Efects of thermophysical, hydrodynamics and thermal characteristics of well stable metallic nanofuids on energy transportation

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Abstract

The current research focuses on the chemical synthesis of ZnO and preparation of ZnO dispersed in EG@DW mixturebased nanofuids for the investigation of enhancement in heat transfer of a square shaped heat exchanger. The steady and uniform dispersion of ZnO nanoparticles was achieved using high probe sonication at varying mass% concentrations like (0.1; 0.075; 0.05 and 0.025 mass%). Also, the ZnO particles were synthesized using sonochemical technique, where sodium hydroxide (NaOH) and zinc acetate ($ZnC_4H_6O_4$) were used as raw materials. For the verification of ZnO synthesis numerous characterizations like UV–Vis, XRD, FTIR, and FESEM analysis were carried out. The changed mass% concentrations of ZnO-EG@DW-based nanofuids were considered to check their efects on thermophysical properties, dispersion, stability, and heat transfer (Nusselt Numbers) at varying Reynolds numbers ranging from 4550 to 20,360 using square heat exchanger. A positive increase in thermal conductivity was observed by the addition of ZnO in the ZnO-EG@DW with (50:50) mixture. The highest thermal conductivity of 1.0 W m⁻¹ K⁻¹ was noticed at a maximum of 0.1 mass% concentration which is greater than base fluid (EG@DW) at 45 °C. The maximum pressure drop of 2400 Pa m^{-1} at Reynold number 20360 and higher friction loss of 0.045 for Reynolds number 4550 was noticed at 0.1 mass% concentration. The highest improvement in heat transfer was recorded at 0.1 mass% is 550–2830 W m⁻¹ K which is 63% greater of the base fluid, while others mass% illustrate improved heat transfer coefficient of about 550–2170, 550–1750, and 550–1500 W m⁻² K⁻¹, respectively, at highest Reynolds (Re) when it compared with the base fuid (EG@DW). All the heat transfer related experiments were executed at 30 °C room temperature where constant heat fux of 10,886 W m−2 and eight varying fow rates were used.

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Graphical abstract

Keywords ZnO synthesis · Nanofluids preparation · Thermal conductivity · Viscosity · Heat transfer coefficient

List of symbols

- EG Ethylene glycol
- *N*_P Nanoparticles
- $\frac{N_{\text{f}}}{\text{DW}}$ **Nanofluids**
- Distilled water
- C_f
Cu Friction coefficient
- Copper
-
- C_p Specific heat
 d_p Nanoparticles *d*_p Nanoparticles diameter
Conc Concentration
- Concentration
- *K* Thermal conductivity
- *R* Ratio of materials
T Temperature
- *T* Temperature
- *h* Convective heat transfer coefficient
- Nu Nusselt numbers
- *Q* Heat fux
- *L* Length of medium

Greek symbols and subscript

-
- *W* Wall
F Fluid Fluid
- **Base fluid**
- $rac{B_f}{T}$ *T* Turbulence
- FF Friction factor
- *Ρ* Density in kg m−3
- μ Dynamic viscosity in N m s⁻¹
- Φ % Volume friction
- *V* Kinematic viscosity $m^2 s^{-2}$
- *ɸ* Volume friction

Introduction

The ever increasing demand for efficient heat transportation solutions has come to be a key challenge for engineering and manufacturing productions especially in power and energy

sectors. The aim of this research is based on the application of diferent mediums in the heat transfer process. Usually, normal or distilled water was used previously as a heat carrier specifcally in oil and gas refneries, coal power plants, nuclear plants, and heating and cooling systems, etc. [[1](#page-20-0)]. However, it is observed that the conventional fuids show few drawbacks, specifcally on their aspect of heat transfer capability. The results arise from conventional fuids subjected to integrally less thermal conductivity which can be a major cause of attaining low heat exchanges [\[2](#page-20-1), [3](#page-20-2)]. Therefore, this loss has encouraged several researchers to determine alternative approaches for the improvement of heat transfer in diferent heat exchangers with their shapes, sizes, surfaces, etc., with improved heat transport characteristics $[4–6]$ $[4–6]$ $[4–6]$.

The heat exchangers with varying geometries are being used in engineering industries for heat transfer, which are commonly known as energy-efficient vehicles with less maintenance cost and can be used for a long duration [[7](#page-20-5)]. These heat exchangers are mostly designed according to the apparatus characteristics comprising the fluid, base fuid temperature, heat fux, fow rates, and their dimension as well [\[8](#page-20-6)]. Due to their particular shape and design, heat exchangers can be classifed into diferent categories such as heat exchangers with wick, thermosyphon heat exchangers, looped heat exchangers, and looped thermosyphon. Each category of the heat exchanger is suitable for specifc applications due to their performance, heat transfer rate, and an impartial system for cooling and due to their installation reasons [\[9](#page-20-7)[–11](#page-20-8)].

The more quality of nanofluids as heat transportation fluids could attain by adding the highly conductive nanoparticles (solids) in varying conventional fuids. Choi et al. [[12\]](#page-20-9) was the frst researcher who coined the term nanofuids, he realized that homogeneous and uniform dispersion of solid nanoparticles in conventional base fuids likewise water, distilled water, Polyethylene glycol (PEG), Palm oil (PO), EG, Diathermic oil (DTO), Transformer oil (TO), glycerin, etc., may illustrate the remarkable improvement in their thermal performances characteristics [[13,](#page-20-10) [14\]](#page-20-11). The metal oxides, ceramics, carbon structure, and various bio-based particles were used earlier to prepare varying nanofuids. These nanofluids were included Al_2O_3 , ZnO, TiO₂, SiO₂, Al/Al_2O_3 , Cu/Cupric, CNTs, SWCNTs, and MWCNTs, etc. [[15](#page-20-12), [16](#page-20-13)]. Several investigations were conducted on heat transfer improvement by mixing diferent nanoparticles in normal or distilled water. The importance of distilled water and ethylene glycol as Nano-coolants with the addition of ZnO nanoparticles [[17](#page-20-14)]. The addition of sonochemically synthesized ZnO nanoparticles in EG+DW base fuid with a stable dispersion and longer suspension stability will be a new approach to see heat transfer development of the square heat exchanger [[18–](#page-20-15)[20](#page-20-16)].

Hosseini et al. $[21]$ used Graphene oxide and Al_2O_3 water-based nanofluids at 0.2 mass% concentration of each in the parabolic trough solar collector (PTSC) for the improvement of thermal efficiency. Varied thermo-physical characteristics like dynamic/kinematic viscosities, densities, specifc heat and thermal conductivities were also analyzed here. The maximum 23.4% in dynamic viscosity and 9.1% enhancement in thermal conductivity were reported here. Furthermore, three diferent 1, 3, and 5 L/mint Graphenebased and Al_2O_3 -based nanofluids were tested for thermal efficiency, where 63.2% for Graphene oxide and 32.1% for Al_2O_3 enhancement at 1 mass% have been reported.

Said, Z et al. [[22\]](#page-20-18) investigated the TiO₂- and Al_2O_3 -based nanofuids for thermal enhancement of the radiator. The solid TiO₂ and Al_2O_3 nanoparticles were dispersed in water and EG at a 50:50 ratio each, respectively. Further, diferent characterizations and thermo-physical properties were analyzed to measure the thermal performance at a 0.3% volume fraction of each. The choice of these metal oxide-based nanofuids was chosen due to their anti-corrosive properties. The highest improvement in heat transfer was observed about 24.21% for Al_2O_3 -based nanofluids, which is more than water-, EG-, and $TiO₂$ -based nanofluids.

