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Omer A. Alawi
Haslinda Mohamed Kamar
Abdul Rahman Mallah
Hussein A. Mohammed
*Edith Cowan University*
Mohd Aizad Sazrul Sabrudin

*See next page for additional authors*

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Graphene Nanoplatelets Suspended in Different Basefluids Based Solar Collector: An Experimental and Analytical Study

Omer A. Alawi 1,*, Haslinda Mohamed Kamar 1,*, Abdul Rahman Mallah 2,©, Hussein A. Mohammed 3,*,©, Mohd Aizad Szazrul Sabrudin 1,4, Omar A. Hussein 5, Salim Newaz Kazi 2,© and Gholamhassan Najafi 6

1 Department of Thermofluids, School of Mechanical Engineering, Universiti Teknologi Malaysia, 81310 UTM Skudai, Johor Bahru 81310, Malaysia; masazul2@graduate.utm.my
2 Department of Mechanical Engineering, University of Malaya, Kuala Lumpur 50603, Malaysia; a.r.mallah@siswa.um.edu.my (A.R.M.); salimnewaz@um.edu.my (S.N.K.)
3 School of Engineering, Edith Cowan University, 270 Joondupal Drive, Joondalup, WA 6027, Australia
4 PROTON Holdings Sdn Bhd, HICOM Industrial Estate, Batu 3, P.O. Box 7100, Shah Alam 40918, Malaysia
5 Department of Mechanical Engineering, College of Engineering-Alsharkat, Tikrit University, Tikrit 34001, Iraq; omar-assi81@tu.edu.iq
6 Department of Biosystem Engineering, Tarbiat Modares University, Tehran 14115-111, Iran; g.najafi@modares.ac.ir
* Correspondence: omeralawi@utm.my (O.A.A.); haslinda@utm.my or haslinda@mail.fkm.utm.my (H.M.K.); hussein.mohammed@ecu.edu.au or Hussein.dash@yahoo.com (H.A.M.)

Abstract: A flat plate solar collector (FPSC) was analytically studied, with functionalized graphene nanoplatelets (f-GNPs) as its working fluid. Four samples (wt % nanofluids) were prepared in different base fluids such as ethylene glycol (EG), distilled water (DW):EG (70:30), and DW:EG (50:50). Experimental results (via DW) were used to verify the effectiveness of the analytical model. Some of the operating conditions were taken into account in this research, including temperatures, power, and mass flow rates. Experimental techniques were used to elucidate the modified nanofluids’ physicochemical properties, such as its particle sizes, stability, and morphology, involving electron microscopes (EMs), UV–VIS, and X-ray techniques. Differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA) were applied to test the thermal analysis. The findings confirmed that the use of f-GNPs nanofluids enhanced the performance of the FPSC relative to the use of base fluids for all testing conditions. The maximum enhancement of the collector’s effectiveness at a mass flow rate of 1.5 kg min\(^{-1}\) and a weight concentration of 0.1 wt %, increased to 12.69%, 12.60%, and 12.62% in the case of EG, DW:EG (70:30), and DW:EG (50:50), respectively. The results also confirmed an improvement in both the heat gain (\(F_R(\tau \alpha)\)) and heat loss (\(F_R U_L\)) coefficients for the f-GNPs nanofluid.

Keywords: flat plate solar collector (FPSC); GNPs; characterization and stability; collector efficiency

1. Introduction

Solar collectors are devices/mediums that convert incident solar energy into thermal energy via working fluids. There are many solar collectors, and the flat plate solar collector (FPSC) represents its most basic form, which carries out the energy conversion process via an absorbing plate [1–4]. An absorbing plate is made of black-colored materials or coated surfaces, which is to increase the absorption of incident solar rays [5]. The absorbed solar rays are then converted to heat via heat transfer fluids (HTFs), channeled to the absorbing plate via a network of collector pipes. Direct absorption solar collectors (DASCs) represent another type of solar collector where the radiation is absorbed via the heat carried by fluids, instead of the heat collected on the plate’s surface [6,7].

The physicochemical intricacies of a standard HTF are admittedly inferior to solid nanoparticles (NPs), which means that rays are translated into heat at lower heat transfer
rates when using the former [8,9]. The efficiency of FPSCs can be enhanced [10,11] by eliminating HTFs from the system and using nanofluids which have superior thermal property.

