Experimental Studies of Nanofluid Thermal Conductivity Enhancement and Applications: A Review*

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ABSTRACT

In many applications, there is a critical need for enhancing the poor thermal conductivity of conventional fluids in order to develop efficient heat transfer fluids. This requirement can be met through dispersing nanometric particles in a given base fluid such as water, ethylene glycol, oil or air. The resulting nanofluids enhanced thermal conductivity of the base fluids. In order to evaluate this enhancement, nanofluid thermal conductivity is required to be measured. Several methods and techniques are covered in the present contribution. In addition, enhancements recorded experimentally are reviewed and summarized. Different parameters affecting on such enhancement are covered, including: nanoparticle concentration, size, shape and thermal conductivity. In addition, base fluid type, nanofluid bulk temperature and dispersion techniques are also covered parameters. However, nanofluids have the potential to contribute in several practical applications including solar thermal, transportation, electronic cooling, medical, detergency and military applications. In the present work, a brief overview of evolution in the use of nanofluids in some applications has been presented. According to this contribution, there is a critical need for further fundamental and applications of nanofluids studies in order to understand the physical mechanisms of using nanofluids as well as explore different aspects of applications of nanofluids.

Nomenclature

 k_{bf} = Base fluid thermal conductivity, W/m.K

 k_{nf} = nanofluid thermal conductivity, W/m.K

 $q = \text{heat flux, W/m}^2$

r = radial coordinate, m

T= temperature, K

t = time, s

 α_{nf} = nanofluid thermal diffusivity, m²/s

 ω = electric current frequency, Hz

Abbreviations

BG= BioGlycol

CNF= Carbon nanofiber

CTAB= Hexadecyltrimethylammonium bromide

DE= Decene

DI= Deionized water

DWCNT= Double-walled carbon nanotubes

DO= Diathermic oil

EG= Ethylene glycol

EO= Engine oil

HTF= High-temperature heat transfer fluid

HTO= Heat transfer oil

MEG= Mono ethylene glycol

MO= Mineral Oil

MWCNT= Multi-wall carbon nanotube

NSAQ= Nanosperse AQ

OAK+= Potassium oleate

PO= Pump oil

TH66= Therminol 66

1. Introduction

Heat transfer is one of the most important processes in many industrial and consumer products. For more than a century, scientists and engineers have made great efforts to enhance the inherently poor thermal conductivity of conventional fluids [1,2]. In 1873, Maxwell [3] proposed an idea of using metallic particles to enhance the electrical or thermal conductivity of matrix materials. He presented a theory for effective conductivity of slurries, by dispersing millimeter- or micrometer-sized particles (typically have size between 0.1 and 100 µm [4]) in liquids. However, major problems such as sedimentation, erosion, and high pressure drop prevented the usual micro-particle slurries to be used as heat transfer fluids. Nanofluids, which is a dilute suspension of nanometer-size particles or fibers (typically less than 100 nm) dispersed in a fluid such as water, oil, and ethylene glycol (EG) [5], have emerged as a potential candidate for the design of heat transfer fluids [6]. According to their potential applications in the heat transfer field, nanofluids have been a subject of intensive investigation [7–12].

According to the definition of micro- and nano-particles size, nanoparticles have surface/volume ratio 1000 times larger than that of microparticles [13]. This in turn, allows improving thermal properties of nanofluids rather than microparticles-colloidal suspensions, since heat transfer occurs on the surface of the particle [14]. Compared with microparticles, nanoparticles stay suspended much longer in base fluids, with very little settling under static conditions, unlike micron-sized suspensions [15]. However, strong van der Waals interactions generate an aggregation tendency between nanoparticles [16]. Therefore, different techniques are utilized

to minimize long-term particles aggregation. This process is quite critical in preparation of nanofluids as particles clustering has been reported as features increasing thermal conductivity of nanofluids [17–19]. Moreover, the number of atoms present on the surface of nanoparticles is very large, as opposed to the interior [20]. These unique properties of nanoparticles can be exploited to develop nanofluids with an unprecedented combination of the two features most highly desired for heat transfer systems: extreme stability and ultrahigh thermal conductivity. Furthermore, because the nanoparticles are so small, they may reduce erosion and clogging dramatically. Other benefits envisioned for nanofluids include decreased demand for pumping power, reduced inventory of heat transfer fluid, and significant energy savings [21].

This discovery brought about a wave of studies in this area, predominantly experimental confirmation of the huge potential of nanofluids as well as efforts to theorize the phenomenon. In this paper, various techniques used to measure thermal conductivity are covered. Then, experimental work carried on studying the thermal conductivity enhancement of nanofluids against their base fluids is reviewed. This review aims to define parameters investigated experimentally through the literature in order to find out points of agreement and conflict in the obtained results to understand the thermal behavior of nanofluids. Moreover, different applications using nanofluid is also reviewed.

2. Thermal Conductivity Measurement Techniques

Measuring the thermal conductivity of nanofluids can be carried out with different methods. The most common techniques for this purpose are the transient ones including: transient hot-wire method [22–37], temperature oscillation method [38,39], and $3-\omega$ method [40–42]. Some other methods such as steady-state parallel-plate technique, micro-hot strip method, and optical beam deflection technique have also been utilized by some researchers [43–45].

2.1 Transient Hot-Wire Method

The transient hot-wire (THW) method is the most widely used experimental technique for measuring fluids thermal conductivity, as it is an easy and low cost method to be implemented. It is a standard transient dynamic technique based on the measurement of the temperature rise in a defined distance from a linear heat source (hot wire) embedded in the test material. A hot wire is placed in the fluid, which functions as both a heat source and a thermometer [46–48]. The ideal mathematical model of the method is based on Fourier's law, assuming the hot wire as an ideal, infinite thin and long heat source in an infinite surrounding from homogeneous and isotropic material with constant initial temperature. According to Fourier's law, when the wire is heated, fluid of higher thermal conductivity corresponds to a lower temperature rise.