Saedodin, S et al. [\[23\]](#page-20-19) has used varying metal oxidebased and turbulator Nano-fuids for heat transfer improvement in a straight tube based on empirical correlation data. The SiO2-/water-based nanofuids were prepared at varying 0.5–2% volume concentrations, where all the volume concentrations have been investigated for the pressure drop and heat transfer improvement. The thermophysical characteristics like dynamic/kinematic viscosity and thermal conductivity were investigated as well. Therefore, based on all these properties the maximum 7.93% heat transfer improvement and 13.96% surge of pressure drop were recorded at a maximum 2% volume concentration. Also, overall 4.2% thermal performance was reported here. Further, this enhancement was compared with metal oxide type nanofluids like $TiO₂$, Al_2O_3 , CuO, and SiO₂ at the same highest 2% volume concentration. In conclusion, the CuO-/water-based nanofuids had better thermal performance as compared to all other nanofluids.

Mahani, R.B et al. $[24]$ $[24]$ studied the heat transfer improvement by combining factor efects, including 3D modeling with varying geometries, and using hybrid nanofluids of Al_2O_3 and Cu at high Reynolds value. The outcomes describe that hybrid nanofuids, Reynolds numbers, and turbulators were infuenced to augment the heat transfer enhancement of the rectangular turbulators. By changing the size of turbulators, nanofuids volume fraction, and Reynolds numbers from lower to higher would cause to increase in heat transfer up to 49.17% of the total. The suggested

factors are more practical and can be benefted in the feld of nanofuids and fuid mechanisms.

Nafchi, P.M., et al. used ethylene glycol (EG) as a base fluid or dispersant as an antifreeze agent in various engineering-based applications like cooling systems, heat exchangers, automobile radiators, solar collectors, refrigeration, etc. Varying thermophysical and thermal properties were analyzed and outcomes are reported accordingly. However, it has been noticed that this base liquid showed less thermal conductivity as a working fuid in thermal engineering systems [[25](#page-20-21), [26](#page-20-22)]. Furthermore, similarly, numerous studies were executed to improve its thermal conductivity and other thermophysical properties such as viscosity, density, and specifc heat that can play a vital role in enhanced heat transportation. For this purpose diferent solid nanoparticles have been used to prepare the ethylene glycol (EG)-based nanofluids at varying mass% concentrations [[27,](#page-20-23) [28\]](#page-20-24).

Also, several researchers have investigated the effects of metal oxide nanoparticles in ethylene glycol (EG) or its mixture on the heat transfer, the results showed improved heat transfer amount with an escalation in concentrations of solid nanoparticles. The addition of highly conductive Nanosized solid particles in ethylene glycol (EG) or its mixture with water gives an escalation in the thermal conductivity, which may lead to augmented heat transfer [[29,](#page-20-25) [30](#page-20-26)]. Further, many researchers have revealed the benefts of using diferent nanofuids in engineering applications such as boiling heat transfer, machining, convective heat transfer, micro-heat exchangers, and cooling [[31\]](#page-21-0).

The ultra-sonochemical is a very common technique that has been used for Nano-size particle synthesis in bulk and their varying nanofuids for diferent purposes. This technique is very useful to produce primarily nanoparticles of regular shape and size, and it is a less time-consuming method as compared to other techniques. Further, solid metal oxide nanoparticles need high sonication to disperse them uniformly and homogeneously in the base fuid. So, for this purpose, the high probe sonication or ultra-sonochemical technique is the best choice, based on these parameters they would prefer this technique throughout the experiment.

Research signifcance

The previous studies showed a lack of longer dispersion and stability of nanofuids, and they used varying stabilizing agents/surfactants that could drop their thermal conductivities. Still, none of the studies has reported varying improved thermophysical and heat transfer properties of the nanofuids together. Also, pure ethylene glycol-based nanofuids showed higher viscosity which needs additional pumping power which may drop down the efficiency of the

system. Similarly, the only distilled water-based nanofuids have dispersion and stability problems. The current study was conducted considering the following key objectives.

- The key objectives of this study are to synthesize ZnO by sonochemical procedure and then using a two-step method for the preparation of surfactant-free stable and homogeneously dispersed ZnO-EG@DW-based nanofuids at diferent mass% concentrations of (0.1; 0.075; 0.05 and 0.025 mass%) EG@DW (50:50)-based fuid.
- The current study intends to achieve supreme heat transfer "h" in a square shaped heat exchanger. Moreover, the mixture of EG@DW with (50:50) ratio was used as a base fuid for nanofuids preparation.
- A high probe sonication procedure was implemented to enhance and prolong the dispersion/suspension and maximum stability of ZnO in the mixture of EG@DW base fuid. This distinct procedure increases the solid particle dispersion in EG@DW without using any surfactant/stabilizing agent, which can lift their stability.
- In the current study, changing flow rate and uniform heat flux conditions and their effects on the increase in heat transfer (Nusselt numbers) at several localities of square heat exchanger in the turbulent regimes were considered for the EG@DW and ZnO-EG@DW nanofuids at (0.1; 0.075; 0.05 and 0.025 mass%) concentrations.

Methodology and materials

Materials

The zinc acetate dihydrate $Zn(CH_3CO_2)$, $2H_2O$ with (99.5%) purity were used as a precursor for ZnO synthesis, sodium hydroxide (NaOH) analytical reagent with (99.%) purity was used as a strong base for chemical reaction with precursor salt, Ethylene glycol (EG) and distilled water (DW) were procured from Sigma Aldrich, Sdn Bhd Selangor Malaysia. Ethylene glycol (EG) with molecular weight (M Wt 62,07) g mol and distilled water were used to synthesis the ZnO nanoparticles by using a one-pot sonochemical procedure. All these chemicals used for synthesis were of analytical grades. Furthermore, a 2-step probe sonication process was adopted to prepare the well dispersed ZnO-EG@DW-based nanofuids at varying (0.1, 0.075, 0.05, and 0.025 mass%) concentrations.

Synthesis of ZnO nanoparticles

The one-pot sonochemical synthesis procedure was used to synthesize the ZnO nanoparticles. Our synthesis process is quite simple and aims at realizing the efficiency of sonochemical waves produced by sonicator and the use of precursors and bases in controlling the size. For ZnO synthesis, 1 M zinc acetate $Zn (CH₃COO)₂H₂O$ precursor were added in a mixture of EG@DW with 50:50 and stirred until completely dissolved. A 2 M sodium hydroxide (NaOH) base was added in EG@DW mixture with 50:50 ratios under continuously stirring to prepare the reaction solution. Further, the precursor solution has been kept under probe sonication and the sodium hydroxide NaOH solution was added drop by drop (1 drop/s) into zinc acetate solution under continuous probe sonication for 2 h shown in Fig. [1.](#page-4-0) The probe amplitude was attuned at 70 with 3/2 s off/on time. The maximum 750 W probe power, 36,000 J delivered energy by the probe, 220 input voltage was selected and the probe temperature was kept at 0 °C. By mixing the NaOH solution into the zinc acetate Zn $(CH_3COO)_2$ H₂O solution, the chemical reaction has started and white precipitate started to form in the EG@DW mixture. After adding the NaOH solution to the zinc acetate solution, both solutions were left for continuous sonication for 3 h. After completing 3 h, these white precipitates were completely transformed into dense white precipitates. After sonication, the dense white precipitates were washed repeatedly with DW and ethanol and fnally dried in a vacuum oven for 2 h at 70 °C. Finally, the dried white powder was kept for calcination in a furnace at 200 °C, and ZnO nanoparticles were achieved with spherical morphology, which later on was characterized and confrmed. The chemical reduction process for ZnO nanoparticles is as follows [\[32\]](#page-21-1).

$$
Zn(CH_3COO)_2.2H_2O \stackrel{\Delta}{\rightarrow} Zn(OH)_2 + 2CCH_3OOH
$$
 (1)

$$
Zn(OH)_2 \stackrel{Na^+OH^-}{\iff} (Zn[OH]_4)^{-2} + Na^+ \tag{2}
$$

Fig. 1 Synthesis fow of ZnO solid nanoparticles

$$
(\text{Zn}[OH]_4)^{-2} \stackrel{\text{H}^+OH^-}{\iff} (\text{Zn}[OH]_4)^{-2} + H^+ \tag{3}
$$

$$
(\text{Zn}[OH]_4)^{-2} \overset{\text{yields}}{\rightarrow} \text{ZnO} + \text{H}_2\text{O} + 2\text{OH}^-
$$
 (4)