The literature has several studies involving the use of carbon-based nanofluids as HTFs within FPSCs. Yousefi et al. [12–14] used a 2 m² FPSC to determine the impacts of Al₂O₃–distilled water (DW), multiwalled carbon nanotubes (MWCNTs)–DW, and pH variation with MWCNTs–DW in the context of energy efficiency. The nanofluids flowed at 1–3 L min⁻¹. The greatest improvement in the FPSC’s thermal efficiency was 28.3% at 0.2 wt % Al₂O₃–DW, with a surfactant [12]. The values of heat gain and loss coefficients increased by 65.51% and 45.84%, respectively, relative to the DW for the 0.4 wt % MWCNTs [13]. The use of 0.2 wt % MWCNTs at a pH value of 3.5 resulted in higher FPSC thermal performance relative to the use of 0.2 wt % MWCNTs at a pH value of 6.5 [14].

The effects of using 0.2 wt % and 0.4 wt % MWCNTs-DW nanofluids on decreasing the dimensions of an FPSC were reported in [15], where the size of the FPSC decreased by ~37% with the use of MWCNTs-DW nanofluid compared to that using distilled water as a base fluid. Said et al. [16] examined different nanofluids (SWCNTs, SiO₂, TiO₂, and Al₂O₃) to improve the effectiveness of FPSCs. SWCNTs-DW performed better than metal oxide nanofluids. When using SWCNTs–DW, the entropy generation and heat transfer coefficient decreased by ~4.34% and increased by ~15.33%, respectively, compared to those by using distilled water. Said et al. [17] increased the energy and exergy efficiencies of the FPSC–SWCNTs by ~95.12% and ~26.25%, respectively, which were higher than the values reported when using water (~42.07% and ~8.77%, respectively). The energy efficiency of an FPSC improved by ~33% via the addition of 0.005 wt % graphene nanoplatelets (GNPs) to DI water as HTFs [18]. Ahmadi et al. [19] reported enhanced FPSC efficiency by ~12.19% and ~18.87% at 0.01 wt % and 0.02 wt % GNPs, respectively. Vincely and Natrajan [20] outlined an improvement in the GO-DW nanofluids’ thermal efficiency at a rate of 7.3% relative to that of the base fluid at a weight percent of 0.02 and a mass flow rate of 0.0167 kg s⁻¹. Enhancements in the performance of the FPSC were 23.47%, 16.97%, 12.64%, 8.28%, 5.09%, and 4.08%, corresponding to the MWCNTs-DW, Gr-DW, CuO-DW, Al₂O₃-DW, TiO₂-DW, and SiO₂-DW, respectively [21]. Akram et al. [22] determined the improvement in the thermal performance of an FPSC to be 78% at a weight percent of 0.1 wt % CGNPs and a mass flow rate of 0.0260 kg s⁻¹. Alawi et al. [23] analyzed the thermo-performance of FPSCs with GNPs as HTFs and reported that relative to the DW, the efficiency of the FPSCs increased by 10.7%, 11.1%, and 13.3% for the PEG-GNPs nanofluid in the case of multiple mass flow rates. In the case of metal oxides, GNPs, and SWCNTs nanofluids, the efficiency was directly proportional to the fraction of NPs, as the improved thermal properties mitigated hydrodynamic losses caused by high viscosity laminar flows [24]. The efficiency of the FPSC was directly proportional to both the weight concentration and specific surface area (SSA), up to 10.53% relative to that of the increase brought about by water [25]. Experimental work using hybrid MWCNT + Fe₃O₄/water nanofluids improved the absorber by 28.09% at 0.3 vol. % at 13:00 h daytime and a Reynolds number of 1413 relative to those using base fluid [26].

Studies on suspending GNPs in distilled water and ethylene glycol lack the primary literature. Therefore, this paper aims twofold with an experimental part and an analytical part. Different tools were used in the experimental section, such as differential scanning calorimetry (DSC), thermogravimetric analysis (TGA), UV–VIS, zeta potential, particle size, HR-TEM, SEM, and energy-dispersive X-ray spectroscopy (EDX) to examine the functionalized graphene nanoplatelets. The analytical section included a variety of operating conditions on three base fluids: the weight percentages of functionalized graphene nanoplatelets (f-GNPs), mass flow rates, input temperatures, and input rates (of heat). A deeper understanding of these parameters’ effects would help significantly enhance the operational efficiency of FPSCs.
2. Methodology

This work is divided into two distinct phases: the experimental phase comprising of preparing f-GNPs nanofluids, the determination of the density and specific heat capacity of the NPs, the evaluation of the long-term stability of the nanofluids using multiple characterization techniques such as UV–Vis, zeta potential, NP analysis, electron microscopy (SEM and HR-TEM), and EDX analysis. This work’s theoretical phase consisted of mathematical modeling via MATLAB for quantifying the thermal performance of base fluids infused with f-GNPs.

