with an increment in  $H$ .

The variations of  $Nu_{\text{hnf}}$  and  $Nu_s$  according to  $Ra$  for various values of the concentration of Ag–MgO hybrid nanoparticles are illustrated in Fig. 8. As shown, using Ag–MgO hybrid nanoparticles declines the rate of heat exchange through the components forming the porous entity. As stated previously, the thermal conductivity and

the dynamic viscosity of the host liquid boost by raising the concentration of the Ag–MgO hybrid nanoparticles suspended. The increment of thermal conductivity and dynamic viscosity of the host liquid with the nanoparticles concentration are known as desirable and undesirable results at the natural convection, respectively. From Table 1, it is seen that the increment of the desirable parameter is significantly less than that of undesirable parameter. Hence, the suspension of the Ag–MgO hybrid



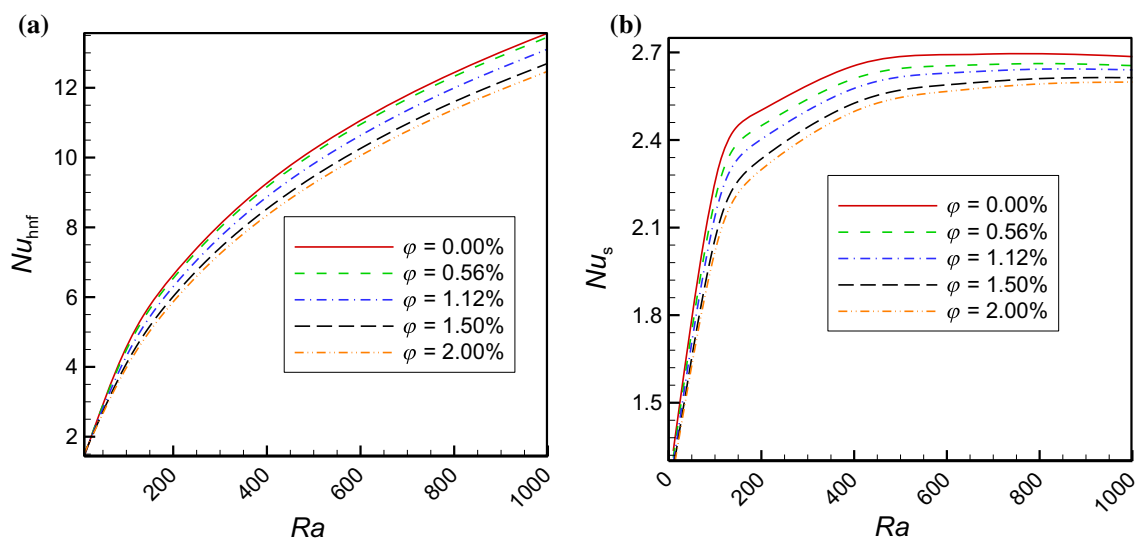


◀ **Fig. 7** Effects of  $H$  on the flow, fluid temperature  $T_{\text{hnf}}$  and solid temperature  $T_s$  fields at,  $Ra = 10^3$ ,  $K_r = R_k = 1$ ,  $d = 0.1$ , **a**  $H = 1$ , **b**  $H = 10$ , **c**  $H = 100$  and **d**  $H = 1000$