Synthesis of ZnO‑EG@DW‑based nanofuids with the help of probe sonication

The above presented synthesis describes the use of sonochemically synthesized ZnO nanoparticles that are physically blended by probe sonication method in a mixture of EG@DW with (50:50) ratio to prepare ZnO-EG@DW-based nanofuids and then run them in a single tube square shaped heat exchanger for to improve the heat transfer. The ZnO nanoparticles were created by using a facile one pot sonochemical method. After synthesis, the ZnO nanoparticles were dispersed in a mixture of EG@DW for 2 h continuously to obtain uniform and homogeneous dispersion without adding any kind of stabilizing agent and surfactant. Four changed (0.1, 0.075, 0.050, and 0.025 mass%) concentrations of the ZnO nanoparticles were dispersed in the EG@ DW mixture and highly sonicated by using probe sonication. Also, the 2 h sonication is the preferred duration for equal/uniform ZnO dispersion in the base fuid, so this way dispersion and suspension increase up to three months without using any surfactant. Figure [2a](#page-5-0) is showing varying base fuids like DW, Ethylene glycol (EG), and a mixture of both DW and EG. The mixture of both EG@DW at (50:50) base fuid was used to prepare the ZnO-EG@DW-based nanofuids at varying mass% concentrations. The 2-step line flow of nanofluids preparation is shown in Fig. [2b](#page-5-0), where the frst step showing the ZnO synthesis by using one pot sonochemical technique and then were blended in base fuid EG@DW under constant high probe sonication. Finally, the four well dispersed and stable at diferent (0.1; 0.075; 0.050; and 0.025 mass%) of ZnO-EG@DW-based nanofuids were prepared.

Test section geometry

A square heat exchanger is a design of stainless steel having 1200 mm long and 0.01 m hydraulic diameter from both ends is a signifcant part of the heat transfer experimental test rig. The heated area of the square heat exchanger was wrapped by electrical insulating tape up to 1200 mm from the start point to end. The maximum delivered power by heater was 600 W capacity; it has been controlled by using voltage regulating supply. A heavy coating of insulating wool was whorled over the square heat exchanger and fully shielded by an aluminum sheet to evade the heat releasing

Fig. 2 a Distilled water (100%), Ethylene Glycol (EG-100%), and Ethylene Glycol and Distilled water (EG@DW-50:50), and **b** Two-step line flow for the preparation of ZnO-EG@DW-based nanofluids

by heat exchanger outer surfaces. K-Type five sensitive thermocouples were attached at various points over the surface of the square heat pipe and separated from each other with an equal distance of 0.2 m of the total length as shown in Fig. [3](#page-6-0)a. Close to the nanofuids fow channel, two additional inlet-outlets, thermocouples were connected with a square heat pipe to enumerate the bulk temperature. Figure [3a](#page-6-0) is showing the 2-D schematic view and complete insulated (Glass wool) square heat exchanger, while Fig. [3b](#page-6-0) is showing a pictorial view of the complete heat transfer test rig showing varying parts along with heat exchangers.

Experimental

The experiments were executed over a well designed heat transfer test rig as given in Fig. [4.](#page-7-0) The test rig includes the main nanofuid fow loop along the diferent bypass, nanofuid pump, fow meter, inlet–outlet, nanofuid, and fuid tank. Further, the pressure meter, valves for heat exchangers, main valve, pressure gauge, K-type thermocouples, chiller, power supply, data logger GRAPHTECH-GL/220, etc., are also the important parts. The voltage regulator, control panel, frequency control, heater control, and diferent indicators on the control panel were all separately connected to the setup. All the parts and the square heat exchanger are shown **Fig. 3 a** The 2-D schematic view and complete insulated (Glass wool) square heat exchanger. **b** Pictorial view of complete heat transfer test rig showing varying parts along with heat exchangers

(a) The 2-D schematic view and complete insulated (Glass wool) square heat exchanger.

(b) Pictorial view of complete heat transfer test rig showing varying parts along with heat exchangers.

in Fig. [4](#page-7-0) schematic/pictorial diagrams. The nanofluid flow loop starts from the maximum 10L capacity of the fuid tank where the nanofuids were stirred throughout the experiment and a chiller was used to keep the temperature constant. The Araki-EX-70R fuid pump was used to pump the nanofuids with a maximum 88 L/min capacity and all about zero discharge heads. The pump flow was controlled by adjusting frequency at main electric control panel according to the experimental requirement. The PX154-025DI/OMEGA company pressure transducer with an accuracy of $\pm 0.075\%$ was used to measure the pressure drop during nanofuids running on the test rig. Preceding to start the experiment, the proper arrangement of the heat exchanger and other parameters is the key importance to provide a constant heat flux to outer surface of the square heat exchanger. The five K-type thermocouples are connected at numerous locations alongside the square heat exchanger and to measure the surface temperature by Wilson plot calculations [[33](#page-21-2), [34](#page-21-3)]. Additionally, the diferent thermo-physical characteristics of newly synthesized ZnO-EG@DW nanofuids based were inspected for the improved heat transfer (Nusselt numbers) studies, fuid friction, total overall pressure drop "∆*P*" and local or average heat transfer. By using the cooling law of Newton for nanofuids, both inlet–outlet temperatures, bulk temperatures, nanofuid internal temperatures, surface temperature, and the total heat transfer coefficient were calculated. Also, constant heat fux and variable fow rate conditions were followed throughout the experiment. Eight varying fow rate were trailed here in this study.

Thermal conductivity analysis for ZnO‑EG@ DW‑based nanofuids

Figure [5](#page-8-0) displays the KD-2 Pro thermal Analyzer setup of Decagon company to calculate the thermal conductivities of varying ZnO-EG@DW mass% concentrations by employing

Fig. 4 Complete schematic design of heat transfer test rig along with varying parts

the ASTM/D-5334 and IEEE- 442/1981 standards. The below shown setup of thermal analyzer tracks the transient line heat sources to read the thermal properties of nanofuids. The experiments were executed to analyze the thermal conductivity for ZnO-EG@DW nanofuids with temperature ranges starting from 20 to 45 °C. A cooling system (chiller) was used to keep the surrounding temperature maintained. Before starting the thermal conductivity experiment the needle sensor of the thermal analyzer has confrmed by calculating the thermal conductivity of the glycerin standard sample provided by the company. The intended value for the standard sample was 0.285 W m⁻¹ K⁻¹ with a precision of \pm 0.34%. For ZnO-EG@DW-based nanofluids, the experiment was conducted repeatedly and the average values were considered for each temperature range. The entire thermal

conductivity was measured using advanced KD-2 pro thermal analyzer. The six varying temperature ranges 20, 25, 30, 35, 40 and 45 °C were chosen to see the thermal conductivity variation of all nanofuids against temperature. Further, the obtained thermal conductivities calculations are directed for diferent time intervals depending upon the surrounding temperature becoming stable before each new calculation at the changed mass% of ZnO-EG@DW nanofuids for diferent ranges of temperatures. This is important to reduce the faults, which happen during the experiment by heat convection due to temperature changes of sensor needle. The ZnO-EG@DW nanofluids at changing mass% concentrations (0.1, 0.075, 0.05, and 0.025 mass%) were tested here for thermal conductivity measurements.