Firstly, the theoretical model was verified using experiments with DW [23] and an average error of ±3.53% was obtained (see Figure 1). The analytical approach (theoretical analyses) involved utilizing different base fluids containing f-GNPs for comparative analysis. Three base fluids were used in the mathematical model, which were ethylene glycol (EG), DW:EG (70:30), and DW:EG (50:50).

Figure 1. Validation of the present theoretical model with the data available in the literature [23].

2.1. Preparation Method and Characterization

The image of the covalent-treated graphene nanoplatelets (f-GNPs) is shown in Figure 2. A high-power ultrasonication probe (Sonics Vibra Cell, Ningbo Kesheng Ultrasonic Equipment Co., Ltd., Ningbo, China) having a 20 kHz frequency power supply, a 1200 W output power, and bath ultrasonication (Powersonic, Digital Ultrasonic, UB410) was used for the preparation of nanofluids. In addition, an industrial hot plate (HTS-1003 Hotplate Stirrer, LMS) was essentially needed for the synthesis process. The light absorbance of a suspension by UV–VIS spectroscopy represented a quantitative characterization of colloidal stability. The UV–VIS used in this work was a Shimadzu UV-spectrometer, operating within 190–3300 nm (UV-750, Lambda PerkinElmer, Waltham, MA, USA). The thermal properties were tested with a thermogravimetric analyzer in the range of 30 °C to 900 °C using PerkinElmer (TGA 4000, Waltham, MA, USA). Litesizer 500-Anton Paar was used for zeta potential and particle size determination following the principle of electrophoretic light scattering (ELS). The nanomaterials’ surface structure and morphology were determined using the Tescan Analytics (SEM; VEGA3) and high-resolution transmission electron microscopy system (HR-TEM; JEM-2100F), respectively.
2.2. Density and Specific Heat Capacity Measurements

The densities and specific heats of the samples were measured and detailed as below:

(i). Solid NPs: DM40-density meter (Mettler Toledo, Columbus, OH, USA) to determine densities and Linseis’ differential scanning calorimeter (1000-/C) to determine specific heats with a temperature range of \(-50\,^\circ\text{C}\) to \(550\,^\circ\text{C}\);

(ii). Base fluids: densities and specific heats were obtained from the National Institute of Standards and Technology (NIST);

(iii). f-GNPs nanofluids: densities and specific heats were determined using the equations reported by Pak and Cho [27].

2.3. Theoretical Approach and MATLAB

The model constructed for determining the energy efficiency of the FPSC infused with nanofluids used the Hottel–Whillier principles reported by Duffie and Beckman [5], which was also slightly modified to align it with the objectives of this work (refer to Table 1). Some of the assumptions made when constructing the model were that it was designed to elucidate the performance of the FPSCs while keeping their fundamental values intact. The model was also designed to simulate the nanofluid-based FPSC in MATLAB, as outlined in Figure 3. However, in the mathematical/analytical model, only one of the tubes was assumed to experience a uniform fluid flow via all of the riser pipes within the parallel channel system.

Table 1. Specifications of the solar collector.

| Specifications                        | Dimension |
|---------------------------------------|-----------|
| Collector occupied area               | 0.6810 m² |
| Absorber area                         | 0.4645 m² |
| Header tube outer diameter            | 22.2 mm   |
| Header tube inner diameter            | 20.9 mm   |
| Riser tube outer diameter             | 12.7 mm   |
| Riser tube inner diameter             | 11.6 mm   |
| Transmittance–absorptance product     | 0.8772    |
| Tilt angle                            | 30°       |
2.4. Data Processing