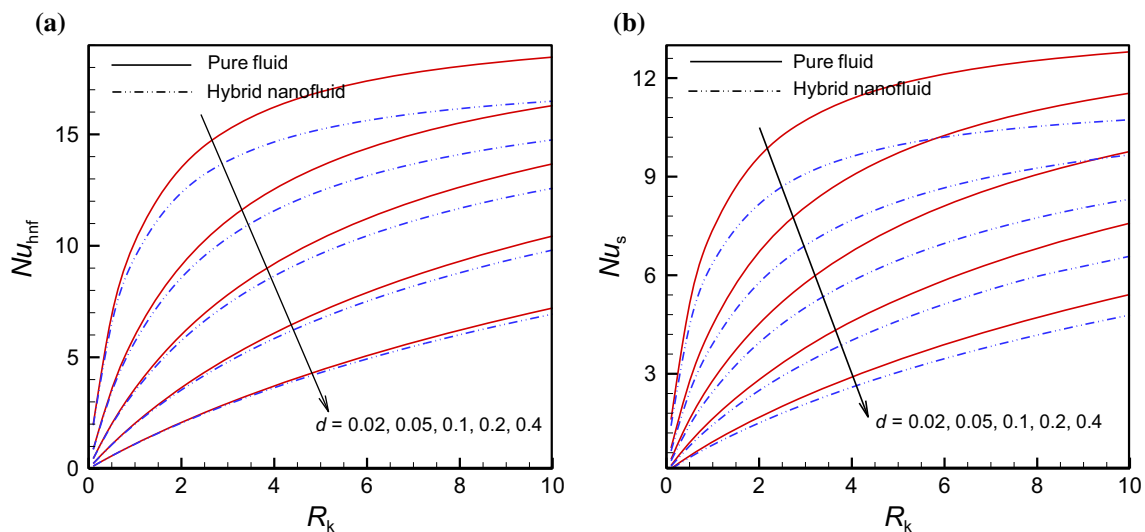
nanoparticles in the host liquid to raise the heat transfer rate cannot be a deliberated operation. In addition, Fig. 8a shows that a growth of  $Ra$  enhances the heat transfer rate carried by fluid phase of the porous medium. However, it can be seen that the increment of  $Ra$  has not significant effects on the  $Nu_s$  when  $Ra > 600$ .

In Fig. 9a, b, the variations of  $Nu_{\text{hnf}}$  and  $Nu_s$  versus  $R_k$  are reported at different values of  $d$  when  $Ra = 10^3$ ,  $\epsilon = 0.5$ ,  $K_r = H=100$ . It is visible that the increment of  $R_k$

increases the heat transfer rates through the fluid and solid phases. This trend is due to that fact that increasing  $R_k$  deals with the increment of the heat exchange through the walls. Additionally, it is observed that whatever the solid wall thickness is less, the heat exchange rate through the solid and fluid phases of the porous medium is more. As said previously, the thermal resistance of the solid wall is completely related to the thickness of the wall and increases with a growth of the wall thickness. Consequently, when the wall is thick, less heat can reach the porous media from the hot bound. It is worth noting here that for a certain thickness of the wall the reduction of  $Nu_s$  due to the use of the hybrid nanoparticles is more than for  $Nu_{\text{hnf}}$ .



**Fig. 8** Variations of  $Nu_{\text{hnf}}$  (a) and  $Nu_s$  (b) according to the  $Ra$  numbers at various values of  $\phi$  when  $\epsilon = 0.5$  and  $R_k = H = K_r = 10$



**Fig. 9** Variations of  $Nu_{\text{hnf}}$  (a) and  $Nu_s$  (b) according to  $d$  at different values of  $R_k$  when  $Ra = 10^3$ ,  $\epsilon = 0.5$ ,  $K_r = H=100$