Fig. 5 The KD-2 pro thermal setup for thermal conductivity measurement

Data reduction

The optimal thermal conductivity for ZnO-EG@DW-based nanofuids has been analyzed by use of Maxwell model offered for thermal conductivity $[35, 36]$ $[35, 36]$ $[35, 36]$ $[35, 36]$ as set in Eq. [5](#page-8-1) and Eq. [6](#page-8-2). Where " k_{eff} " is the overall thermal conductivity, " \varnothing " is the volume fraction of the solid nanoparticles, " k_s or k_p " is the thermal conductivity of solid nanoparticles, and " k_f " is the thermal conductivity of the fuid.

$$
\frac{\text{keff}}{\text{kf}} = ks + 2kf + 2\emptyset (kf - ks)/ks + 2kf - \emptyset (kf - ks) \tag{5}
$$

$$
\frac{\text{keff}}{\text{kf}} = kp + (n - 1_{\text{kf}} - (n - 1)(kf - kp)\emptyset / kp + (n - 1)kf + (kf - kp)\emptyset
$$
\n(6)

The average and local heat transfer (h) were evaluated using Eq. [7](#page-8-3). Where "q" is known as the value of heat fux, Heat exchanger wall temperature is "Tw" and the bulk temperature is "Tb." The amount of bulk temperature was calculated by following Newton's cooling law.

$$
h = \left(\frac{q''}{Tw - Tb}\right) \tag{7}
$$

The total delivered power of closed square exchanger and its area was considered to fnd the value of overall expanded heat fux by the heater. Where "*Q*" is the power which is the product of both $V \times I$ that has been applied to the heater and similarly the "A" is a dynamical area of the heat exchanger as shown in Eq. [8.](#page-8-4)

$$
q'' = \left(\frac{Q}{A}\right) \tag{8}
$$

The square pipe (heat exchanger) area could be calculated by using Eq. [9.](#page-8-5)

$$
A = 4AL \tag{9}
$$

where "*A*" is the area of the square pipe and "*L*" is the total length of the square heat exchanger.

In the present research, the maximum power delivered to the heat exchanger surface by the heater is 600 W. To fnd the value of Nu, Eq. [6](#page-8-2) was used throughout the experiment, where "h" is known as heat transfer coefficient, " D_h " is known as the hydraulic diameter while "k" defnes the entire thermal conductivities. Furthermore, varying empirical correlations that were presented by Petukhov [[37\]](#page-21-6), and Gnielinski [\[38](#page-21-7)] have been used to calculate Nusselt numbers "Nu" of the single nanofuids. These Nusselt numbers "Nu" could be assessed as well by using Eqs. [10](#page-8-6), [12](#page-8-7), [13](#page-8-8), and [14](#page-8-9).

$$
Nu = \left(h * \frac{D_h}{K}\right) \tag{10}
$$

Equation [11](#page-8-10) was used to measure the value of Reynold numbers "Re." Different flow rates were selected and later were converted to velocities. The "*ρ*" is the overall density of the Nano-fuid, "*v*" is the value of velocities which is based on flow rates, " μ " is the kinematic and dynamic viscosities of the nanofuids, and "*D*" is the overall diameter of the heat exchanger.

$$
\text{Re} = \left(\frac{\rho v D}{\mu}\right) \tag{11}
$$

The Nusselt numbers "Nu" were calculated by using var-ying empirical correlations offered by Petukhov [\[37](#page-21-6)], Dittus-Boelter [[39\]](#page-21-8), and Gnielinski [\[38](#page-21-7)]. Where "*f*" is the friction of ZnO nanoparticles fowing inside the heat exchanger, "Pr" is showing the Prandtl numbers and "Re" is the specifc Reynold numbers.

$$
\text{Nu} = \left(\frac{\left(\frac{f}{8}\right)(\text{Re} - 1000) \text{Pr}}{1 + 12.7 \left(\frac{f}{8}\right)^{0.5} \left(\text{Pr}^{\wedge} 2/3 - 1\right)} \right)
$$
(12)

$$
\text{Nu} = \left(\frac{\left(\frac{f}{8}\right)\text{RePr}}{1.07 + 12.7\left(\frac{f}{8}\right)^{0.5}(\text{Pr}^{\wedge} 2/3 - 1)}\right) \tag{13}
$$

$$
Nu = (0.023 \text{ Re}^{0.8} \text{Pr}^{0.4})
$$
 (14)

The value of friction coefficient "f" was analyzed inside the square heat exchanger by using Eqs. [15](#page-9-0), [16](#page-9-1), and [17](#page-9-2) that was predicted by Petukhov [[37](#page-21-6)] and Blasius [[40](#page-21-9)]. Equation [12](#page-8-7) represents the overall friction loss of the square heat exchanger, whereas "∆*P*" is representing the complete pressure drop of the square shaped heat exchanger, "*L*" is the heat exchanger total length, "*D*" is the total diameter " ρ " is showing to the density and "v" denotes velocities inside the square heat exchanger.

$$
f = (0.79 \, \text{Ln} \, \text{Re} - 1.64)^{-2} \tag{15}
$$

$$
f = \frac{\Delta P}{\left(\frac{L}{D}\right)\left(\frac{\rho v^2}{2}\right)}\tag{16}
$$

$$
f = 0.3164 \, r e^{-0.25} \tag{17}
$$

Equation [18](#page-9-3) and Eq. [19](#page-9-4) were used to calculate the required pumping power of the square shaped heat exchanger. Equation [18](#page-9-3) can be mainly used when the Reynolds numbers "Re" are constant. Therefore, "*ρ*" is the density of nanofuids, "*D*" is the diameter of the square heat exchanger, "*μ*" is the viscosity of the Nano-fluids, " $\rho_{\rm bf}$ " is the base fluid density, " ρ bf" is the Nano-fluids density, " $\mu_{\rm bf}$ " is the base fluid viscosity and " μ_{nf} " is the Nano-fluids viscosity.

$$
W = 0.158 \left(\frac{4}{\pi}\right)^{1.75} \left(\frac{Lm^{2.75} \mu^{0.25}}{\rho^2 D^{4.75}}\right)
$$
 (18)

$$
\frac{\text{Wnf}}{\text{Wbf}} = \left(\frac{\rho bf}{\rho nf}\right)^2 \left(\frac{\mu nf}{\mu bf}\right)^3\tag{19}
$$

Characterizations of ZnO‑EG@DW‑based stable nanofuids

The optical behavior and energy band gaps of ZnO particles were examined by using a UV–Vis spectrum analyzer (UV 1800 Shimadzu corp 08,579). The absorbance level at all mass% of newly Sonochemically synthesized ZnO-EG@DW nanofluids as a function of energies band gaps (E_g) from pH level 8–11 has been recorded by UV–Vis spectrum analyzer. The absorbance level in % could resolute by UV–Vis spectrum analyzer and (hν) was calculated by equating wavelength λ where h $\nu = h * c/\lambda$, where "*h*" is the Planck constant (6.62 × 10⁻³⁴), λ = total wavelength 800–200 nm *c* = light speed 3×10^8 in m/s [[41](#page-21-10)]. The morphological erection of the sonochemically synthesized ZnO was calculated using XRD analysis (Bruker Advanced X-rays D8 Solutions). The XRD results obtained have matched the (JCPDS) X-ray data fles. The crystalline size of ZnO nanoparticles was achieved by calculating the expansion of the diferent XRD peaks with the desired positioning by using the Scherer formula:

$$
LC = \frac{180}{\pi} * k \cdot \frac{\lambda}{\text{Cos}} \mathcal{G} \sqrt{\text{FWHM}^2} \cdot s^2
$$
 (20)

Where π is equal to 3.142, λ is the wavelength of Cu/K_α radiations and κ is the constant of the Scherer rate (0.89), s is broadening of the instrument (0), while (FWHM) is known as the full width half/maximum of the surface plane (101), and plan angle is θ (101). The FWHM and plane angle have been directly taken out from the (EVA) software, and analyzed by XRD [[42\]](#page-21-11).