The mathematical model which describes the relation between thermal conductivity $k_{\rm nf}$ and measured temperature T using the THW method is explained and summarized as follows [47]. Assuming a thin, infinitely long line source dissipating heat into a fluid reservoir, the energy equation in cylindrical coordinates can be written as:

$$(1/\alpha_{nf})(\partial T/\partial t) = (1/r)\partial [r(\partial T/\partial r)]/\partial r$$
 (1)

The initial condition can be written as shown in Eq. (2):

$$T\big|_{t=0} = T_0 \tag{2}$$

while the boundary conditions are defined by Eq. (3) and Eq. (4) as follows:

$$\lim_{r\to 0} \left(r\left(\partial T/\partial r\right)\right) = \left(q/2\pi\right)\left(1/k_{nf}\right) \tag{3}$$

and

$$\left(\frac{\partial T}{\partial r}\right)_{r=\infty} = 0 \tag{4}$$

If the temperatures of the hot wire at times t_1 and t_2 are T_1 and T_2 , then by neglecting higher-order terms, the thermal conductivity can be approximated as [5]:

$$k_{nf} = (q/4\pi) \ln(t_1/t_2)/(T_1 - T_2)$$
 (5)

Therefore, in order to determine k_{nf} experimentally using THW method according to Eq. (5), a constant electric power supply is used to heat the wire with a constant heat flux, q, at a time step, t. A Wheatstone-bridge circuit is used to determine the temperature increase of the wire from its change in resistance. Although the THW is an easy, fast response and low cost method; its accuracy can be affected by nanoparticle interactions, sedimentation and/or aggregation, and natural convection during extended measurement times. In addition, the assumptions of an infinite wire-length and the ambient acting like a reservoir may also introduce errors [42,49].

2.2 Temperature Oscillation Method

This method is based on the oscillation method proposed by Roetzel et al. [50] and further developed by Czarnetzki and Roetzel [38]. Applying this method requires measuring the temperature response of the nanofluid sample when a temperature oscillation or heat flux is imposed. The measured temperature response of the sample is an indication of averaged or localized thermal conductivity in the direction of sample chamber height [51]. The experimental set up of this method is explained in details by Paul et al. [42].

$2.33-\omega$ Method

The $3-\omega$ method is quite similar to the THW method, as it uses a radial flow of heat from a single element which acts both as the heater and the thermometer. However, the main major difference is the use of electric current frequency dependence response instead of the time dependent response which is utilized by the TWH method. When a sinusoidal current at frequency, ω , passes through the metal wire, a heat wave can be generated at a frequency of 2ω , which is deduced by the voltage component at frequency 3ω . More details about this method are available in [42,52].

2.4 Other Thermal Measurement Methods

The short-hot-wire method is an improved design of the hot-wire method, in which boundary effects can be taken into account, It has been applied by [53,54]. Another modification of THW is carried out by Mintsa et al. [55], who inserted a mixer into his THW experimental devices to avoid nanoparticle aggregation/deposition in the suspensions. In order to avoid interference between the detector and heater, Ali et al. [56]

separated them by combining the THW method with a laser beam displacement method.

2.5 Optical Measurement Methods

In order to improve the thermal conductivity measurement accuracy, optical measurement techniques have been proposed as non-invasive methods [57–61]. The accuracy improvement resulted from separating detector and heater from each other avoiding the unavoidable interference between them in the THW method. In addition, optical techniques provide faster measurement, within a few microseconds compared with 2 to 8 s of measurement using the THW method. This fast response helps in avoiding natural convection effects.

One of the proposed optical techniques is thermal-lensing (TL) method, which is applied by Rusconi et al. [57,62]. In this method, a laser-diode module was used as a heater and a photodiode was used as a thermometer to measure the temperature difference as optical signals.

The forced Rayleigh scattering (FRS) method is an extension of quasi elastic Rayleigh light scattering technique [63,64]. In the FRS method, two intersecting laser beams are absorbed by the sample in order to generate a spatially periodic temperature distribution. Analyzing the time dependence of the light scattered by the thermal fluctuations inside the sample was required in order to measure thermal diffusivity. The FRS was also used by many researchers to measure the thermal conductivity of nanofluids [36,58,65].

Optical beam deflection was another optical technique which was used to measure thermal conductivity of nanofluids [44,66]. In this method, two parallel lines using a square current were used to heat the nanofluid sample. Dual photodiodes were used to capture light signals that indicate the temperature change of nanofluids.

Other optical techniques were applied by different researchers, such as the Transient Optical Grating method [59], Laser Flash method [61], Modern Light Flash technique [67]. However, there is a critical need to apply different measurement techniques for the same nanofluids in order to compare the accuracy and reproducibility of their experimental results.

3. Experimental Studies on Conduction Heat Transfer of Nanofluids

3.1 Liquid-Based Nanofluids

Liquid-based nanofluid is used to enhance thermal properties of a "liquid" base fluid. Since thermal conductivity is the most important parameter responsible for enhanced heat transfer, many experimental works have been reported on this aspect. Choi et al. [68] developed a new project to design and analyze a microchannel heat exchanger that uses liquid-nitrogen as the cooling fluid which used to cool high-heat-load x-ray optical elements. They focused on the thermal conductivity of the fluid itself rather than on channel size to develop a new heat transfer fluid concept that enables heat transfer enhancement without a large pumping power increase and without cryogenic coolants. A summary of maximum measured thermal conductivity enhancement for nanofluids is given in Table 1.