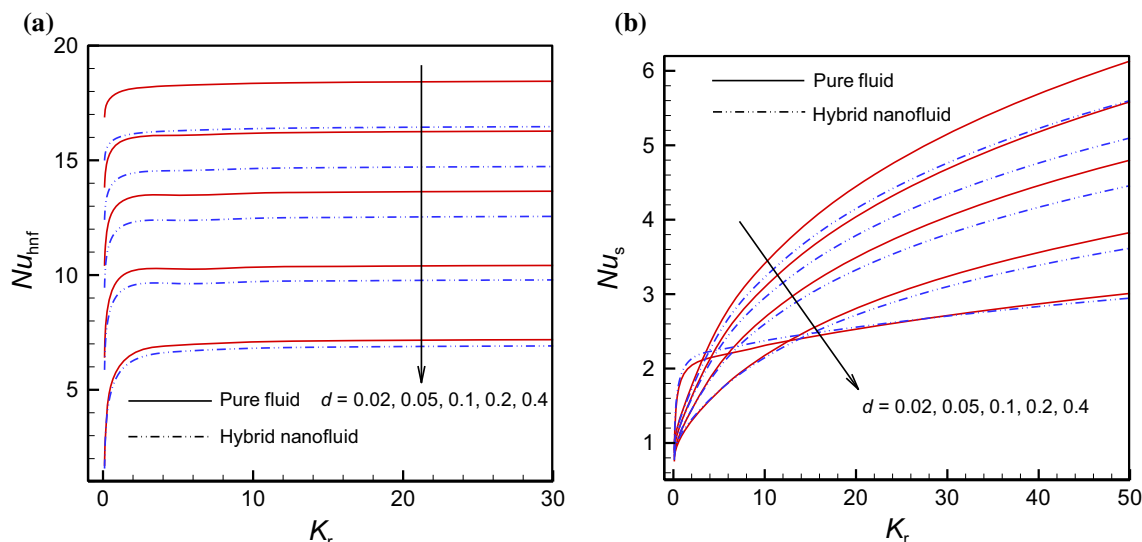


Fig. 10 Variations of  $Nu_{hnf}$  (a) and  $Nu_s$  (b) according to  $d$  at different values of  $K_r$  when  $Ra = 10^3$ ,  $\varepsilon = 0.5$ ,  $R_k = H = 10$

Figure 10a, b displays the influences of the thickness of the solid wall on  $Nu_{hnf}$  and  $Nu_s$  as functions of  $K_r$ . It can be concluded that the heat rate carried by hybrid nanofluid occupying the pores ( $Nu_{hnf}$ ) is increasing when  $K_r$  increases from 0.1 to 3, after which this rate approximately remains constant with a growth of  $K_r$ . In addition, the heat exchange rate through the solid phase  $Nu_s$  increases as  $K_r$  is increasing, while retaining the solid wall thickness fixed. It is obviously seen that the increment of the thickness lessens  $Nu_{hnf}$  for all the values of  $K_r$ . This is despite the fact that the decreasing behavior  $Nu_s$  with the increase in the thickness  $d$  only can be seen for thicknesses of less than or equal to 0.2. As shown in Fig. 10b, depending on the value of  $K_r$ , an increase in the wall thickness to 0.4 can increase or decrease  $Nu_s$ . Here it can also be seen that dispersing

Ag–MgO hybrid nanoparticles declines  $Nu_s$  with the exception  $d = 0.4$ . Using the hybrid nanoparticles slightly raises  $Nu_s$  when  $K_r < 18$ , while the nanoparticles can enhance  $Nu_s$  for  $K_r > 42$ .

From Fig. 11a, b, the increase in  $H$  declines and elevates  $Nu_{hnf}$  and  $Nu_s$ , respectively, keeping the thickness of solid wall constant. The increment in  $H$  augments the heat transfer between the fluid and solid phases of the porous medium, consequently, the thickness of thermal boundary layer increases. Hence, the temperature gradient inside the boundary layer declines which means the reduction of  $Nu_{hnf}$ . As illustrated in Fig. 11b, it can be observed that  $Nu_s$  increases when  $H$  grows. This result can easily be justified due to this fact that the temperature gradient of the porous matrix near the vertical bounds amplifies as a result of the