The FESEM investigation was engaged to see the morphology (Shape) and size of ZnO nanoparticles in presence of diferent elements by (EDX mapping using Zeiss-Supra 35-VP) energy dispersion X-rays spectrometer. Further, the investigation for attached functional groups with ZnO material was carried out by using FTIR analysis. This analysis was verifed by using FTIR spectrum analyzer Perkin Elmer with a range from 4000 to 400 cm⁻¹ [\[6](#page-20-4)].

UV–Vis spectrum analysis

UV–Vis spectra analysis is the most important and extensively used method for the confrmation of diferent nanoparticles. The absorbance spectrum of Sonochemically synthesized ZnO nanoparticles has been tested within the specifc range of wavelengths from 200 to 800 nm by using a UV-1800 SHIMADZU crop spectrometer with a cuvette length of 10 mm maximum, 190–110 nm minimum, and maximum wavelength capacity [\[43\]](#page-21-12). UV–Vis spectra test was executed to notice the optical conduct for Sonochemically produced ZnO. All the ZnO particles displayed a robust absorption level at 376 nm, which is dependent onto blue-lateral shifting of ZnO nanoparticles in bulk as given in Fig. [6.](#page-10-0) The UV–Vis spectrum analysis for Sonochemically synthesized ZnO nanoparticles has been carried out at 0.05 mass% concentration of nanoparticles in the base fuid, where the prominent peak at 376 nm confirms the ZnO synthesis. Figure [7](#page-10-1) shows the absorption levels at four changing (0.1, 0.075, 0.05, and 0.025 mass%) of ZnO-EG@DW type nanofuids, where it can be seen that the absorbance level is increased with an increase in mass% concentration of the solid nanoparticles into base fuid [[44,](#page-21-13) [45\]](#page-21-14).

Figure [7](#page-10-1) describes the maximum absorption level of the Sonochemically synthesized ZnO-EG@DW nanofuids at varying mass% (0.1, 0.075, 0.05, and 0.025) concentrations at normal temperature. The noticeable peak at 372 nm at varying mass% of ZnO-EG@DW reveals the confrmed preparation of ZnO particles by using probe sonication method. Consequently changed (0.1, 0.075, 0.05, and 0.025 mass%) of ZnO-EG@DW nanofuids display the diferent absorption levels due to the presence of maximum solid ZnO in the base fuid, where it can see a rise in mass% of ZnO increased the absorption level. The maximum absorption of 2.8% was observed at 0.1 mass% concentration of the ZnO-EG@DWbased nanofluids.

Fig. 6 The UV–Vis spectrum of ZnO nanoparticles

Fig. 7 UV–Vis spectrum analysis for absorbance ratio of ZnO-EG@ DW-based nanofluids mass%

XRD investigation

The XRD of sonochemically synthesized ZnO nanoparticles was conducted at Makmal Sinaran-X, University Malaya 301, Aras#1, Bangunan Utama (J-22), Jabatan Geologi by using apparatus of Malvern Analytical Model: EMPY-REAN. The white color ZnO nanoparticles were synthesized by adopting a one-pot sonochemical-assisted method where a pH of 11 was considered to produce crystalline nanoparticles. Figure [8](#page-10-2) describes the X-ray difraction spectra of sonochemically synthesized ZnO nanoparticles for nanofuids prepared with a pH value of 11. The ZnO nanoparticles cannot be synthesized very well with pH value 6 because of their low concentrations of OH− ions and H+ ions in the solution. Chand, P. et al. [[46\]](#page-21-15) describe that the ZnO nanoparticles having $pH \geq 8$ give the crystalline nature of the nanoparticles with diferent difraction angles of 2θ. The three diferent prominent peaks are considered to be the 100, 002, and 101 correspondingly to the planes. The 101 peak is to consider the preferred orientation of ZnO nanoparticles with≥8 pH values. The maximum intensity peak for ZnO nanoparticles was noticed at pH 11 due to enough presence of OH− ions to formulate the ZnO nanoparticles. All the detected peaks with pH 11 show the successful synthesis of ZnO nanoparticles and all the bases and precursors are fully decomposed and no impurity has been found during the X-ray-difraction analysis.

FTIR analysis

The one-pot facile sonochemical synthesis method was used to synthesis the ZnO nanoparticles where the pH level was adjusted at 11 to obtain the crystalline nanoparticles with specific morphology. Figure [9](#page-11-0) displays the Fourier transform infrared (FTIR) spectrum of sonochemically synthesized ZnO nanoparticles with pH 11. Fernandes D. et al. [[47\]](#page-21-16) explain that the energy bands from 3543 to 3393 cm⁻¹ correspond to the vibration mode for O and H. When the pH increased the O and H peaks became narrower due to the additional presence of O and H ions from NaOH which strongly reacts with the zinc acetate precursor solution at higher pH values. The maximum vibration and stretch mode has been observed between 1420 and 1570 cm−1. The symmetrical vibration and stretching occur between 1450 and 1330 cm^{-1} due to the presence of C and O. The prominent peaks of ZnO nanoparticles that appear between 550 and 400 cm^{-1} ranges describe that $\text{Zn}(\text{OH})_2$ are completely transformed into ZnO nanoparticles. So, the nanoparticle

Fig. 8 The X-ray difraction (XRD) spectra of Sonochemically synthesized ZnO nanoparticles

Fig. 9 The Fourier transforms infrared spectroscopy (FTIR) ZnO nanoparticles

size affects the shifting rate of peaks due to which FTIR aid FESEM images [\[48](#page-21-17)].

FESEM analysis

Figure [10](#page-11-1) displays the FESEM image of Sonochemically produced ZnO nanoparticles. The FESEM image of ZnO nanoparticles displays the homogeneous structure when the pH value raises to alkaline conditions 10 and 11. It can be seen from the FESEM image that most of the ZnO nanoparticles were spherical, where this synthesis technique gives ZnO particles less agglomeration. This low agglomeration happens due to the high centrifuge for washing of the sample at the end. The high-speed centrifuge removes the impurities like hydroxyl ions and increases the interaction between Zn and O ions [[49](#page-21-18)]. Where the OH− ions aid the nucleation process and growth of the formation of ZnO nanoparticles, and the increase in pH value up to 10 and 11 the particle size decreased [\[50](#page-21-19)]. Smaller ZnO particle size is helpful for ZnO-EG@DW-based nanofuids dispersion and can enhance the heat transfer properties.

Results and discussion

We executed the experiments in diferent stages. 1. In the frst stage, we used a single pot sonochemical synthesis technique including (Chemical reaction, sonication, drying, centrifuge, and calcination) to produce ZnO nanoparticles in bulk. 2. In the second stage, we have done the necessary characterizations to see the proper ZnO synthesis. 3. In the third stage, we prepared ZnO-EG@DW nanofuids

Fig. 10 FESEM analysis showing the ZnO morphology

by following a two-step technique. 4. In the fourth stage, all thermo-physical characteristics like dynamic/kinematic viscosities, densities and thermal conductivity, were tested. 5. In the fnal stage, our targeted properties including stability/ dispersion, thermal properties, and hydrodynamic characteristics like (Heat transfer coefficient "h," Nusselt Numbers "Nu," Friction loss, pressure drop, and required pumping power) were investigated in detail.

For the thermal and hydrodynamic characteristics, the steady heat flux and changing flow conditions were used. Eight varying fow rates were considered to see the consequence of heat transfer improvement in square shaped heat exchanger geometry. Also, their hydrodynamics efects were calculated accordingly. Finally, the outcomes were equated with previously presented varying correlations that showed an association.