According to the previous literature, there are different types of nanoparticles that were commonly used, including oxides, nanotubes, metals and carbides. These nanoparticles were dispersed in different base fluids. The number of contributions mentioned in the present work versus different nanoparticles is shown in Fig. 1, while Fig. 2 represents the number of publications per year, mentioned in this paper and concerned with experimental studies on thermal conductivity in nanofluids

Table 1 Summary of maximum measured thermal conductivity enhancement for nanofluids

		Dontialos Tura /		Looding	Enhancement*			Pa	rameters Investigat	ed		
Ref.	Year	Particles Type / Size (nm)	Base-fluid	Loading (% vol.)	Enhancement* (%)	Particle Concentration	Particle Size	Particle Shape	Particle Thermal Conductivity	Base Fluid Type	Temperature	Preparation Technique
[22]	1993	$Al_2O_3 / 13$	Water	4.33	32							
		SiO ₂ / 12		2.3	1.1	\checkmark						\checkmark
		TiO ₂ / 27		4.35	11.6							
[69]	1997	$Al_2O_3 / 33$	Water	5	29							
		CuO / 36		5	60	\checkmark						\checkmark
		Cu / 35	HE-200 Oil	0.052	44							
[70]	1998	$Al_2O_3 / 13$	Water	4.33	32		\checkmark		✓			
		TiO ₂ / 27		4.35	11.6		v		V			
[71]	1999	$Al_2O_3 / 38$	Water	4.3	10							
			EG	5	18							
		CuO / 24	Water	3.41	12	\checkmark	\checkmark		\checkmark	\checkmark		
			EG	4	23							
[43]	1999	Al ₂ O ₃ / 28	Water	5.5	16							
			EG	8	41							
			EO	7.4	30		,					
			PO	7.1	20		\checkmark					\checkmark
		CuO / 23	Water	9.7	34							
			EG	14.8	54							
[72]	2000	Cu / 100	Water	7.5	78							
			HE-200 Oil	7.5	43	\checkmark	\checkmark					\checkmark
[73]	2001	Cu / < 10	EG	0.3	40	√			√			
[74]	2001	MWCNT /	Oil	1	160			√	•			
		Ø25×50,000				<u> </u>		<u> </u>				
[23]	2002	SiC (sphere)/26	DI	4.2	15.8							
		SiC (cylinder) /		4	22.9							
		600 SiC	EG	3.5	13	\checkmark		\checkmark		\checkmark		
		(sphere)/26										
		SiC (cylinder) / 600		4	23							
[75]	2002		EG	4	17							
		CeO ₂ / 29		4	18							
		TiO ₂ / 40		4	13	✓	\checkmark		\checkmark			✓
		CuO / 33		4	17	-	-		•			•
		Fe ₂ O ₃ / 28		4	16							

	-			T 31	T-1	Parameters Investigated								
Ref.	Year	Particles Type / Size (nm)	Base-fluid	Loading (% vol.)	Enhancement* (%)	Particle Concentration	Particle Size	Particle Shape	Particle Thermal Conductivity	Base Fluid Type	Temperature	Preparation Technique		
		ZnO / 56		4	21									
[76]	2002	Al ₂ O ₃ / 60.4	Water	5	23									
			EG	5	30	,								
			PO	5	38	\checkmark				\checkmark		V		
			Glycerol	5	27									
[77]	2003	MWCNT /	DI	1	7									
		Ø15×30,000	EG	1	12.7	\checkmark		\checkmark		\checkmark				
			DE	1	19.6									
[78]	2003	Au / 10-20	Water	0.00026	21									
			Toluene	0.011	8.8	\checkmark					\checkmark			
		Ag / 10-20	Water	0.001	16.5									
[39]	2003	Al ₂ O ₃ / 38.4	Water	4	24	,			,		,			
		CuO / 28.6		4	36	\checkmark			\checkmark		\checkmark			
[79]	2004	MWCNT / Ø20-	Water	0.84	21									
		60 ×(few				\checkmark					\checkmark			
		tens×10³)												
[24]	2005	DWCNT / Ø5	Water	1	8									
		MWCNT /		0.6	34			\checkmark				\checkmark		
		Ø130×10000												
[80]	2005	Al ₂ O ₃ / 11	Water	1	9									
		$Al_2O_3 / 47$		1	2		\checkmark				\checkmark			
		$Al_2O_3 / 150$		1	0.5									
[25]	2005	Fe / 10	EG	0.55	18	✓						✓		
[26]	2005	MWCNT /	EG	1	12.4	,		,		/				
		Ø20~50	EO	1	8.5	\checkmark		\checkmark		\checkmark				
[27]	2005	TiO ₂ / 15	DI	5	30									
		(sphere)												
		$TiO_2 / \emptyset 10 \times 40$		5	33	\checkmark		\checkmark						
		(cylinder)												
[81]	2005	Al ₂ O ₃ / 10	Water	0.5	100		√				√			
[82]	2006		EG	2.5	125						•			
[]		40	_ ~			\checkmark	\checkmark							
[83]	2006	Ag-Cu / 10	PO	0.003	15									
[00]				0.006	33									
				0.009	12	\checkmark								
				0.015	0.02									
[28]	2006	MWCNT / Ø40	Water	0.49	80			√			√			
[-0]						<u> </u>		▼			у			

		Doubleles War /		I og din r	Enhance	-		Pa	rameters Investigat	ed		
Ref.	Year	Particles Type / Size (nm)	Base-fluid	Loading (% vol.)	Enhancement* (%)	Particle Concentration	Particle Size	Particle Shape	Particle Thermal Conductivity	Base Fluid Type	Temperature	Preparation Technique
[84]	2006	MWCNT / Ø10-	Water	1	11.3							
		30×10,000-										
		50,000				✓		./		✓		
		CuO / 33		1	5	•		\checkmark		V		
		SiO ₂ / 12		1	3							
		CuO / 33	EG	1	9							
[85]	2006	CuO / 29	Water	6	52	✓					\checkmark	
		$Al_2O_3 / 36$		10	30	V					,	
[44]	2006	Au / 4	Ethanol	0.018	1.3							
		Au / 2	Toluene	0.024	1.4	✓		✓				
		C ₆₀ -C ₇₀		0.378	0.816	V		V				
		fullerenes										
	2006	TiO ₂ / 34	Water	6.8	6	✓						
[87]	2007	TiO ₂ / 20	Water	2	4.2	✓	✓					
[29]	2007	$Al_2O_3 / 38$	Water	3	8							
			EG	3	10.6							
		$TiO_2 / 10$	Water	3	11.4							
			EG	3	15.4							
		$TiO_2 / 34$	Water	3	8.7							
			EG	3	12.3							
		$TiO_2 / 70$	Water	3	6.4	\checkmark	\checkmark			\checkmark		
		5 0 / 10	EG	3	7.5							
		ZnO / 10	Water	3	14.2							
		ZnO / 30	Water	3	11.5							
		7.0.460	EG	3	21							
		ZnO / 60	Water	3	7.3							
- FO 01	200=	11.0.706	EG	3	10.7							
[88]	2007	$Al_2O_3 / 36$	Water	6	28	✓	\checkmark				\checkmark	
[04]	2005	Al ₂ O ₃ / 47	D.	6	26		•				•	
[31]	2007	$TiO_2 / 25$	DI	1	14.4							
		$Al_2O_3 / 48$	EC	1	4	✓	\checkmark		\checkmark			
		Fe / 10	EG	0.3	16.5	-	-		•			
[22]	2005	W0 ₃ / 38	m-1	0.3	13.8							
[32]	2007	Au / 1.65	Toluene	0.003	8							
		$Al_2O_3 / 20$	Water	14.6	22	\checkmark		✓	\checkmark			
		$TiO_2 / 40$		2.6	6.5	•		•	•			
		CuO / 33		4.18	16.5							

