



Research report

Heat transfer characteristic of advanced loop thermosyphon with check value (ALT/CV) containing nanofluids

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Abstract

This report is focus on heat transfer characteristic advanced loop thermosyphon with check value (ALT/CV) containing nanofluids. It will highlight theories for investigating heat transfer characteristic. Points of importance will be emphasize, with significance given to the heat transfer characteristics of ALT/CV and their use in this experiment. The chapter I present work aims to study the effects of operating temperature, loop and check valve on thermal performance of nanofluids in an ALT/CV. The chapter II present theoretical consideration to study the thermosyphon, heat transfer characteristics, dimensionless, nanofluids and surfactant. The chapter III describes the results and discussions of silver nanofluids properties when preparing for filled as working fluids in ALT/CV. It is divided into a number of sections to determine the thermal properties of silver nanofluids containing surfactant. The chapter IV describes the heat transfer rate behaviour of an advanced loop thermosyphon with check valve(ALT/CV) which is filled with silver nanofluids and containing oleic surfactant (OA) and potassium oleate surfactant (OAK+). Also included are the explanation of dimensionless and the behaviour characteristics in the ALT/CV. The filling ratios are 30, 50 and 80% with respect to evaporator volume. The heat was supplied of 20%, 40%, 60%, 80% and 100% of heater (2,000 Watt). Five working fluids are: deionized water, deionized water based silver nanoparticles concentration of 0.5 wt% (NP), NP containing 0.5, 1 and 1.5 wt% of OA and OAK+ respectively. The dimensionless parameters on ALT/CV are such as $\frac{Lo_{size}}{D_i}$,

Pr , Bo , Ja , Co , Cd , Ga_m , Pe_m , Ar , Gr and Z . Then, all of dimensionless was to create a correlation equation with Kutaeldze number for predicting heat transfer of ALT/CV, shows as below;

$$q = 3.24 \left[\frac{Lo_{size}^6 \cdot Pr^{6.2} \cdot Bo^{6.4} \cdot Ja^{6.2} \cdot Co^{5.2} \cdot Cd^{3.8} \cdot Ga_m^{1.5}}{Pe_m^{2.7} \cdot Ar^{0.4} \cdot Z^{0.2} \cdot Gr^{0.6}} \right]^{1.57} \times \left[\rho_v h_{fg} \left(\frac{\rho_v - \rho_l}{\rho_v^2} \right) \right]^{0.25}$$

The correlation equation is used to calculate and construct a design for the oven installed with ALT/CV (OALT/CV) shows in Chapter V.

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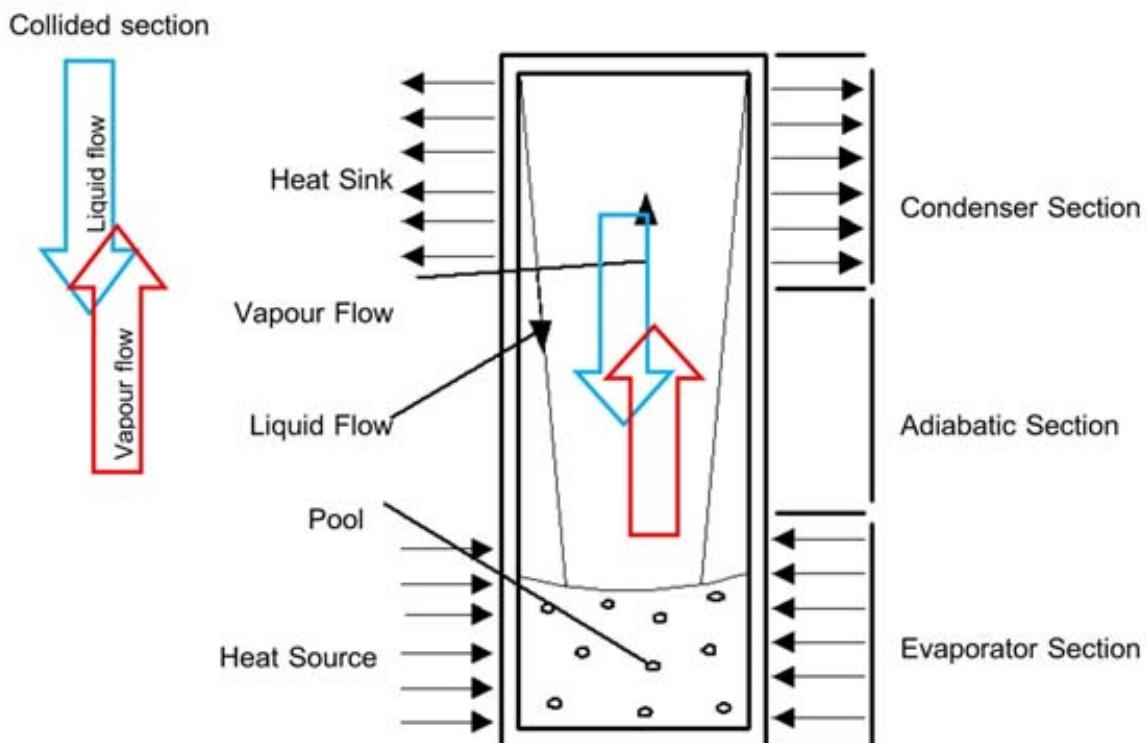
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CHAPTER I INTRODUCTION