Dispersion and stability of ZnO‑EG@DW‑based nanofuids

The different thermophysical properties of ZnO-EG@ DW nanofluids are extremely affected by their stability and dispersion in the EG@DW. Similarly, as stated in the literature, a maximum dispersion and stability of around 4–6 weeks have been reported for ZnO-based nanofuids, where diferent surfactants and dispersants were used to escalate the solid nanoparticles stability and longer suspension [[49–](#page-21-18)[51\]](#page-21-20). The current study emphasizes to improve the stability and dispersion of the ZnO-EG@DW-based Nano-fuids without using surfactant. Stability represents the aggregate of suspended particles at a signifcant rate. Particles in the dispersion may adhere together and form aggregates of increasing size which may settle out due to gravity. The rate of aggregation is in general determined by

the frequency of collisions and the probability of cohesion during collision. In fact, the stability of a particle in solution is determined by the sum of van der Waals attractive and electrical double layer repulsive forces which endures between particles as they approach each other due to the Brownian motion [[51](#page-21-20), [52\]](#page-21-21). At a large attractive force, the particles collide and make unstable suspension. At sufficiently high repulsion of particles, the suspensions remain in a stable state. Thus, with the dominated repulsive forces between particles nanofuids or colloids could be stable. Two types (steric, electrostatic) or repulsive forces are recognized [\[53\]](#page-21-22).

Figure [11a](#page-13-0) and b is presenting the physical observation and stability graph obtained by UV–Vis spectroscopy of Sonochemically dispersed ZnO-EG@DW-based nanofuids at varying (0.1, 0.075, 0.05, and 0.025 mass%) concentrations. No sedimentation occurred for stated nanofuids for 3 months after the preparation for all mass% concentrations. This longer dispersion is mainly accredited to a smaller size, the viscosity of the base fuid (EG@DW), and sonication time during preparation. Figure [11](#page-13-0)a reveals the reliability of ZnO nanoparticles dispersion and stability in the EG@DW mixture exposed to high Probe sonication for (0.1, 0.075, 0.05, and 0.025 mass%) of ZnO-EG@DW-based nanofuids. The sonicator setting was made according to (Total I/P voltage 220 v, Frequency 20 kHz, Pulse Amp 80, Pulse duration $3-2$ s on and off, total delivered energy 36000 J, Probe Temp 0 °C, Total Power 750 W) has been adjusted for the probe sonication of ZnO-EG@DW-based nanofuids for 2 h continuously. The pictorial images in Fig. [11](#page-13-0)b display the uniform dispersion of ZnO nanoparticles in the EG@DW mixture and look very stable until 11 weeks for all the mass% concentrations after their preparation and nanoparticles are well dispersed in EG@DW without any surfactant. Sonication time increases the dispersion of ZnO nanoparticles in base fuid due to which it becomes stable. From week 11 to onward all the sample starts from fewer sediments and high sedimentation was noticed for 0.025% sample because of lower amounts of ZnO nanoparticles. Figure [11c](#page-13-0) is expressing the comparison of ZnO-EG@DWbased nanofuids at 0.1 mass% concentration before use and after use after the day of preparation for equal time slots of (30, 60, and 90) days to check their stable characteristics for heat transfer studies. The UV–Vis analysis was conducted to check the behavior of ZnO-EG@DW nanofuids with different time passages like (on the day of preparation, after (30 days, 60 days, and fnally after 90 days). It can be seen in Fig. [11](#page-13-0)c that the absorbance % and the prominent peak at 288 nm are stable for all sonication time, which is the strong justifcation that the nanofuids didn't lose its properties even before and after the use. Therefore, the same nanofuids easily could be re-used by again sonication.

Thermophysical characteristics of ZnO‑EG@DW nanofuids

Figure [12](#page-14-0)a displays the thermal conductivity enrichment for the ZnO-EG@DW-based nanofuids at changed (0.1, 0.075, 0.05, 0.025 mass%) concentrations at numerous temperatures ranges from 20 to 45 °C. The thermal conductivity of all the mass% of ZnO-EG@DW nanofuids exhibited growing trends with an increase in temperature value, which has already an agreement in earlier researches [[54,](#page-21-23) [55](#page-21-24)], whereas the EG@DW displays declining trends with an increase in temperature. The same declining behavior has already been reported in earlier experiments steered by Asadi et al. [[55,](#page-21-24) [56](#page-21-25)] and Aberomand et al. [\[57\]](#page-21-26). The mounting inclination has been detected at all mass% of ZnO-EG@DW nanofuids. An linear surge in ZnO particles collisions and their Brownian motions could be the obvious key reason for enhancing the overall thermal conductivity of ZnO-EG@DW nanofuids with a temperature change. The thermal behavior of the ZnO-EG@DW nanofluids with different mass% concentrations is shown in Fig. [12a](#page-14-0). where the thermal conductivities of all mass% of ZnO-EG@DW nanofuids present a growing trend as the mass% concentration increases. This growing trend was noticed for all the temperature values. Also, the graph shows the addition of ZnO in EG@DW led to an improvement in the thermal conductivity of the base fuid, which was previously reported by several researchers. So the use of these nanofluids/Nano-coolants with solid mass% concentrations could be commended for diferent cooling applications.

The dynamic viscosity of EG@DW and ZnO-EG@DW nanofuids moderates with the rise in temperature of the fuid, and the viscosity outcomes for all (0.1, 0.075, 0.05, and 0.025) mass% of ZnO-EG@DW nanofuids and EG@ DW as well against diferent mass% of ZnO nanoparticles are given in Fig. [12b](#page-14-0). The key reason for this increasing behavior of viscosity value is due to an increase in mass% of particles in base fuid (EG@DW) while the viscosity ratio for all mass% exhibits the declining trend against the increase in temperature value. This declining behavior of the viscosity of ZnO-EG@DW type nanofuids is due to their weak internal molecular interaction force [[58,](#page-21-27) [59\]](#page-21-28). The detailed dynamic viscosities at all concentrations of ZnO-EG@DW type nanofuids and EG@DW base are illustrated in Tabl[e1](#page-14-1), where the observed minimum and maximum temperature range was from 20 to 45 °C. Similarly, Fig. [12c](#page-14-0) shows the same declining trend for the kinematic viscosity of EG@DW and varying concentrations of ZnO-EG@DW type nanofuids against the temperature values. While the increase in viscosities value is due to the increase in mass% of ZnO solid nanoparticles in the base fuid EG@DW. The 0.1 concentration of the ZnO-EG@DW showed highly dynamic and kinematic viscosities due to the existence of

Fig. 11 a ZnO sedimentation efecting to the dispersion and stability of the nanofuids, **b** Dispersion and stability behavior for ZnO-EG@DW nanofuids, **c** UV–Vis showing the stable characteristics of ZnO-EG@DW-based Nano-fuids after and before use on test rig

the maximum concentration of ZnO in EG@DW. The behavior of all mass% concentrations of the ZnO-EG@DW-based nanofuids and base fuid EG@DW against the temperature variations are given in Table. [2.](#page-14-2)

Figure [12d](#page-14-0) shows the density behavior of EG@DW and varying concentrations of ZnO-EG@DW-based nanofuids. After adding the ZnO nanoparticles to EG@DW, the overall density of the nanofluids has changed according to mass% of nanoparticles. Also, due to an increase in the temperature value the density of all fuids starts to decrease slightly. When the temperature was low, all fuids showed maximum density, the maximum density was noticed for

Fig. 12 a Variations in thermal conductivities for at varying ZnO-EG@DW mass%., **b** Dynamic viscosity variations against varying temperatures at varying mass% of the ZnO-EG@DW nanofuids, **c** Kinematic viscosity variations against varying temperatures for

changed concentrations of the ZnO-EG@DW nanofuids, and **d** Density behavior against the varying temperatures of the EG@DW mixture and all concentrations of ZnO-EG@DW-based nanofuids

Table 1 Dynamic viscosity profle at varying concentrations of ZnO-EG@DW-based nanofuids and EG@DW mixture

Temp	0.025% mass%	0.05% mass%	0.075% mass%	0.11 mass%	EG@DW 50:50
20	3.8066	3.6669	3.7345	4.2306	4.4475
25	3.2956	3.1650	3.2054	3.6158	3.9572
30	2.8319	2.7125	2.758	3.1027	3.2682
35	2.4524	2.3668	2.4038	2.6871	2.8195
40	2.1311	2.0572	2.1034	2.3398	2.4516
45	1.9038	1.8447	1.8763	2.0841	2.1699

Table 2 Kinematic viscosity profle at all concentrations of ZnO-EG@DW-based nanofuids and EG@DW mixture

higher concentration of the ZnO-EG@DW-based nanofuids. Though the EG@DW and the lowest concentration of ZnO-EG@DW-based nanofuids showed lower density as compared to other fuids, these density variations are credited to the occurrence of the ZnO. The detailed density behavior of all mass% concentrations of the ZnO-EG@DW-based nanofuids and base fuid EG@DW for diferent values of temperature is shown in Tabl[e3](#page-15-0).