		Particles Type /		Loading	Enhancement*			Parameters Investigated						
Ref.	Year	Size (nm)	Base-fluid	(% vol.)	(%)	Particle Concentration	Particle Size	Particle Shape	Particle Thermal Conductivity	Base Fluid Type	Temperature	Preparation Technique		
		CNF /	•	0.89	41.4									
		$150\emptyset \times 10,000$												
[89]	2007	MWCNT / Ø10- 30×10,000- 50,000	Water	1	7									
		CuO / 33		1	5									
		$SiO_2 / 12$		1	3.2									
		CuO / 33	EG	1	9.1	\checkmark	\checkmark	\checkmark	\checkmark	\checkmark		\checkmark		
		MWCNT / Ø10- 30×10,000- 50,000	МО	0.5	8.7									
		C ₆₀ –C ₇₀ fullerenes / 10		5	6									
[90]	2008	Al ₂ Cu / 31	Water	2	96									
		Al ₂ Cu / 101		2	61									
		$Ag_2Al/33$		2	106									
		Ag ₂ Al / 120		2	75									
		Al ₂ Cu / 31	EG	2	84	\checkmark	\checkmark			\checkmark				
		Al ₂ Cu / 101		2	56									
		$Ag_2Al/33$		2	96									
		Ag ₂ Al / 120		2	62									
[34]	2008	TiO ₂ / 15	EG	5	18									
		Al / 80		5	45	\checkmark			\checkmark		\checkmark			
[40]	2008	Al ₂ O ₃ / 45	DI	4	13.3									
		•	EG	4	9.7	\checkmark								
[91]	2008	Fe ₃ O ₄ / 6.7	Kerosene	< 2%	300	√						√		
[35]	2009	Al ₂ O ₃ / 282	Water	4	17.7	·						•		
		$Al_2O_3 / 282$	EG	3	16.3	\checkmark	\checkmark							
[92]	2009	TiO ₂ / 21	Water	2	7	√					√			
[55]	2009	Al ₂ O ₃ / 36	Water	18	31	•					•			
		$Al_2O_3 / 47$		18	31	✓	✓				✓			
		CuO / 29		16	24	·	·				•			
[93]	2009	TiO ₂ / 21	DI	3	7.2	√					√			
[94]	2009	Al ₂ O ₃ / 15-50	Water	4	10.1	<u>·</u> ✓					-	√		
[95]	2010	•	Water	3	11.5	<u>·</u> ✓					√	· ·		
[96]	2010	Al ₂ O ₃ / 12	Water	4	5.4	<u>·</u> ✓	√			√	<u> </u>	<u> </u>		

	•	Particles Tyme /	· ·	Loading	Enhancement*			Pa	rameters Investigat	ed		
Ref.	Year	Particles Type / Size (nm)	Base-fluid	(% vol.)	(%)	Particle Concentration	Particle Size	Particle Shape	Particle Thermal Conductivity	Base Fluid Type	Temperature	Preparation Technique
			EG	4	14.3							
		$Al_2O_3 / 10$	EG-Water	3	11.3							
		$Al_2O_3 / 50$	(50:50 wt.%)	3	10.4							
[97]	2010	Fe ₃ O ₄ / 15	Kerosene	1	34.6	✓					✓	
[98]	2011	SiC / 100	DI	3	7.2	✓						
[99]	2012	SWCNT / 100- 600	Water	0.3	12.1	✓					✓	
[100]	2012	SWCNT / 100- 600	EG	0.21	15.5	✓					✓	
[101]	2012	CuO / 25	Water	7.5	32.3	/						
			MEG	7.5	21.3	\checkmark						✓
[102]	2012	MWCNT / Ø5- 20	НТО	2	15	✓					✓	
[103]	2013	SiO ₂ / 10	Water	1.2	11	/						
		SiO ₂ / 60		1.2	13	\checkmark						
[104]	2013	Al ₂ O ₃ / 36.5	EG-Water	8	17.89	,					,	
		CuO / 27	(50:50 wt.%)	8	24.56	\checkmark					\checkmark	
[105]	2014	γ-Al ₂ O ₃ / 13	Water	6	14.5							
		SiO ₂ / 15		6	10.8							
		TiO ₂ / 13.5		4	15.1	\checkmark						
		α -Al ₂ O ₃ / 24.4		6	18.6							
[106]	2014	Sn-SiO ₂ / 50- 100	ТН66	5	13	✓			✓			
[107]	2015	Al ₂ O ₃ / 40	Water	4	14.4	✓					✓	✓
[108]	2015	DWCNT / Ø3 +		1	33							
		ZnO / 10-30 (50:50 vol.%)	(60:40 wt.%)			\checkmark					\checkmark	
[109]		NiFe ₂ O ₄ / 8	DI	2	17.2	✓					✓	
[110]		AG / 5-25	DI	0.5	16	✓						✓
[111]	2015	Al ₂ O ₃ / 13	EG-Water (60:40 wt.%)	2	8.4							
			EG-Water (50:50 wt.%)	2	12.6	\checkmark				✓	\checkmark	
			EG-Water (40:60 wt.%)	2	16.2							
[112]	2015	MgO / 40	EG-Water (40:60 wt.%)	3	34.43	✓			✓		✓	