The total collector heat loss encompasses the top heat loss via a glass cover and a back as well as the edge heat loss via the back and edge insulations. The losses were measured at a similar mean plate temperature \( T_{\text{pm}} \). The total heat loss from the collector was written as follows \([5,28]\):

\[
Q_{\text{loss}} = U_L A_c (T_{\text{pm}} - T_a), \quad (1)
\]

\[
Q_{\text{loss}} = Q_t + Q_b + Q_e, \quad (2)
\]

where \( U_L \) is the total loss coefficient of the collector, \( A_c \) is the area of the collector, and \( (T_{\text{pm}} - T_a) \) is the temperature gradient between the mean plate and its surrounding atmosphere and the subscripts \( t, b, \) and \( e \) represent the top, back, and edge, respectively.
The following empirical equation can be used for manual and analytical calculations to determine the top collector loss coefficient (\(U_t\)), [5]:

\[
U_t = \left| \frac{N}{T_{pm} - T_a} \right|^{-1} + \frac{1}{h_w} + \frac{\sigma (T_{pm} - T_a) \left( 1 + \frac{T_{pm}^2 + T_a^2}{2} \right)}{\varepsilon_p + 0.00911N\varepsilon_{gp} + 2N + f + 1 + 0.133\varepsilon_{ap}} - N, \tag{3}
\]

where \(ff\) is written as \((1 - 0.089h_{wind} + 0.1166h_{wind}^2\varepsilon_p) (1 + 0.078661N)\), \(CC\) is described as \(520 \times (1 - 0.000051\phi^2)\), \(ee\) is written as \(0.430 (1 - 100/T_{pm})\), and \(N, \sigma, \varepsilon_g, \varepsilon_{ap}, \) and \(h_w\) are the number of glasses, the Stefan–Boltzmann constant, the emittance of glass, the emittance of the absorber plate, and the wind-heat transfer coefficient, respectively.

The following equations can be used to determine the back and edge heat losses [5,28]:

\[
Q_b = k_b L_b A_b (T_{pm} - T_a), \tag{4}
\]

\[
Q_e = k_e L_e A_e (T_{pm} - T_a), \tag{5}
\]

where \(k_b, L_b,\) and \(A_b\) are the thermal conductivity, thickness, and back insulation area, respectively, while \(k_e, L_e,\) and \(A_e\) are the thermal conductivity, thickness, and area of edge insulation, respectively.

The following equation can be used to determine the useful energy (\(Q_u\)) [29–31]:

\[
Q_u = \dot{m}C_p(T_o - T_i), \tag{6}
\]

where \(C_p\) is the specific heat capacity of the HTFs.

The specific heat and density of the nanofluid can be determined using Equations (7) and (8) [27,32]:

\[
C_{p_{nf}} = \frac{(1 - \phi)\rho_{bf}C_{p_{bf}} + \phi\rho_{np}C_{p_{np}}}{(1 - \phi)\rho_{bf} + \phi\rho_{np}}, \tag{7}
\]

\[
\rho_{nf} = (1 - \phi)\rho_{bf} + \phi\rho_{np}, \tag{8}
\]

where the subscripts \((nf, bf, and np)\) represent the nanofluid, base fluid, and NPs, respectively.

Another indication for the rate of useful energy gained by the difference between the absorbed power and heat losses of the FPSC is as following [29–31]:

\[
Q_u = F_R A_c (G_T (\tau_a) - U_L (T_i - T_a)), \tag{9}
\]

where \(F_R\) is the heat removal factor, \(G_T\) is the global solar radiation, \(\tau_a\) is the product of absorptance and transmittance, \((T_i - T_a)\) is the difference between the input/ambient temperatures, and \(\eta\) is the thermal efficiency of the collector, generally referred to as the Hottel–Whillier–Bliss equation, which can be written as [29–31]:

\[
\eta = \frac{Q_u}{A_c G_T} = \frac{\dot{m}C_p(T_o - T_i)}{A_c G_T}, \tag{10}
\]

\[
\eta = F_R (\tau_a) - F_R U_L \frac{T_i - T_a}{G_T}. \tag{11}
\]

3. Results and Discussion

3.1. Thermal Analysis

The TGA and DSC analyses of the f-GNPs are shown in Figure 4. At the range of 0–100 °C, the initial weight loss reflected in the f-GNPs was individually 11% in virtue
of adsorbed moisture. The second slight weight loss for the f-GNPs was reported by 16% from 100 to 500 °C, which is attributable to the occurrence of –OH and C=O groups. In the range of 500–800 °C, the f-GNPs suffered another slight weight loss of nearly 24%, which is ascribable with the breakdown of the graphitic carbon structures in the air.