1. Background and explanation of the problem

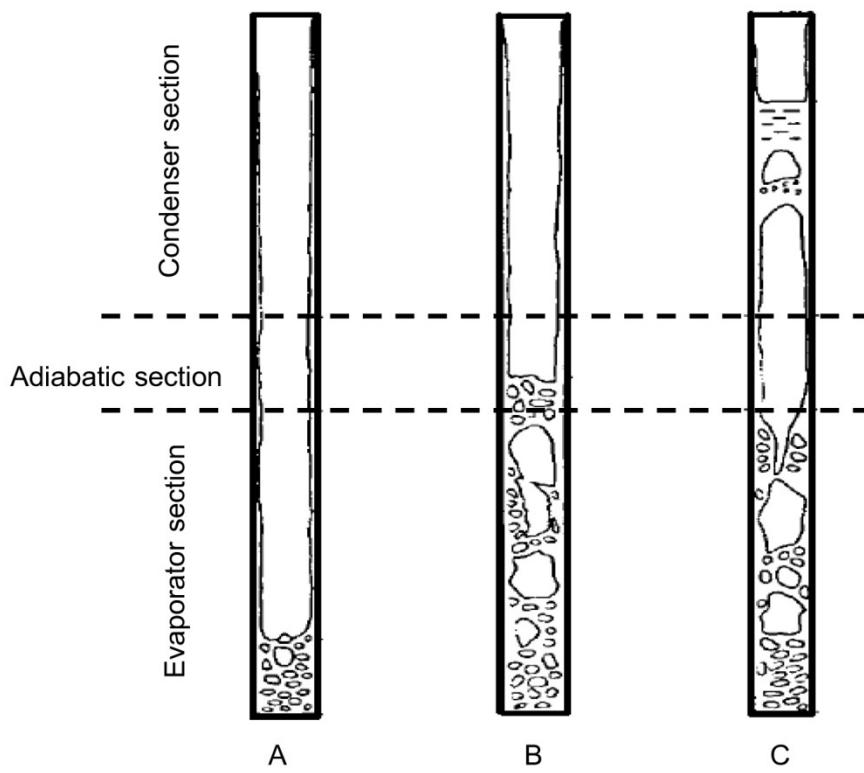
A traditional two-phase closed thermosyphon (TPCT) is essentially a gravity-assisted wickless heat pipe, which utilizes the heat of evaporation and condensation as a working fluid. Contrary to the conventional heat pipe that uses the capillary force to return the liquid to evaporator, the TPCT uses gravity to return liquid to condensate. Since the evaporator of a TPCT is located in the lowest position, the gravitational force will support the capillary force's duty [1] shown in Figure 1. The TPCT has a number of advantages these include it's simple structure, has very small thermal resistance, high efficiency and low manufacturing costs. It has, therefore, been widely used in various applications. These include industrial heat recovery, electronic component cooling, turbine blade cooling and solar heating systems.