	-	Particles Type /		Loading	Enhancement*	-		Pa	rameters Investigat	ed		
Ref.	Year	Size (nm)	Base-fluid	(% vol.)	(%)	Particle Concentration	Particle Size	Particle Shape	Particle Thermal Conductivity	Base Fluid Type	Temperature	Preparation Technique
[113]	2016	SiC / 30	DO	8.0	7.36	✓					✓	✓
[114]	2016	ZnO / 50	EG	2.4	13	✓					✓	✓
[115]	2016	CuO / 55-66	Water	2	24							
			EG	2	21	\checkmark				\checkmark	\checkmark	\checkmark
			EO	2	14							
[116]	2016	AG / 20	НТО	0.171	41	✓					√	
[117]	2016	TiO ₂ / 5	EG	7	19.52	✓					√	
[118]	2016	AG / 10	НТО	0.6	36.3	✓					✓	
[119]	2016	S-SWCNT /	Water	048	8.1							
		Ø1-2 x1000-										
		3000										
		L-SWCNT /		0.48	16.2							
		Ø1-2 x5000-				\checkmark					\checkmark	
		30,000										
		MWCNT /		0.48	5							
		Ø10-										
		30x30,000										

^{*}Enhancement % is calculated based on: Enhancement (%) = $[(k_{nf} - k_{bf})/k_{bf}]$ *100

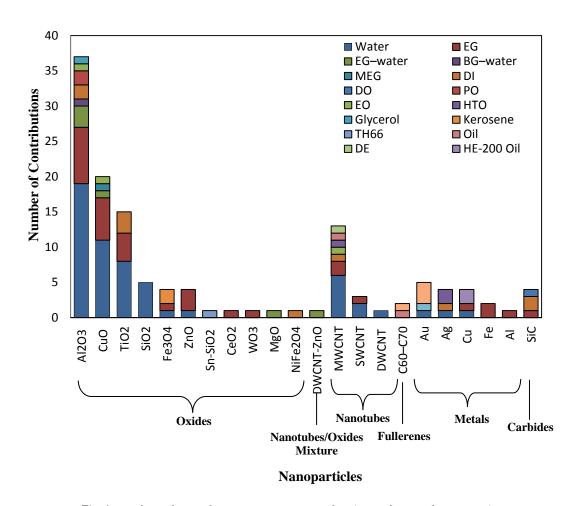


Fig. 1 Number of contributions vs. nanoparticles (according to this review)

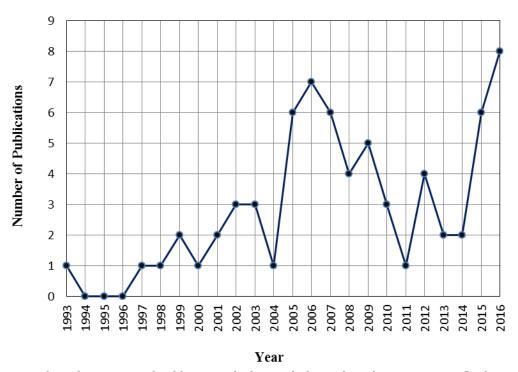


Fig. 2 Number of experimental publications dealing with thermal conductivity in nanofluids per year (according to this review)

3.2 Gas-Based Nanofluids

In such type, nanoparticles are dispersed in gaseous base fluids. This mixture is also known as aerosols [120,121]. Studying heat transfer characteristics of aerosols has a great importance not only for thermal applications, but also for environmental and climate studies [121–123]. Although natural convection heat transfer in gas-based nanofluids is investigated [124–127], limited research work was done in the field of thermal conductivity enhancement and forced convection of aerosol nanoparticle systems [128].

In the field of thermal conductivity enhancement, Bibire et al. [129] carried out a mathematical analysis to study thermal and electrical conductivities of atmospheric aerosols. They applied the scale relativity model [130]. They found that the enhancement in the effective conductivity is inversely proportional to atmospheric nanoparticle diameter for a given atmospheric nanofluid.

4. Parameters Affecting Nanofluids Thermal Conductivity

According to literature listed in

, there are different investigated parameters affecting thermal conductivity of nanofluids, as illustrated in Fig. 3. In this section, such parameters will be discussed based on past work observations.

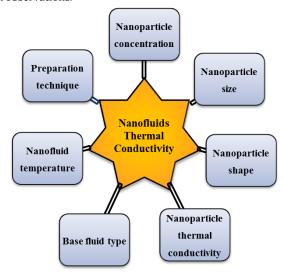


Fig. 3 Parameters affecting the thermal conductivity of nanofluids

4.1 Nanoparticle Concentration

Many researchers investigated the effect of volumetric loading of nanoparticles in suspension on thermal conductivity enhancement. Most of them found that the more nanoparticles concentration, the more enhanced thermal conductivity of the suspension. Some of researchers found the relation to be nonlinear [25,79,86,87], while others found it linear [22,23,26–29,31,32,34,35,39,40,55,69–78,82,84,85,88–102,104–119,131–133].

However, this relation is associated with a limited nanoparticle loading. This fact was observed by Ceylan et al. [83], who tested suspended Ag-Cu alloy nanoparticles (10 nm) in PO. They found that the thermal conductivity enhancement increases with increasing volumetric loading until a certain peak point. Beyond this point the thermal conductivity decreases until it reaches its value for the base fluid. On the other hand, the proportionality relationship between

nanoparticles volume fraction and thermal conductivity enhancement could not be proven by Putnam et al. [44]. They investigated thermal conductivity of Au suspensions in ethanol and toluene and fullerene in toluene for very low loadings (<<1% vol.). Their results showed fluctuating small enhancements. This may be attributed to their inability to synthesize and study well-dispersed nanoparticle suspensions, especially for nanofluids with particles concentrations higher than 1% vol.