![Graph showing thermal properties of functionalized graphene nanoplatelets (f-GNPs)](image)

**Figure 4.** Thermal properties of the functionalized graphene nanoplatelets (f-GNPs) by differential scanning calorimetry (DSC) (a) and thermogravimetric analysis (TGA) (b).

### 3.2. Stability and Morphological Analysis

Figure 5 illustrates the UV–VIS spectra of the f-GNPs suspended in DW at different sonication times. The sharp absorption peak located at a wavelength of ~283 nm is correlated with the \( \pi \rightarrow \pi^* \) transition of the C=C double bonds for the GNP. The enlarged peak intensity resulting from the increasing particle percentages of the nanofluids directly complied with the Beer–Lambert law.
Zeta potential is defined as the measure of the effective electrical charge on NPs’ surface. Its magnitude represents particle stability; higher magnitudes mean increased stability from the (more) pronounced electrostatic repulsion between NPs [33,34]. Generally, NPs with zeta potentials of $\geq 30$ mV or $\leq 30$ mV are expected to be stable for the long term. Dispersions with zeta potentials of $\leq 25$ mV or $\geq 25$ mV will agglomerate due to interparticle (secondary) interactions [35]. The stability of the f-GNP suspensions in distilled water was elucidated using its zeta potential values and size distributions (Figures 6 and 7). Figure 6 shows the zeta potential and polydispersity index (PDI) of the f-GNPs at a pH value of 7. Both factors are representative of the nature of the electrostatic interactions between the colloidal NPs and can be indicative of the state of stability of the overall solution(s). A more negatively charged ($-39.4$ mV) solution was reported postsonication for 1 h with the f-GNPs at a temperature of 25 °C. The size distributions of the f-GNPs solutions were determined using the concept of dynamic light scattering (DLS), and the results are shown in Figure 7, where the average particle size of the f-GNPs was 442 nm. The size distribution of the f-GNPs was 83.9–1318.6 nm, and its PDI was 0.255 (low).

![Figure 5. UV–VIS variations for pristine GNPs and f-GNPs at 1 and 3 h sonication periods.](image)

![Figure 6. Zeta potential distribution of f-GNPs nanofluids at 25 °C.](image)
The pristine GNPs had a multilayered structure with a smooth surface and edges (low edge defects) within aggregations, as per Figure 8a. Contrarily, surfactal features of the f-GNPs were rough and riddled with defects and wrinkles due to the prevalence of covalent functionalization (see Figure 8b). Postfunctionalization, the acid treatment, and the carboxyl group on the edges and surfaces of GNPs resulted in crumpled and wrinkled sheets, showcasing a minimal blur effect. The morphological and functional group analyses confirmed the strong reactions between the acid molecules and the GNPs–COOH, while the defective folded flakes and rough edges represented the successful occurrence of the covalent functionalization method. The lines evident in the HR-TEM images were wrinkles on the GNPs surface, caused by the inherent stability in the 2D structures. The functionalization method resulted in increased amounts of these lines during sonication, which was further exacerbated from prior wrinkling/waviness.

The dispersion and the stabilizing effect of the f-GNPs nanofluids were determined using SEM. The surfactant-stabilized nanofluids were placed onto a silicon wafer for SEM imaging purposes. Figure 9 shows that the GNPs were flat and smooth flakes with sharp corners of various in-plane sizes. The SEM micrographs suggested that the covalent synthesis of the f-GNPs resulted in a significantly wrinkled structure. The results of the f-GNPs analyses using the energy-dispersive X-ray spectroscopy is detailed in Figure 9, where it encompassed carbon (C), oxygen (O), silicon (Si), sulfur (S), and zirconium (Zr).

It can be seen that the carbon content was 88.95% and the atomic oxygen content was 10.94%. The atomic contents of Si, S, and Zr corresponded to 0.05%, 0.05%, and 0.01%, respectively. The results confirmed the formation of high-quality materials and are in excellent agreement with other works reported in the literature [36–38].