12

13 Figure 1 Schematic of the TPCT [2]

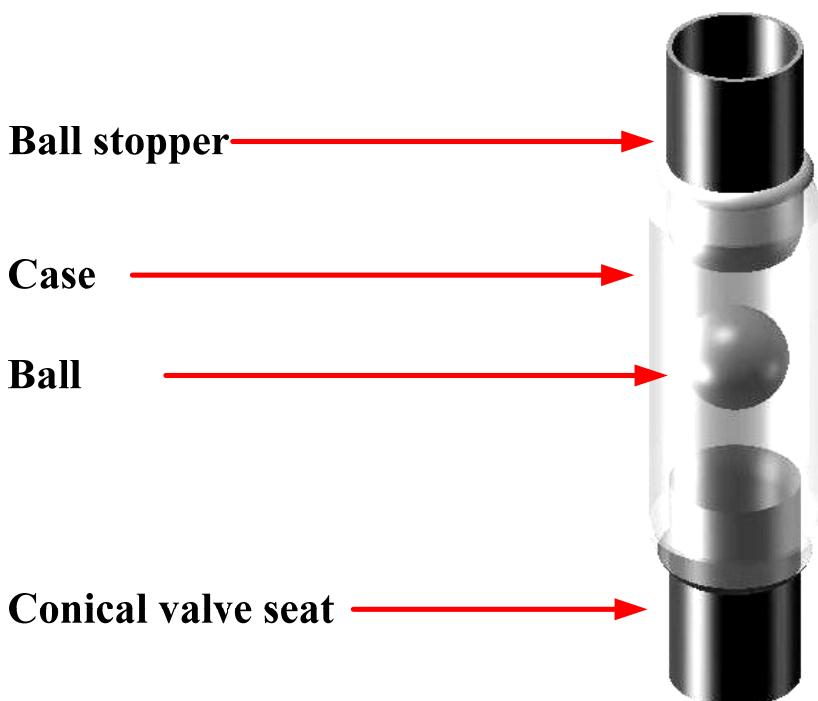
14 However, the traditional TPCT had disadvantage that is collision of liquid and vapour
15 movement of inside them. Moreover, the point of importance, traditional TPCT had disadvantage
16 behaviour caused to decrease heat transfer/heat flux [3, 4]. Figure 2A showed the condensate on the
17 wall surface of the cooled section flow down on the form of liquid film to the heated section through
18 the adiabatic section. Then break down into rivulet at a comparatively small heat flux. The film
19 breakdown occurs; the heat transfer in the breakdown area is decreases and then the wall temperature
20 rise. The wall temperature, however, does not rise continuously but reaches an equilibrium. This is
21 because the breakdown heat flux is relatively small and because heat conduction through the tube wall
22 appears gradually to prevail. This region is called “dry out” [5, 6]. The flow patterns of this region are
23 presumed to correspond to Figure 2(B) and (C), and then the heated wall temperature, which had had
24 a uniform temperature distribution, rose sharply at a lower half position of the heated section when the
25 critical condition was reached. Since the rate of temperature rise of this region was faster than that of
26 the dry-out region, this region is called the “burn-out” [5, 7, 8].



27

28 Figure 2 Flow pattern in traditional TPCT [5]

29 The check valve (CV) in Figure 3 is a buoyancy valve that consists of a stainless steel ball
30 and copper tube, in which a ball stopper and conical valve seat are provided at the ends of the check-
31 valves case: a conical valve seat is provided at the bottom of the case and a ball stopper is provided at
32 the top of the case, respectively. The ball can move freely between the ball stopper and the conical
33 valve-seat. The conical valve seat contacts the stainless-steel ball in order to prevent a reversal of the
34 flow of the working fluid. The ball stopper allows the working fluid to travel to the condenser section
35 for transferring heat. Thus, the CV was controlled vapour and fluid flow.