4.2 Nanoparticle Size

Particles size plays a great role in enhancing thermal conductivity of nanofluids. It represents the most significant difference between nanofluids and micron-sized suspensions. The effect of nanoparticles size is not limited to suspension stability, but extends to include thermal properties. The effect on enhancing thermal conductivity of nanofluids was studied frequently. It was found by Lee et al. [71] that the effective thermal conductivity of nanofluids increases with decreasing particle size. They observed this relation through comparing their results with reported data provided by Masuda et al. [22]. This fact was confirmed later by Wang et al. [43], who compared their results with Masuda et al. [22] and Lee et al. [71] results. Another proof of the validity of this conclusion comes through comparing Xuan and Li [72] results with Eastman et al. [69] ones, as the larger particle size of Cu used by Xuan and Li [72] leaded to drop the enhancement from 44% to 12% despite the relatively higher concentration than used by Eastman et al. [69]. In the same context, achieving a near amount of enhancement by 100 nm compared to 35 nm, it was required to raise the concentration from 0.052 vol.% to 7.5 vol.%. This inverse relation between particle size and thermal conductivity enhancement was proven by many authors [29,31,55,75,80-82,87,88,90,96,119].

However, this relation is not always true, especially if particle shape parameter is interfered. Based on [23] and [89] results, large cylindrical-shaped and MWCNT nanoparticles can enhance the conductivity more than smaller spherical-shaped ones if dispersed in the same base fluid. Moreover, Pak and Cho [70] recommended selecting larger particles to enhance heat transfer performance, based on their results. This finding was confirmed later by and Hwang et al. [89] Beck et al. [35]. Hwang et al. [89] results showed that CuO (33nm) enhanced thermal conductivity higher than SiO₂ (12nm) for the same base fluid and concentration. Beck et al. [35], who studied thermal conductivity of Al₂O₃/water and Al₂O₃/EG nanofluids for particles sizes ranging from 8 to 282 nm, also found that the enhancement of thermal conductivity decreases as the particle size decreases below about 50 nm. They attributed this to a decrease in the thermal conductivity of the nanoparticles as a result of increased phonon scattering effect.

4.3 Nanoparticle Shape

The effect of nanoparticles shape was studied by Xie et al. [23]. Their results indicated that the cylindrical-shaped nanoparticles showed higher enhancement than spherical-shaped ones for the same base fluid, despite their larger average size. This conclusion was confirmed later by [26–28,77,84] results. Moreover, the increase in length-to-diameter ratio of the dispersed nanotubes leads to the increase of the thermal conductivity enhancement [24,32]. In the same

context, the effective thermal conductivity measured by Choi et al. [74] showed that nanotubes (MWCNT) yield an anomalously nonlinear increase in the conductivity compared to predicted linear behavior. On the other hand, fullerenes, which are carbon molecules in the form of a hollow sphere, ellipsoid, tube and many other shapes [134], showed lower enhancement. This result was concluded by Putnam et al. [44] who dispersed C₆₀-C₇₀ fullerenes in toluene and compared their effect on enhancement of thermal conductivity against dispersing Au nanoparticles. Their results indicated that fullerenes showed lower enhancement than Au at loadings << 1% vol. However, these data was not confirmed for higher volumetric loadings. Another lower enhancement of fullerenes in mineral oil nanofluids was observed by Hwang et al. [89] when compared its enhancement with MWCNT in the same base fluid, even at higher concentration.

4.4 Nanoparticle Thermal Conductivity

Select nanoparticles having higher thermal conductivity was recommended by Pak and Cho [70] to enhance heat transfer performance. This finding was confirmed later by [32,34,39,71,89] results. In the same context, by comparing [111] and [112] results, it is clear to find that for the same base fluid, MgO showed significant higher thermal conductivity enhancement compared to the A1203 system, despite the larger MgO particle size (40nm) compared to (13 nm) for A1203 nanoparticles. This comparison indicates that the particle thermal conductivity has a stronger effect than particle size. According to Eastman et al. [73], nanofluids containing metallic particles can achieve a large improvement in effective conductivity compared to either base fluids or nanofluids containing oxide particles. Moreover, Cingarapu et al. [106] found that addition of ceramics encapsulated phase change nanoparticles enhances thermal conductivity of nanofluids than conventional HTFs. They also observed that such type of particles improved the heat transfer and thermal storage properties of HTFs. However, Yoo et al. [31] reported that determining nanofluids thermal conductivity was not primarily affected by suspended nanoparticles thermal. Thermal conductivities of some materials used as nanoparticles are listed in Table 2.

Table 2 Thermal conductivities of some materials used as nanoparticles

Particle	Thermal conductivity (W/mK)	Reference
Al ₂ O ₃	40	[135]
CuO	76.5	[89]
Fe_2O_3	6	[136]
MgO	54.9	[137]
SiO_2	1.34-1.38	[89,138]
TiO_2	8.4	[29]
ZnO	29	[29]
Ag	429	[139]
Al	238–273	[138,139]
Au	310	[139]
Cu	401	[139]
Fe	75–80	[138,139]
MWCNT	2000~3000	[140,141]
$C_{60}-C_{70}$	0.4	[89]
(Fullerene)	0.4	
SiC	490	[142]

4.5 Base Fluid Type

For the nanofluids using the same nanoparticles, the thermal conductivity improvement was found to be inversely proportional to the base fluid thermal conductivity, regardless nanoparticles shape [29]. This conclusion was achieved and confirmed through observing higher thermal conductivity enhancement for nanofluids with EG as a base fluid compared to others that with water base fluid [23,71,76,77,84,89,96].

In contrast of that conclusion, observations reported by [26,115] indicated that at the same volume fraction, using EG as the base fluid leaded to achieving higher enhancements compared to the enhancement in case of using EO. This conflict was reconfirmed through measurements obtained by [90,115], as water-based nanofluids they tested showed higher enhancement In thermal conductivity compared to EG-based ones at the same volume fraction.

With the beginning of the 2000s, a new research trend emerged. It was based on investigating the use of mixed base fluids instead of conventional ones. In that trend, Beck et al. [96] discussed the use of EG-water mixture as a base fluid with mixing ratio of 50:50% by weight. As a result of improved thermal conductivity of such new base fluid, adding nanoparticles showed more enhancement than dispersing in conventional base fluids.