3.3. Thermal Performance of Nanofluids

The effects of using the f-GNPs nanofluids with multiple weight fractions (0.025, 0.05%, 0.075%, and 0.1%) wt % as HTFs within the FPSC system are shown in Figures 10–13. Zero-loss efficiency enhancements at a flow rate of 0.5 kg min⁻¹ were 5.81%, 7.16%, 9.49%, and 10.26%, respectively, and the enhancements at a flow rate of 1 kg min⁻¹ were 6.19%, 7.55%, 9.88%, and 10.65%, respectively, for the f-GNPs-EG nanofluids in the case of multiple weight concentrations (herein the values of the weight concentrations being 0.025, 0.05%, 0.075%, and 0.1%) relative to the base fluid, as per Figure 10. The efficiency enhancements at a flow rate of 1.5 kg min⁻¹ were 8.15%, 9.53%, 11.90%, 12.69% for the f-GNPs-EG
nanofluids in the case of multiple weight concentrations, respectively. For DW:EG (70:30)-based f-GNPs nanofluids in the case of multiple weight concentrations, the efficiency enhancements at a flow rate of 0.5 kg min\(^{-1}\) were 5.70%, 7.03%, 9.31%, 10.06%, and the efficiency enhancements at a flow rate of 1 kg min\(^{-1}\) were 6.13%, 7.47%, 9.78%, and 10.55%, respectively, as can be seen in Figure 11. The enhancements at a flow rate of 1.5 kg min\(^{-1}\) were 8.09%, 9.46%, 11.82%, 12.60% for DW: EG (70:30) based f-GNPs nanofluids in the case of multiple weight concentrations, respectively. Finally, the enhancements at a flow rate of 0.5 kg min\(^{-1}\) are 5.73%, 7.06%, 9.35%, and 10.12% and the enhancements at a flow rate of 1 kg min\(^{-1}\) were 6.14%, 7.49%, 9.81%, and 10.58%, respectively, for the DW:EG (50:50)-based f-GNPs nanofluids in the case of multiple weight concentrations, respectively, as per Figure 12, while the enhancements at a flow rate of 1.5 kg min\(^{-1}\) were 8.11%, 9.48%, 11.84%, and 12.62% for the DW:EG (50:50)-based f-GNPs nanofluids in the case of multiple weight concentrations, respectively. These observations are in excellent agreement with the results reported in [39–41], where the higher flow rates of the f-GNPs resulted in the increased absorption of a heat flux, which enhanced the overall performance of the FPSC. Heat absorption was uniform via the fluid layer, when the f-GNPs flowed at a lower mass fraction, which means that the heat loss rate at the boundaries was lower than that in the case of a higher f-GNPs weight fraction. The majority of heat absorption occurred at the top layer of the nanofluid, resulting in a high-temperature region near the top wall, which increased heat losses and subsequently decreased the efficiency of the FPSC [42].

![HR-TEM image of pristine GNPs](image1.png) (a)  
![HR-TEM image of functionalized GNPs](image2.png) (b)  