Hydrodynamic characteristics of all concentrations of ZnO‑EG@DW‑based nanofuids inside a square heat exchanger

To study the total pressure drop inside the square heat exchanger, we intended the friction factor for EG@DW mixture and all concentrations of ZnO-EG@DW-based nanofuids, respectively. In the current study, the results that

Table 3 Densities of all concentrations of ZnO-EG@DW-based nanofuids and EG@DW mixture

Temp	0.025/ mass%	0.05/ mass%	0.075/ mass%	0.1/ mass%	EG@DW 50:50
20	1.0656	1.0640	1.0649	1.0706	1.072
25	1.0626	1.0614	1.0622	1.0678	1.0694
30	1.0597	1.0585	1.0593	1.0647	1.0662
35	1.0569	1.0555	1.0564	1.0617	1.0629
40	1.0539	1.0523	1.0534	1.0585	1.0597
45	1.0508	1.0494	1.0503	1.0555	1.0563

arise from the calculation of the friction factor in Fig. [13](#page-15-1)a, where the outcomes show a falling trend with a growth in Reynolds numbers, all concentrations showed a decrease in friction factor due to an increase in velocity which afects the Reynolds increase. The friction was evaluated considering the overall pressure drop "Δ*P*," length of the heat exchanger "*L*," hydraulic diameter of the heat exchanger "*D*," viscosity "*V*," and density "*ρ*" of the running fuid. To calculate the changes in pressure drop across square heat exchanger, calculations have been done to fnd the friction loss for base fuid in the square heat exchanger by applying Eq. [16](#page-9-1). The assessed results were compared with Blasius's [[40](#page-21-9)] and Petukhov model [[60\]](#page-21-29) Models where the total difference was

<

(21)

5% when it has compared with Blasius and Petukhov correlations results. These correlations showed that the square shaped heat exchanger is reliable to notice the total pressure drop across the test section for changed Reynolds values stated in the research. The entire pressure drop across the square shaped heat exchanger of ZnO-EG@DW-based nanofuids at diferent concentrations according to Reynolds (Re) is given in Fig. [13](#page-15-1)b. During the experiment, it has been observed that there is a slight surge in pressure drop of ZnO-EG@DW nanofluids as in comparison to EG@DW. This increase in pressure drop was reported due to the existence of ZnO in EG@DW, which may increase the friction loss resulting in an increase in the overall pressure drop.

Fig. 13 a Comparison of friction loss of ZnO-EG@DW nanofuids and EG@DW, **b** Comparison of experimental and model data between pressure drop and Reynolds numbers

Average heat transfer (h) and average Nusselt (Nu) numbers analysis of all concentrations of ZnO‑EG@ DW‑based nanofuids

Detailed investigations describe the positive effects varying mass% concentrations(0.1, 0.075, 0.05, and 0.025) of ZnO-EG@DW-based nanofuids on average heat transfer/ Nusselt numbers for specifed Reynolds ranging from 4550 to 20,360. The changing fow value and persistent heat fux conditions were chosen throughout the experiment. The experimental process has been conducted to assess the average heat transfer (h) and their special efects on to average Nusselt (Nu) numbers in the square shaped heat exchanger of ZnO-EG@DW-based nanofuids. The calibration and necessary setting of the heat transfer experimental setup was maintained on a priority basis before running the experiment. To realize the overall temperature of the heat exchanger surface and its heat conduction at wall of the heat exchanger, as well as the heat convection allowed by ZnO-EG@DW-based nanofuids were studied here. Wilson plot [\[33\]](#page-21-2) procedure was trailed here, by figuring the resistance at the various locations of the square heat exchanger, also heat transfer in the square heat exchanger. We measured the thermo-physical characteristics of distilled water and varying mass% concentrations of ZnO-EG@DW nanofluids by seeing numerous factors like friction, pumping power, heat transfer (h), pressure drop and Nusselt (Nu) numbers, etc. Likewise, the average heat transfer was recorded for ZnO-EG@DW-based nanofuids running in a square heat exchanger. Figure [14a](#page-17-0) is describing the typical heat transfer for EG@DW mixture and all concentrations of ZnO-EG@ DW-based nanofuids. Further, the growing trend shows an improvement in the heat transfer of at varying concentrations of ZnO-EG@DW nanofuids, although the base fuid (EG@ DW) bounces lesser heat transfer coefficient (h) which is credited to the nonexistence of ZnO particles in the EG@ DW mixture. Also, it can be seen, the highest enhancement in average value of heat transfer was perceived for at 0.1 concentration of ZnO-EG@DW-based nanofuids which are nearly about to 600–1800 W m⁻¹ K⁻¹, similarly rest all mass% concentrations showed improved heat transfer outcomes compared to EG@DW mixture but less enhanced than 0.1 mass% concentration. The outcomes accomplish the full existence of ZnO particles in base will surge to the heat transfer as well as Reynolds increased.

Figure [14b](#page-17-0) is presenting a comparison between the current study on heat transfer improvement and the past studies, where diferent metal oxides were used in diferent base fuids. The Reynold range was considered within 400 to 25,000. It can be seen in Fig. [14](#page-17-0)b the use of ZnO in EG@DW mixture is leading to improved heat transfer. This improvement is mainly due to good stability, High flow rate,

and higher mass% concentration of the ZnO-EG@DW-based nanofluids.

Based on heat transfer coefficient (h) measurements it's easier to evaluate average Nusselt (Nu) development at all (0.1, 0.075, 0.05, and 0.025 mass%) concentrations of ZnO-EG@DW-based nanofuids. Similarly, Fig. [14](#page-17-0)c is presenting the increase in average Nusselt (Nu) at all concentrations of ZnO-EG@DW nanofluids with growth Reynolds values varying from 4550 to 20,367. Average Nusselt enhancement for EG@DW is comparatively less than ZnO-EG@DW-based nanofuids, where 0.1 mass% gives the highest improvement against the Reynolds increases. The enhancement of Nusselt numbers relies on the heat transfer coefficient, which mainly increases due to the occurrence of ZnO. After adding the ZnO nanoparticles in the EG@DW the improvement in the Nusselt (Nu) numbers was observed, which has been attributed to the increase in ZnO particles mass%. Also, the enrichment in Nusselt numbers is credited to the increase in Reynolds values, as the Reynolds increasing the average Nusselt numbers were increased accordingly.

The outcomes arising from experimental data were compared with some empirical correlations data, like Petukhov, Gnielinski, and Dittus to fnd the improvement in average Nusselt numbers of the ZnO-EG@DW-based nanofuids. These empirical correlations are mentioned in Eq. [6](#page-8-2) and [7](#page-8-3) as well, also these correlations can be used for only the turbulent type flow of the nanofluids. Figure [14](#page-17-0)d shows the comparison graph of experimental data with correlations data, where the mounting trends in average Nusselt numbers with variation in Reynolds. Also, the experimental results show positive results compared with diferent model empirical correlations. Therefore, the presented experimental study and setup can be used to study the heat transfer characteristics of the ZnO-EG@DW nanofuids in square heat exchangers.