However, using mixed base fluids requires intensive study to select the appropriate mixing ratio, as it dramatically affects thermal conductivity. As an evidence on this fact, Abdolbaqi et al. [131] observed that the enhancement in thermal conductivity by dispersing Al_2O_3 (13 nm) in BG-water base fluid mixed at a ratio of 40:60 wt.% was 24% at 2 vol.% with temperature of $80^{\circ}C$. This enhancement dropped to 13% by opposing the base fluid mixing ratio to be 60:40 wt.%. This result also agrees with Usri et al. [111] results, who tested Al_2O_3 (13 nm) in EG-water base fluid, as they observed that increasing the EG concentration decreases the thermal conductivity. Thermal conductivities of some base fluids that used in this review as are listed in Table 3.

Table 3 Thermal conductivities of some base fluids

Base fluid	Thermal conductivity (W/mK)	Reference
EG	0.257	[138]
EO	0.139-0.146	[115]
Ethanol	0.161-0.171	[143,144]
Glycerol	0.285	[138]
Kerosene	0.145-0.168	[143,145]
Toluene	0.133	[142]
Water	0.608	[146]

4.6 Nanofluid Temperature

The effect of nanofluid temperature on improving their thermal conductivity has been studied by many researchers. Patel et al. [78] observed that the increments in thermal conductivity of the nanofluids were directly proportional to temperature. This result was in agreement with observations reported later by [34,39,55,80,81,88,95,99,100,104,109,111–113,115–118,131]. Even in case of dispersing nanotubes in base fluids, the thermal conductivity enhancement increased with increasing temperature nonlinearly [28,79,102,119,147]. However, Experiments carried out using MWCNT at

temperatures 60-70°C showed destabilization of the nanofluid. However, Li and Peterson [85] concluded that dependence of the effective thermal conductivity on the bulk temperature is much weaker than the dependence on the volume fraction. This conclusion was confirmed later by [107,108] results. In the same context, Beck et al. [96] results indicated that the thermal conductivity enhancement behavior over temperature variation followed closely that of the base fluid. This observation was concluded through studying different base fluids including: water, EG and EG-water mixture. Results reported by [97,114] was in agreement with that conclusion.

In contrast with previous literature, Duangthongsuk and Wongwises [92] results indicated that nanofluid thermal conductivity enhancement slightly decreases with increasing temperature in reverse relation. This result agreed with Turgut et al. [93] findings, despite the insignificant dependence on temperature.

4.7 Preparation Technique

Nanofluid dispersion technique plays a great role in stabilizing nanofluid and in improving its thermal conductivity, as it might change the morphology of the nanoparticles [43,72]. This section includes investigating the effect of adding different additives to control the pH value of the prepared nanofluid. In addition, the effect of sonication time will be covered.

Investigating nanofluids pH value was started in early stage. The effect of adding acid (HCl) or base (NaOH) on electrostatic repulsive forces among the particles was firstly reported by Masuda et al. [22]. This effect kept the nanoparticles dispersed. Xie et al. [76] results showed that the increase in the difference between the pH value and isoelectric point1of Al₂O₃ resulted in the enhancement of the effective thermal conductivity. This may be the reason for indicating much higher thermal conductivity than that of Wang et al. [75] for Al₂O₃-EG nanofluid although the particle size used by Xie et al. [76] was double that of the particles of Wang et al. [75]. Moreover, Xie et al. [76] results indicated that the thermal conductivity enhancement decreases with an increase in pH value. Zhu et al. [94] observed that that enhancement in thermal conductivity increased with increasing the pH value up to 8-9, then the relation turns to be inversely proportional as reported before by Xie et al. [76]. They also recommended using a chemical dispersant combined with adjusting the pH of the suspension to improve the thermal conductivity. In the same context, Abareshi et al. [95] observed the best crystallinity at a pH of

On selecting additive type side, Assael et al. [24] also found that the dispersant type affects the enhancement achieved, as they observed larger enhancement in case of adding Nanosperse AQ (NSAQ) compared Hexadecyltrimethylammonium bromide (CTAB) addition. While Parametthanuwat et al. [110] found that the addition of potassium oleate surfactant (OAK+) with 1 wt.% improved the working properties, especially with Ag nanoparticles. Moreover, using oleic acid boosted the rise in the heat transfer rate.

The sonication (or homogenization) time is a critical parameter should be considered in preparing nanofluids, as it plays a major role in stabilizing the suspension [24,110,113-

115,131]. Khedkar et al. [101] observed that increasing the sonication time enhances the thermal conductivity of the nanofluids until certain limits. They attributed this result to the increased Brownian motion of small particles and agglomeration. However, a moderate sonicating time should be applied, as very low time leaves the nanoparticles bundles untangled, while intensive sonication breaks down the agglomerates size. In case of using nanotubes, longer sonication time decreases the nanotubes length, which leads to quick suspensions precipitation [24,148,149]. These results are in agreement with observations reported by Hong et al. [25], who found that thermal conductivity enhancement increased with increasing the sonication time up to 50 minutes while saturation was shown after this point [25]. In 2015, Buonomo et al. [107] proposed a simple procedure to estimate the minimum sonication time required to obtain a stable nanofluid mixtures. Philip et al. [91] achieved an extraordinary thermal conductivity enhancement of 300% via applying a magnetic field on Fe₃O₄ nanoparticles dispersed in base fluid in order to form the linear chain-like structures.

5. Applications of Nanofluids in Practical Community

Applications of nanofluids have a wide spectrum in the practical field, so that they require separate review articles to be covered. However, this section focuses on some applications where nanofluids contribution to the thermal conductivity enhancement is significant. Fig. 4 shows different practical applications covered in the present work.

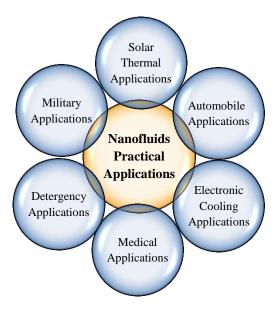


Fig. 4 Practical applications of nanofluids

4.1 Solar Thermal Applications

Nanofluids can be used in order to improve the performance of solar thermal devices ranging from the solar water heaters to Concentrated Solar Power (CSP) plants. This is as a result of many advantages of nanofluids over conventional HTFs. These include: small particles size allow them to be fluidized to pass through pumps, micro-channels and piping without any adverse effects. In addition, nanoparticles represent an absorption medium allowing the fluid to absorb solar energy directly [150]. Moreover, by studying photo thermal characteristics of the nanofluids [151-155], they

¹ the pH value at which a molecule carries no net electrical charge [248].

showed improved optical properties compared to that of their base fluids [156], such as low emittance in the infrared region [150].