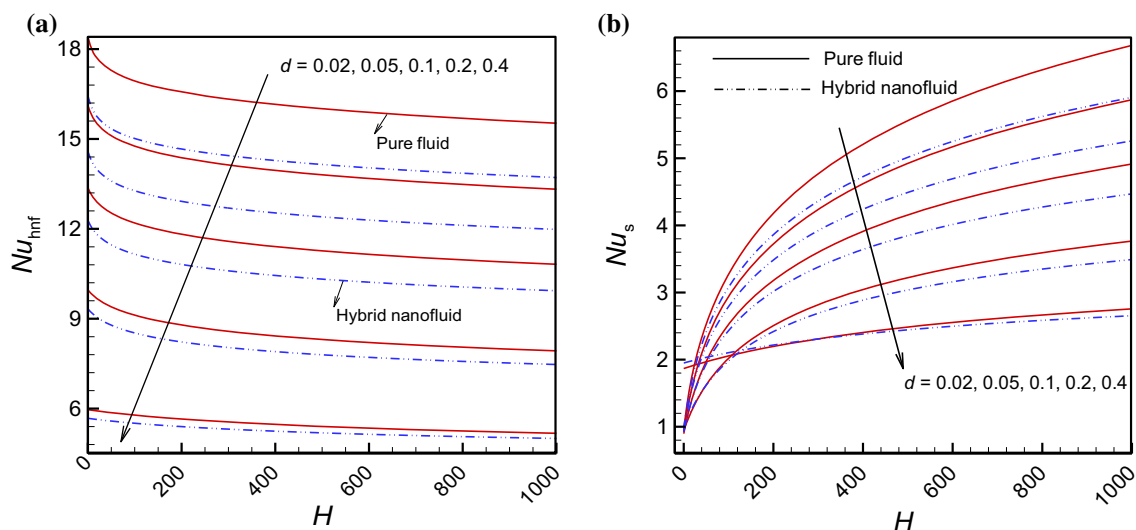


Fig. 11 Variations of  $Nu_{hnf}$  (a) and  $Nu_s$  (b) according to  $d$  at various values of  $H$  when  $Ra = 10^3$ ,  $\varepsilon = 0.5$ ,  $R_k = 10$  and  $K_r = 1$

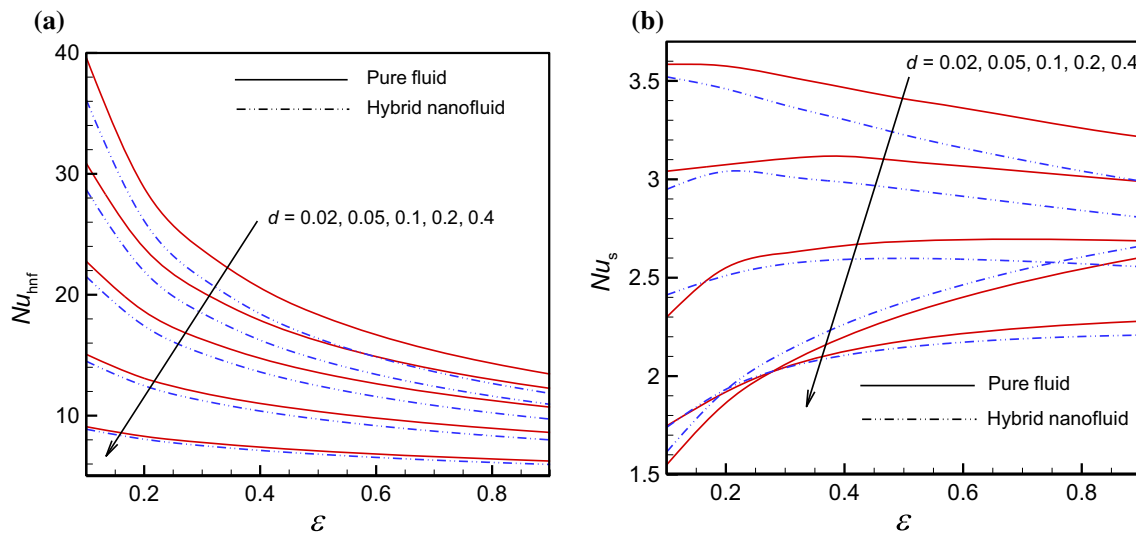


Fig. 12 Variations of  $Nu_{hnf}$  (a) and  $Nu_s$  (b) according to  $d$  at different values of  $\epsilon$  when  $Ra = 10^3$ ,  $R_k = H = K_r = 10$

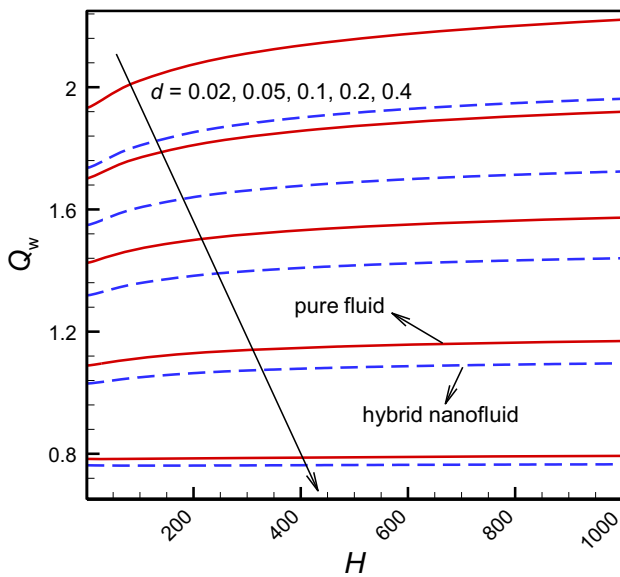


Fig. 13 Variations of  $Q_w$  according to  $d$  at various values of  $H$  for  $Ra = 10^3$ ,  $\epsilon = 0.5$ ,  $R_k = 10$  and  $K_r = 1$