**Figure 8.** HR-TEM image of pristine GNPs (a) and functionalized GNPs (b) dispersed in DW.
Figure 9. Cont.
Figure 9. Cont.
Figure 9. SEM and energy-dispersive X-ray spectroscopy (EDX) mapping analysis of the GNPs; (a) SEM microimage at 2 µm; (b) SEM microimage at 5 µm; (c) EDX mapping analysis; (d) EDX elemental analysis; (e) electron image; (f) carbon mapping; (g) oxygen mapping; (h) silicon mapping; (i) sulfur mapping; (j) zirconium mapping.
Figure 10. The theoretical efficiency for the f-GNPs–EG nanofluids with different weight concentrations: (a) 0.5 kg min$^{-1}$; (b) 1 kg min$^{-1}$; (c) 1.5 kg min$^{-1}$.
Figure 11. The theoretical efficiencies for the DW:EG (70:30)-based f-GNPs nanofluids with different weight concentrations: (a) 0.5 kg min$^{-1}$; (b) 1 kg min$^{-1}$; (c) 1.5 kg min$^{-1}$.
Figure 12. The theoretical efficiencies for the DW:EG (50:50)-based f-GNPs nanofluids with different weight concentrations: (a) 0.5 kg min$^{-1}$; (b) 1 kg min$^{-1}$; (c) 1.5 kg min$^{-1}$.
Figure 13. The surface plots of collector efficiency against mass concentration and reduced temperature factor at 1.5 kg min $^{-1}$: (a) EG + GNP, (b) DW:EG (70:30) + GNP, (c) DW:EG (50:50) + GNP.
The heat gain \((F_R(\tau a))\) and heat loss \((F_R U_L)\) coefficients are listed in Tables 2–4, and the values are in a good agreement with those reported in [43–45] on the increase of the \((F_R(\tau a))\) and \((F_R U_L)\) factors of the f-GNPs nanofluids. At a flow rate of 1.5 kg min\(^{-1}\) and concentrations of 0.025–0.1 wt %, the heat gain parameter for EG increased by 8.01%, 10.28%, 11.91%, and 12.48%, respectively (Table 2), while the heat losses increased by 17.38%, 17.47%, 23.11%, and 34.25% at a flow rate of 1.5 kg min\(^{-1}\) and a NPs' concentration of 0.025 to 0.1 wt %. Moreover, it can be seen in Table 3 that the flow of the f-GNPs at 1.5 kg min\(^{-1}\) and NPs concentration of 0.025 to 0.1 wt % increased the \((F_R(\tau a))\) and \((F_R U_L)\) factors of DW:EG (70:30) by 7.96%, 10.21%, 11.83%, and 12.40%, respectively, and the corresponding values of \((F_R U_L)\) improved by 17.26%, 17.35%, 22.95%, and 34.01%, respectively, at similar operating conditions. Finally, Table 4 shows that the heat gains for DW:EG (50:50) at a flow rate of 1.5 kg min\(^{-1}\) and f-GNPs concentrations of 0.025 to 0.1 wt % increased by 7.97%, 10.23%, 11.85%, and 12.42%, respectively, while the respective values of the heat loss parameter increased by 17.30%, 17.39%, 22.99%, and 34.08% at 1.5 kg min\(^{-1}\) and NPs concentrations of 0.025 to 0.1 wt %.

### Table 2. Heat absorbed and heat removal factors at a different flow rate for the EG-based f-GNPs nanofluids.

| \(m\) (kg min\(^{-1}\)) | EG | 0.025% f-GNPs | 0.05% f-GNPs | 0.075% f-GNPs | 0.1% f-GNPs |
|--------------------------|----|---------------|--------------|--------------|------------|
|                          | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) |
| 0.5                      | 0.6802 | 6.1547 | 0.7276 | 6.6331 | 0.7427 | 6.6384 | 0.7536 | 6.9783 | 0.7574 | 7.6509 |
| 1                        | 0.7000 | 6.3315 | 0.7513 | 7.0471 | 0.7669 | 7.0527 | 0.7781 | 7.4035 | 0.7821 | 8.0977 |
| 1.5                      | 0.7068 | 6.3930 | 0.7634 | 7.5044 | 0.7794 | 7.5101 | 0.7909 | 7.8702 | 0.795 | 8.5826 |

### Table 3. Heat absorbed and heat removal factors at a different flow rate for the DW:EG (70:30)-based f-GNPs nanofluids.

| \(m\) (kg min\(^{-1}\)) | DW:EG (70:30) | 0.025% f-GNPs | 0.05% f-GNPs | 0.075% f-GNPs | 0.1% f-GNPs |
|--------------------------|---------------|--------------|--------------|--------------|------------|
|                          | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) |
| 0.5                      | 0.6935 | 6.2736 | 0.7409 | 6.7520 | 0.7560 | 6.7573 | 0.7669 | 7.0973 | 0.7707 | 7.7698 |
| 1                        | 0.7069 | 6.3959 | 0.7582 | 7.1115 | 0.7738 | 7.1171 | 0.785 | 7.4679 | 0.789 | 8.1621 |
| 1.5                      | 0.7114 | 6.4576 | 0.7680 | 7.5490 | 0.7841 | 7.5546 | 0.7956 | 7.9148 | 0.7996 | 8.6271 |

### Table 4. Heat absorbed and heat removal factors at a different flow rate for the DW:EG (50:50)-based f-GNPs nanofluids.