Research papers [157–176] investigated the thermal performance enhancement achieved by using nanofluids in different types of solar collectors. In addition, the optical properties of nanofluids used in solar collectors were studied in [177–185]. Recent CSP systems require high operating temperature and high heat storage capacity [52]. Since nanofluids have improved heat transfer and thermal storage properties [106], they can be used as a HTF in CSP plants rather than conventional HTFs. This, in turn, can improve efficiencies and reduce the costs of CSP plants [186–188]. However, many authors are concerned with using nanofluid in CSP systems and performed investigations on the achievable enhancement in the performance of these systems [189–198].

4.2 Automobiles Applications

In recent years, the energy crisis and fuel economy created a competition between automobiles manufacturers. According to this, designers have to improve the aerodynamic designs of vehicles in order to reduce the amount of energy required to overcome the drag force. Unfortunately, they face the fact that more than 50% of the total vehicle energy output is lost in overcoming the aerodynamic drag. The large radiator position in the vehicle front is partly responsible for this fact [199]. Therefore, it is required to replace poor cooling medium, such as EG-water mixture, with nanofluids to remove heat from relatively smaller size [200,201].

Many researchers have been attracted to investigate, either experimentally or numerically, the use of nanofluids as engine coolant [202-208]. They reported excellent enhancements in the thermal properties of nanofluids, such as EG-based nanofluids, compared to conventional coolants, e.g. 50/50 mixture of EG and water. In addition, some authors indicated that the frontal area of the radiator can be reduced up to 10% due to the use of nanofluid coolants. This in turn can lead to reduce aerodynamic drag and save fuel up to 5% [204]. Engine cooling is not only the application of using nanofluids in automobiles, but also they can be used to cool other moving parts in an automobile. Tzeng et al. [209] dispersed nanoparticles into engine transmission oil. They reported that the thermal performance of nanofluids has a clear advantage as they produced the lowest transmission temperatures at both low and high speeds.

4.3 Electronic Cooling Applications

Heat transfer at medium and low temperatures is also affected by the improvements raised by nanofluids. Nanofluids are used as the working fluid in heat pipes and thermosyphons, which can be utilized for compact device cooling, e.g. electronic devices. Some researchers investigated the use of nanofluids in heat pipes [210–217], while others studied the effect of using nanofluids on the performance of thermosyphons [218–226]. They observed obvious enhancements in thermosyphons performance, as a result of the reduction of thermosyphons thermal resistance due to the use of nanofluids. Other researchers investigated the use of nanofluids in electronic devices of cooling systems [227,228]. They also reported higher cooling performance in their coolers.

4.4 Medical Applications

Recently, nanofluids contributed in a wide range of applications in medical applications and biomedical industry [229]. Nanofluids have been used in nano-medicine applications as iron based nanoparticles can be used as nanodrug delivery vehicles [230-237]. Nanofluids can be also used in cancer therapeutics. They can be utilized in cancer imaging and drug delivery, by using magnetic nanofluids which guide the particles up the bloodstream to a tumor with magnets. In addition, they can be used to kill cancerous cells without affecting the nearby healthy cells by producing higher temperatures around tumors [238-241]. On the other hand, nanofluids can also be used in Cryosurgery, which is a procedure that uses freezing to destroy undesired tissues. This procedure can be regarded as a novel method of cancer treatment [241,242]. In addition, nanofluids can be used to avoid risk of organ damage by cooling around the surgical region in surgery operations [243].

4.5 Other Applications

Many other fields were invaded by nanofluids. They can be used in detergency. Nanofluids differ from conventional simple liquids as they have different behavior of spreading and adhesion on solid surfaces [244–247]. According to this fact, nanofluids arise as excellent candidates in lubrication, processes of soil remediation, oil recovery and detergency. Nanofluids are also applied in some military applications. In such applications, high heat flux cooling fluids are required in order to remove a large amount of heat from both military mechanical and electrical devices, e.g. submarines and high power laser [243].

6. Conclusions

The present work reviewed recent research progress achieved in enhancing thermal conductivity using nanofluids. In addition, some practical applications that used the improvements resulting from the use of nanofluids were presented including solar thermal applications, automotives, electronic cooling, medical, detergency and military applications. According to this contribution, it can be found that the main parameters affecting heat transfer properties of the base fluid are nanoparticles concentration, size, shape, thermal conductivity, base fluid type, nanofluid temperature and preparation technique. The key parameter which had the most significant effect was nanoparticle concentration. It was found in most literature that it has a direct proportional relationship with thermal conductivity enhancement. However, this relation was found to be limited and extra particle loadings dramatically affect such enhancement. For particles size, it was quite agreed that is the relation was proportional with thermal inversely conductivity improvement for spherical particles, while large size cylindrical-shaped can particles enhance effective conductivity than small spherical ones. The particle shape was also found to be a critical parameter. Nanotubes were found to increase thermal conductivity compared to spherical particles, while fullerenes showed lower enhancement. Selecting particles with higher thermal conductivity, including using metallic particles, was recommended by many authors to increase the nanofluid thermal conductivity. In addition, it was found by some researchers that the thermal conductivity improvement was found to be inversely proportional to the base fluid thermal conductivity, while other authors reported an opposite finding. Moreover, using mixed base fluids has been emerged as a new research trend. However, mixing fluids was found to be investigated intensively, as small difference in mixing ratio affects thermal conductivity significantly. Nanofluid temperature effect was found to be much weaker than volumetric concentration, but it also found to increase thermal conductivity by raising the fluid temperature. Agglomeration leads to losing the advantage of using nanofluids. Therefore, using appropriate surfactants and sonication are required to improve nanoparticles dispersion

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