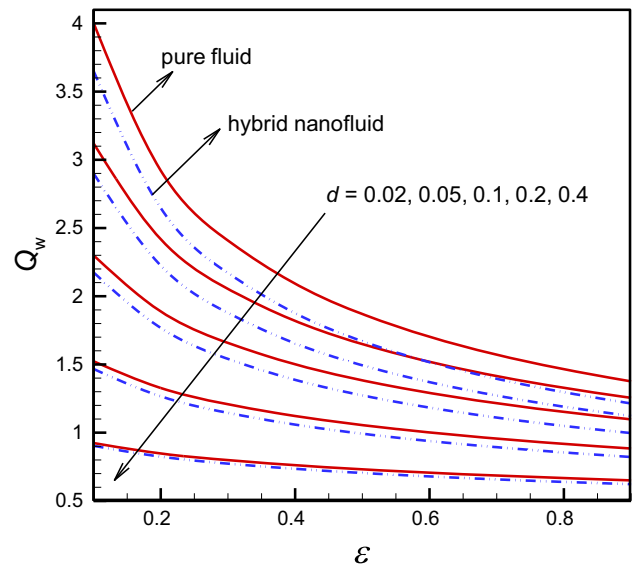


Fig. 14 Variations of  $Q_w$  according to  $d$  at different values of  $\epsilon$  when  $Ra = 10^3$ ,  $R_k = H = K_r = 10$

enhancement of  $H$ . For a cavity with  $d = 0.4$  and  $H < 300$ , dispersing the hybrid nanoparticles Ag–MgO in the host fluid can amplify the heat transfer rate through the solid phase. However, if  $H > 300$ , the use of the nanoparticles debilitates this rate.

The results represented in Fig. 12a illustrate that for different values of  $d$ , the increase in porosity  $\epsilon$  scripps  $Nu_{hnf}$ . Nevertheless, as shown in Fig. 12b, the different trends can be observed for the variations of  $Nu_s$  versus porosity  $\epsilon$  as the thickness of solid wall varies. Although in the low value of  $d$  ( $d = 0.02$ ), the increase in porosity

declines  $Nu_s$ , when  $d = 0.2$  or  $0.4$ ,  $Nu_s$  continuously augments with increasing  $\epsilon$ . Further, there is no a certain trend for the variations of  $Nu_s$  with increasing  $\epsilon$  when  $d = 0.05$  and  $0.1$ .

Figure 13 shows the variations of heat transfer rate across the solid wall  $Q_w$  versus  $H$  for various values of  $d$  when  $Ra = 10^3$ ,  $\epsilon = 0.5$ ,  $R_k = 10$  and  $K_r = 1$ . According to the above presented results, it is expected that the increment of the solid wall thickness declines the heat exchange rate  $Q_w$ . This fact has obviously been illustrated in Fig. 13. Also  $Q_w$  increases with the increment of  $H$ . It is clear that the effect of  $H$  on the increase in the rate of  $Q_w$  is

more essential when the solid wall thickness is low. Figure 14 displays the variations of heat transfer rate across the solid wall  $Q_w$  versus  $\varepsilon$  for various values of  $d$  when  $Ra = 10^3$ ,  $R_k = H = K_r = 10$ . As shown, the increment in  $\varepsilon$  strongly declines  $Q_w$ . In addition, it can be said that the decreasing trend  $Q_w$  by using the Ag–MgO hybrid nanoparticles is more essential at high values of  $\varepsilon$ .