| \(m\) (kg min\(^{-1}\)) | DW:EG (50:50) | 0.025% f-GNPs | 0.05% f-GNPs | 0.075% f-GNPs | 0.1% f-GNPs |
|--------------------------|---------------|--------------|--------------|--------------|------------|
|                          | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) | \(F_R(\tau a)\) | \(F_R U_L\) |
| 0.5                      | 0.6899 | 6.2429 | 0.7373 | 6.7211 | 0.7525 | 6.7266 | 0.7633 | 7.0666 | 0.7672 | 7.7391 |
| 1                        | 0.7051 | 6.3817 | 0.7564 | 7.0973 | 0.7720 | 7.1028 | 0.7832 | 7.4536 | 0.7872 | 8.1478 |
| 1.5                      | 0.7102 | 6.4251 | 0.7668 | 7.5365 | 0.7828 | 7.5423 | 0.7944 | 7.9022 | 0.7984 | 8.6146 |

### 4. Conclusions

This study discussed the development of well-dispersed graphene nanoplatelets (f-GNPs) for the analytical assessment of the impact of utilizing nano-coolants as absorbing mediums upon the energy efficiency of an FPSC. The covalent functionalization approach’s effectiveness was determined using HR-TEM and SEM. The effects of many variables were elucidated during the analyses, encompassing base fluids, weight concentrations, fluid flow rates, input fluid temperatures, and input heat rates. The analysis leads to the following conclusions:

i. The UV–VIS spectra showed a sharp absorption peak at a wavelength of ~283. In the temperature range of 500–800 °C, f-GNPs suffered from a weight loss of nearly 24%, which was attributable to the breakdown of the air’s graphitic carbon nanostructures.
ii. The zeta potential of the prepared f-GNPs had a negative charge of $-39.4 \text{ mV}$. The EDX analyses of the GNP s confirmed its high carbon content of 88.95% and lower atomic oxygen content of 10.94%. The atomic contents of Si, S, and Zr were 0.05%, 0.05%, and 0.01%, respectively.

iii. The zero-loss efficiency at a flow rate of 1.5 kg min$^{-1}$ was 0.6992, 0.7038, and 0.7026 for EG, DW: EG (70:30), and DW: EG (50:50), respectively. This means that the increased flow rate improved the performance of the collector. $F_R \tau_\alpha$ and $F_R U_L$ increased in tandem, with the flow rate increasing from 0.5 to 1.5 kg min$^{-1}$.

iv. At a flow rate of 1.5 kg min$^{-1}$, the efficiency enhancements were 8.15%, 9.53%, 11.90%, and 12.69% for the EG-based f-GNPs nanofluids with multiple weight concentrations, respectively, while the enhancements were 8.09%, 9.46%, 11.82%, and 12.60% for the DW:EG (70:30)-based f-GNPs nanofluids for multiple weight concentrations, respectively. The enhancements were 8.11%, 9.48%, 11.84%, and 12.62% for the DW:EG (50:50)-based f-GNPs nanofluids in the case of multiple weight concentrations, respectively. The results revealed that GNP s-based basefluids could function as a kind of good and alternative conventional working fluid in heat transfer applications.

v. At a flow rate of 1.5 kg min$^{-1}$ and 0.1 wt %, $F_R \tau_\alpha$ and $F_R U_L$ increased by 12.48% and 34.25%, 12.40% and 34.01, 12.42% and 34.08% for the EG, DW:EG (70:30)-based, and DW:EG (50:50)-based f-GNPs nanofluids, respectively.

In future work, more research has to be conducted using CNTs-based hybrid nanofluids and graphene-based nanofluids. The nanofluids with heat transfer promoters need to be investigated further to produce maximum cost-effective thermal performance. Future research should be devoted to developing FPSCs with built-in PCM as a latent heat storage, when the nanofluids are used as HTFs. In addition, a porous medium has to be analyzed numerically and experimentally with different porosity materials and nanofluids.

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Nomenclature and Abbreviations

| Symbol | Description |
|--------|-------------|
| $A_c$  | surface area of collector, m$^2$ |
| Al$_2$O$_3$ | aluminium oxide |
| CNTs | carbon nanotubes |
| $C_p$ | specific heat coefficient, kJ kg$^{-1}$ K$^{-1}$ |
| CuO | copper oxide |
| DASC | direct absorption solar collector |
| DSC | differential scanning calorimetry |
| DW | distilled water |
| EDX | energy dispersive X-ray |
| EG | ethylene glycol |
| ELS | electrophoretic light scattering |
| FPSC | flat plate solar collector |


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