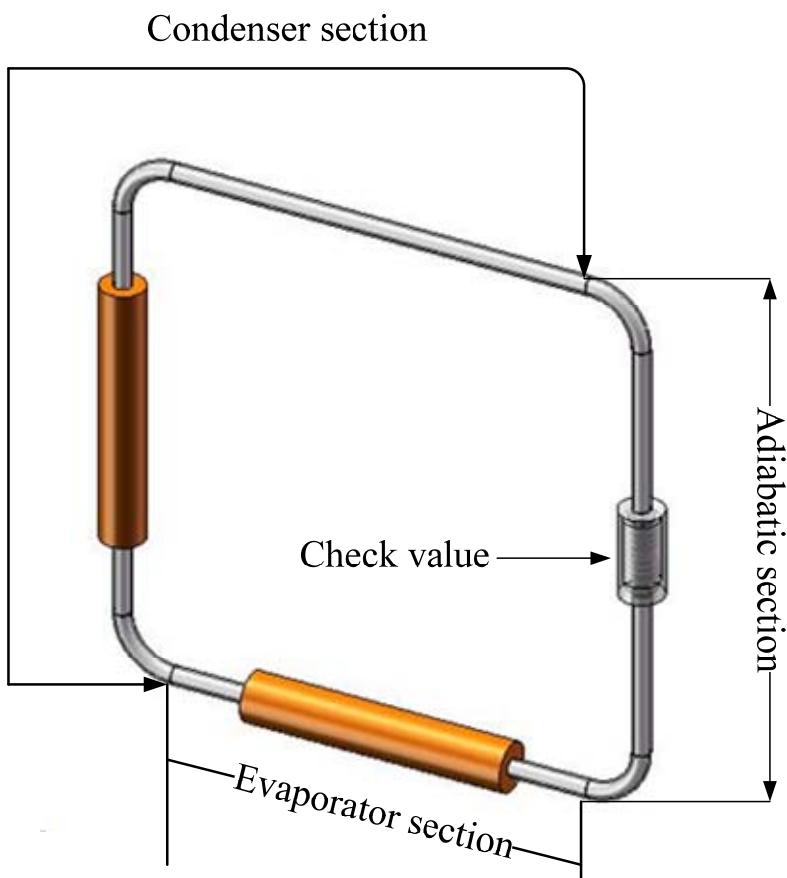
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37 Figure 3 The check valve

38 For these reasons, advanced loop thermosyphon with check valve (ALT/CV) designed to
39 solve the weak points in the limitation of heat being received in the evaporator area and the limitation
40 of the condensing area. The TPCT made adjustments to for the problems which used an advanced
41 loop thermosyphon with check valve (ALT/CV). The new type of thermosyphon (ALT/CV), shown in
42 Figure 4 utilizes evaporation and the condensation of a working fluid inside to transport heat with a
43 loop and control flow with check valve. Expectedly, the heat transfer rate of ALT/CV was superior in
44 heat transfer rate over of all experimental conditions studied.

45 Finally, cooling is one of the most important challenges facing numerous industrial sectors.
46 Despite the considerable amount of research and development focusing on industrial heat transfer
47 requirements, major improvements in cooling capabilities are still insufficient. This is due to the fact
48 that conventional heat transfer fluids possess poor heat transfer properties. One common method used
49 to overcome this problem is to increase the surface area available for heat exchange. This leads to
50 impractical enhancement in the size of the heat management system. Thus, there is a current need to
51 improve the heat transfer capabilities of conventional heat transfer fluids. However, traditional fluids
52 have poor heat transfer properties compared to most solids [9-11]. A new class of heat transfer fluid
53 called “Nanofluids” which are engineered by suspending ultra fine metallic or non-metallic particles
54 of nanometer dimensions in traditional fluids cooling [11, 12]. The applications would benefit from a
55 decrease in the thermal resistance of the heat transfer fluid. Using nanofluids leads to smaller electric
56 potential to reduce thermal resistance and improved heat transfer fluid. Nanofluids can be used in the
57 transportation, electronics, medical, food and manufacturing sectors. In addition, nanofluids consist of
58 such particles suspended in liquid, typically convectional heat transfer liquids to enhance the thermal
59 conductivity and convective heat transfer performance of the base liquids [10, 12].

60 More exploratory research is indeed required to benchmark the scope and applicability of
61 these fluids in engineering systems. This present work aims to study the effects of operating
62 temperature, loop and check valve on thermal performance of nanofluids in an advanced loop
63 thermosyphon with check valve(ALT/CV). Thus, the ALT/CV used in this study was a special type
64 that uses nanofluids in the ALT/CV to transfer heat from the evaporator to condenser, without any
65 external energy requirement. The primary task is to create the correlation of the Kutateladze number.
66 The ALT/CV was designed and tested in order to increase the heat transferred in oven application,
67 which intensifies the ALT/CV’s effectiveness.



68

69 Figure 4 Schematic of the ALT/CV

70

71 **2. Objectives of this study**

72 To study the thermal properties of silver nanofluids

73 To study the effect of loop check valve, evaporator area, condenser area, operating temperature and
74 working fluids of heat transfer rate of an advanced loop thermosyphon with check valve containing
75 nanofluids contained oleic acid and potassium oleate surfactant.