## Conclusions

Current investigation studies the conjugate natural convection inside a porous enclosure filled with Ag–MgO hybrid nanofluid using LTNE model. Two solid walls located in two sides of cavity introduced as conductive interface between hot and cold walls, and besides, the top and bottom bounds have been insulated. The governing differential equations are written using the Darcy law and then transformed into a non-dimensional form for better representation of the results. Governing equations have been solved by the finite element method. Various amounts of the Darcy–Rayleigh number ( $Ra = 10$ – $10^3$ ), volume fraction of hybrid nanoparticles ( $\varphi_{hnf} = 0$ – $0.02$ ), porosity ( $\varepsilon = 0.1$ – $0.9$ ), interface parameter ( $H = 1$ – $10^3$ ), thermal conductivity ratio ( $K_r = 0.1$ – $10$ ), and the width of the solid wall ( $d = 0.1$ – $0.4$ ) are examined to perform calculations. The most significant findings are listed below:

- Using the hybrid nanoparticles decreases the flow strength. The heat transfer rate increases when  $R_k$  rises. When  $R_k$  is low, the influences of the hybrid nanoparticles on the temperatures fields and the relative decreasing of the flow strength are, respectively, more and less than those compared to the high values of  $R_k$ ,
- As  $Ra$  enhances, the flow strength becomes more. When  $Ra$  is low, the hybrid nanoparticles more impress the thermal fields. Besides, increment of  $\varepsilon$ , increases strongly the size and the strength of the vortex created within the porous media.
- When  $K_r$  augments from 0.1 to 1, the strength of recirculation formed slightly increases, whereas the augmentation of  $K_r$  from 1 to 10 declines that. When  $K_r$  is low, the heat transfer rate is low and by increasing  $K_r$ , thermal fields become closer to each other. The strength and size of the recirculation formed decreases as  $d$  increases. The effect of hybrid nanoparticles on thermal fields with the thinner solid walls is more than that the thicker ones.
- An increment in  $H$  leads to the heat exchange enhancement. The influences of hybrid nanoparticles on the solid phase isothermal fields are significant at high magnitudes of  $H$  and on the reduction of two

phases temperatures difference reduces with increasing  $H$ .

- Using hybrid nanoparticles declines the heat exchange rate. By raising the hybrid nanoparticles concentration,  $Nu_{hnf}$  and  $Nu_s$  decreased for constant  $Ra$ . Besides, increase in  $Ra$  enhances the  $Nu_{hnf}$  and  $Nu_s$ . The increment of  $R_k$  and  $d$  augment the heat transfer. For a certain  $d$ , the reduction of  $Nu_s$  due to using the hybrid nanoparticles is more than that for  $Nu_{hnf}$ . Growth of  $Ra$  enhances the heat exchange rate. However, the increment of  $Ra$  has not significant influences on  $Nu_s$  when  $Ra > 600$ .
- $Nu_{hnf}$  enhances when  $K_r$  increases from 0.1 to 3, after which this rate approximately remains constant. Moreover,  $Nu_s$  increase with augment of  $K_r$ , while  $d$  retaining fixed. The increment of  $d$  lessens  $Nu_{hnf}$  for all values of  $K_r$  and not has specific trends for  $Nu_s$ . Dispersing hybrid nanoparticles declines  $Nu_s$  except for  $d = 0.4$ , raises  $Nu_s$  when  $K_r < 18$ , while it can enhance  $Nu_s$  for  $K_r > 42$ .
- The increase in  $H$ , respectively, declines and elevates  $Nu_{hnf}$  and  $Nu_s$ , keeping  $d$  constant. For all values of  $d$ , the increase in  $\varepsilon$  scrimps  $Nu_{hnf}$ . Although in low value of  $d$  ( $d = 0.02$ ), the increase in  $\varepsilon$  declines  $Nu_s$ , but at higher values,  $Nu_s$  continuously augments.
- For different values of  $d$ , the increase in  $\varepsilon$  scrimps  $Nu_{hnf}$ ; however, various trends can be seen for the variations of  $Nu_s$  versus  $\varepsilon$  as  $d$  varies. The increment of  $d$  and also  $\varepsilon$  declines the heat transfer rate. In addition,  $Q_w$  increases with the increment of  $H$ .

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