Local heat transfer coefficient (h)/local Nusselt (Nu) numbers measurement of ZnO‑EG@DW‑based well stable nanofuids in a square shaped heat exchanger

For the heat transfer measurements at varying points of the square shape heat exchanger (local heat transfer, h) the heat exchanger was calibrated with five K-type sensitive thermocouples on its outer surface. These thermocouples were placed apart from each other at a distance of 0.2 m along the square heat exchanger to measure the local heat transfer values at every point. The investigation was conducted to quantify the effects of $ZnO-EG@DW$ nanofluids with $(0.1, 1.1)$ 0.075, 0.05, and 0.025) mass% on local heat transfer (h) in the square test section individually. The local heat transfer measurement for all mass% concentrations on fve diferent points of the square heat exchanger has been conducted at

Fig. 14 a Average heat transfer development in ZnO-EG@DW nanofuids with changed mass%., **b** Comparison of heat transfer improvement in % of the current study with past, **c** Growth in Nusselt num-

bers vs Reynolds numbers of the ZnO-EG@DW-based nanofuids, and **d** Comparison between experimental and varying model data results

constant heat fux and changing fow rate settings throughout the experiment. Figure [15a](#page-18-0)–d indicates the local conduct of the heat transfer (h) for all mass% of ZnO-EG@DW nanofuids at various points of the square test section. All the mass% concentrations were treated individually in a heat exchanger with numerous flow rates or specified range 4550-20,367 of Reynolds were also exposed. Through the experiment, the outcome arises that at the 0.025 mass% of ZnO-EG@ DW-based nanofuids exhibited low local heat transfer (h) which is 550–1500 W m⁻² K⁻¹ for Reynolds increasing from 4550 to 20,367. In the case of 0.05 mass% of ZnO-EG@ DW nanofuids, the local heat transfer (h) was observed at 550 to 1750 W m⁻² K⁻¹ at identical Reynolds numbers. The 0.075 mass% of ZnO-EG@DW nanofuids show the heat transfer (h) 550–2170 W m⁻² K⁻¹ against the same Reynolds numbers. Finally, the maximum improvement in local heat transfer was observed at about 550 to 2830 W m^{-2} K^{-1} for the 0.1 mass% of ZnO-EG@DW nanofluids against the same Reynolds values ranging from 4550 to 20,367. At all mass% of the ZnO-EG@DW-based nanofuids exhibited a supreme heat transfer (h) value at the frst point of the heat exchanger which is closest to the inlet point of the test section. As the distance increases from the inlet point, the heat transfer decreases according to diferent fow rates and their respected Reynolds. The supposition from the stated research describes that the presence of ZnO nanoparticles in EG@DW 50:50 and their diferent mass% addition up to 0.1 mass% exposed to positive growth in local heat transfer coefficient (h) in square heat exchangers.

Fig. 15 a-d Variations in local heat transfer coefficient (h) augmentation for all mass% of ZnO-EG@DW-based nanofluids

According to local heat transfer (h) measurements of changing mass% of ZnO-EG@DW-based nanofuids, it is quite easy to calculate the variations in local Nusselt (Nu) on diferent points alongside the square heat exchanger. Based on heat transfer results, the local Nusselt numbers improvement was assessed for changing mass% of ZnO-EG@DWbased nanofuids accordingly. Figure [16a](#page-19-0)-d demonstrates the diferent trends of Nusselt (Nu) for specifed Reynolds ranging from 4550 to 20,367 at all mass% like (0.1, 0.075, 0.05, and 0.025) of the ZnO-EG@DW-based nanofuids. Through the experiment, it was observed local Nusselt (Nu) improves with an increase in Reynolds with the same hydraulic diameter of a square heat exchanger. These improved results are attributed to an increase in mass% of ZnO in EG@DW, and surge in Reynolds. From Fig. [16](#page-19-0)a-d, it is observed that all mass% of ZnO-EG@DW nanofuids; the local Nusselt (Nu) are grown with the rise in mass% and Reynolds (Re). While Local Nusselt numbers for all mass% decrease as the distance increases from the input point of the square shape heat exchanger. Also, at the lowest mass% of ZnO-EG@DW nanofuids, the local Nusselt (Nu) was recorded 8–13 consistent with the heat (h) at Reynolds continued from 4550 to 20,367. At the 0.05 mass% of ZnO-EG@DW-based nanofuids showing a rising trend of local Nusselt (Nu) from 6 to 15 at the similar Reynolds (Re). For 0.075 mass% of ZnO-EG@ DW-based nanofuids, the local Nusselt (Nu) was perceived from 6 to 18 at the similar Reynolds (Re). Moreover, the 0.1 mass% of ZnO-EG@DW nanofuids showed a surprising enhancement in local Nusselt (Nu) about 6–25 against the specifed Reynolds (Re) numbers mentioned earlier. This decreasing behavior of local Nusselt (Nu) at lowest mass% of ZnO-EG@DW-based nanofuids for all Reynolds (Re) is

Fig. 16 a–d Local Nusselt (Nu) numbers calculation at varying mass% of ZnO-EG@DW nanofuids

attributed to the fewer amount and agglomeration of ZnO for low Reynolds. While, high Reynolds numbers stop the agglomeration of ZnO in EG@DW, which enhances the ZnO in EG@DW due to proper mixing. Finally, it has been perceived that the improvement in Nusselt numbers (Nu) is credited to a surge in heat transfer coefficient, which is ascribed to the maximum presence of ZnO nanoparticles in the base fuid.

Conclusions

The thermophysical characteristics, hydrodynamic and heat transfer (h) improvement of the ZnO-EG@DW-based nanofuids at varying mass% concentrations fowing in a square heat exchanger with constant heat fux boundaries, variable Reynolds conditions in turbulent fow regimes were studied in the current investigation. Based on experimental results the following important fndings have been concluded.

- The distinctive single pot sonochemical synthesis technique for the synthesis of ZnO nanoparticles was followed by a standard high probe sonication procedure. The spherical shape of ZnO nanoparticles was produced which was confrmed by FESEM analysis.
- The ZnO-EG@DW nanofluids were prepared using a 2-step nanofuids preparation technique at four diferent mass% of 0.1, 0.075, 0.05, and 0.025, All the characterization tests of ZnO nanoparticles were carried out using UV–visible spectrum XRD, FTIR, and FESEM to confrm a precise synthesis.
- The experimentation was targeted to measure the thermophysical and heat transfer properties of ZnO-EG@ DW-based nanofuids at varying 0.1, 0.075, 0.05, and 0.025 mass% concentrations against Reynolds (Re) numbers varied from 4550 to 20,367.
- All the mass% concentrations of ZnO-EG@DW-based nanofuids were tested for thermal conductivity measurement by using an advanced thermal analyzer (KD-2 pro). An improvement in thermal conductivity was noticed when the mass% of ZnO increases in EG@DW. The substantial augmentation in thermal conductivity was noted at 0.1 mass% concentration of the nanofuids which is greater than to the base fuid, while all other mass% shows thermal conductivity higher than base fuid and less than 0.1 mass%.
- Based on constant heat flux and variable Reynold (Re) numbers the significant improvement of heat transfer (h) was perceived at all mass% concentrations of the ZnO-EG@DW-based stable nanofuids. The highest development in heat transfer coefficient (h) was observed at 0.1 mass% concentration that is about 550– 2830 W m⁻² K⁻¹ that is 63% and higher than the base fuid while, 0.25 mass% of the Nano-fuids demonstrated less improvement as compared to others.
- The ZnO-EG@DW-based nanofluids showed enhanced Nusselt (Nu) from 6 to 27 at the 0.1 mass% concentration

at the maximum value of Reynold (Re) numbers 20367. This improvement is mainly credited to the occurrence of the maximum concentration of solid ZnO nanoparticles in the base fluid.

• Thus, the improved thermal conductivity and overall heat transfer properties of ZnO-EG@DW-based nanofuids could be a good choice for heat exchangers for household and industrial applications.

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Declarations

Conflict of interest The author does not have any kind of confict.

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