76 To study the effect of dimensionless parameters on the heat transfer rate of an advanced loop
77 thermosyphon with check valve containing nanofluids contained oleic acid and potassium oleate
78 surfactant.

79 To create a correlation equation for predicting the heat transfer rate of an advanced loop
80 thermosyphon with check valve containing nanofluids contained oleic acid and potassium oleate
81 surfactant.

82 The equation in 3 was used to calculate and construct a design for the prototye of oven an advanced
83 loop thermosyphon with check valve (OALT/CV) containing nanofluids contained oleic acid and
84 potassium oleate surfactant.

85

86 **3. Scope of study**

87 Table 1 Controlled and variable parameters about Silver nanofluids properties[13]

The controlled parameters	<ul style="list-style-type: none">• Deionized water mixed Silver nanofluids concentration of 0.5 wt% (NF)• Operating temperature of 20 °C to 80 °C• Shear rate ranges were 100 s⁻¹ to 103 s⁻¹
Independent Variable	<ul style="list-style-type: none">• Working fluid was:<ul style="list-style-type: none">◦ Deionized water (DI-water)◦ DI-water containing surfactant◦ NF◦ NF containing surfactant
Dependent Variable	<p>Surfactant were:</p> <ul style="list-style-type: none">• Concentration of Oleic acid (OA) was 0.5, 1, and 1.5 wt%• Concentration of Potassium oleate (OAK⁺) was 0.5, 1, and 1.5 wt% <p>The dependent variable was:</p> <ul style="list-style-type: none">• Thermal conductivity, specific heat, density, viscosity, contact angle and application of thermal enhancement• Rheological behaviour

89 Table 2 Controlled and variable parameters about Silver nanofluids thermal properties[13]

The controlled parameters	<ul style="list-style-type: none">• Cylinder diameter and height of 1,500 mm and 3,000 mm.• The volumetric flow rate of 0.2–1.5 l/min
Independent Variable	<ul style="list-style-type: none">• Heater (2,000 Watt) with diameter and height of 500 mm and 2,000 mm.• The heat was supplied of 20%, 40%, 60%, 80% and 100% of heater.• Working fluid was:<ul style="list-style-type: none">◦ Deionized water (DI-water)◦ DI-water containing surfactant◦ NF◦ NF containing surfactant
	<p>Surfactant were:</p> <ul style="list-style-type: none">• Concentration of OA was 0.5, 1, and 1.5 wt%• Concentration of OAK^+ was 0.5, 1, and 1.5 wt%
Dependent Variable	Heat transfer enhancement

91 Table 3 Controlled and variable parameters about ALT/CV thermal performance and behaviour

The controlled parameters	<ul style="list-style-type: none"> • ALT/CV was made from copper tube with inside diameter of 12.70 mm. • The temperature of cooling fresh air is at 25°C. • The ambient temperature is at 25°C. • The velocity of air inlet of condenser section is at 0.6 m/s.
Independent Variable	<ul style="list-style-type: none"> • The heat supplied 20%, 40%, 60%, 80% and 100% from heater 2,000 Watts. • The filling ratio was 30%, 50% and 80% respect to the evaporator section volume. • The ALT/CV had three sections, evaporator, adiabatic, and condenser, of loop equal size of 30 cm, 40 cm, and 50 cm, within installation fin of 8 FPI at condenser section. • Working fluid was: <ul style="list-style-type: none"> ◦ Deionized water (DI-water) ◦ DI-water containing surfactant ◦ NF ◦ NF containing surfactant
	<p>Surfactant were:</p> <ul style="list-style-type: none"> • Concentration of OA was 0.5, 1, and 1.5 wt% • Concentration of OAK^+ was 0.5, 1, and 1.5 wt%
Dependent Variable	<p>The dependent variable was:</p> <ul style="list-style-type: none"> • Heat transfer rate (W) • Heat flux (W/m^2) • Thermal resistance ($W/^\circ C$)

92

93

94 Table 4 Controlled and variable parameters about OALT/CV

The controlled parameters	<ul style="list-style-type: none"> • The heat supplied was LPG burner. • The filling ratio was 80% respect to the evaporator section volume. • ALT/CV was made from copper tube with inside diameter of 12.70 mm. • The ALT/CV had three sections, evaporator, adiabatic, and condenser, of loop equal size of 30 cm, 40 cm, and 50 cm, with installation fin of 8 FPI at condenser section.
The variable parameters	<p>Working fluid was:</p> <ul style="list-style-type: none"> • Deionized water (DI-water) • DI-water containing surfactant • NF • NF containing surfactant
Dependent Variable	<p>Surfactant were:</p> <ul style="list-style-type: none"> • Concentration of OA was 0.5, 1, and 1.5 wt% • Concentration of OAK^+ was 0.5, 1, and 1.5 wt% <p>The dependent variable was:</p> <ul style="list-style-type: none"> • Total colour • Texture • Effectiveness • Sensory score

95

96

97 **4. Expected Benefits**

98 The thermal properties of silver nanofluids, will be clarified;

99 The effect of loop, filling ratios and check valve of heat transfer rate of an advanced loop
100 thermosyphon with check valve containing nanofluids contained oleic acid and potassium oleate
101 surfactant, will be clarified.

102 The effect of dimensionless parameters of heat transfer rate of an advanced loop thermosyphon with
103 check valve containing nanofluids contained oleic acid and potassium oleate surfactant, will be
104 clarified.

105 The correlation equation to predict the heat transfer rate of an advanced loop thermosyphon with
106 check valve containing nanofluids contained oleic oleic acid and potassium oleate surfactant, will be
107 achieved.

108 The prototype of an advanced loop thermosyphon with check valve oven containing nanofluids
109 contained oleic acid and potassium oleate surfactant (OALT/CV), will be achieved.

110 In the future. an advanced loop thermosyphon with check valve in the study can be using
111 practically in the many industries.

112

113

CHAPTER II THEORETICAL CONSIDERATION

114

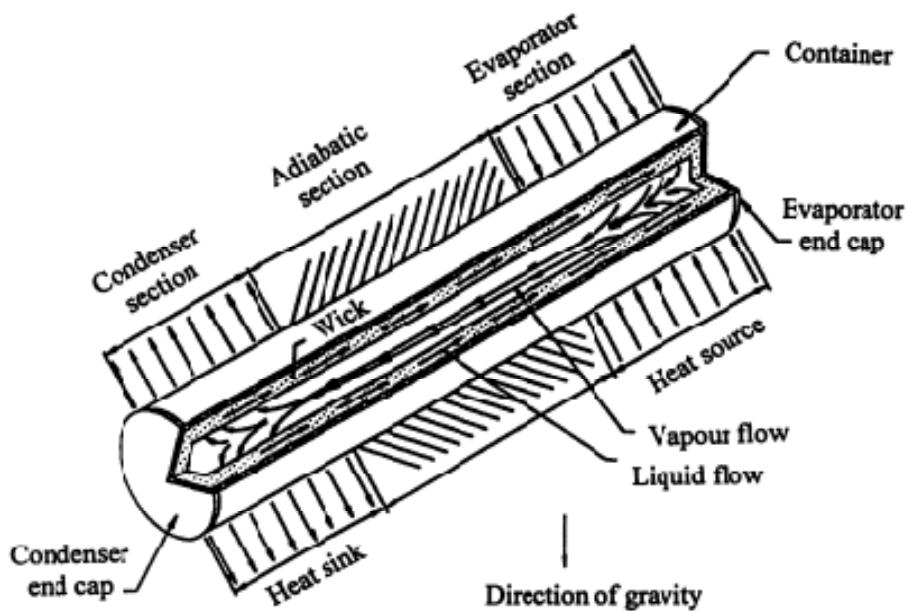
1. Thermosyphon

115

Recently, investigators have been paid great attention to heat pipes and thermosyphons.

116

Their mode of operation has also been investigated. The operation of a heat pipe is easily understood by using a cylindrical geometry, as shown in Figure 5. The components of a heat pipe are a sealed container (pipe wall and end caps), a wick structure, and a small amount of working fluid in liquid state, which is in equilibrium with its own vapour. The length of the heat pipe is divided into three parts: the evaporator section, adiabatic section and condenser section. In the selection of a suitable combination of three basic components, inevitably a number of conflicting factors may arise.



122

123

124 Figure 5 Schematic of a conventional heat pipe showing the principle of operation and circulation of
125 the working fluid [14]

126

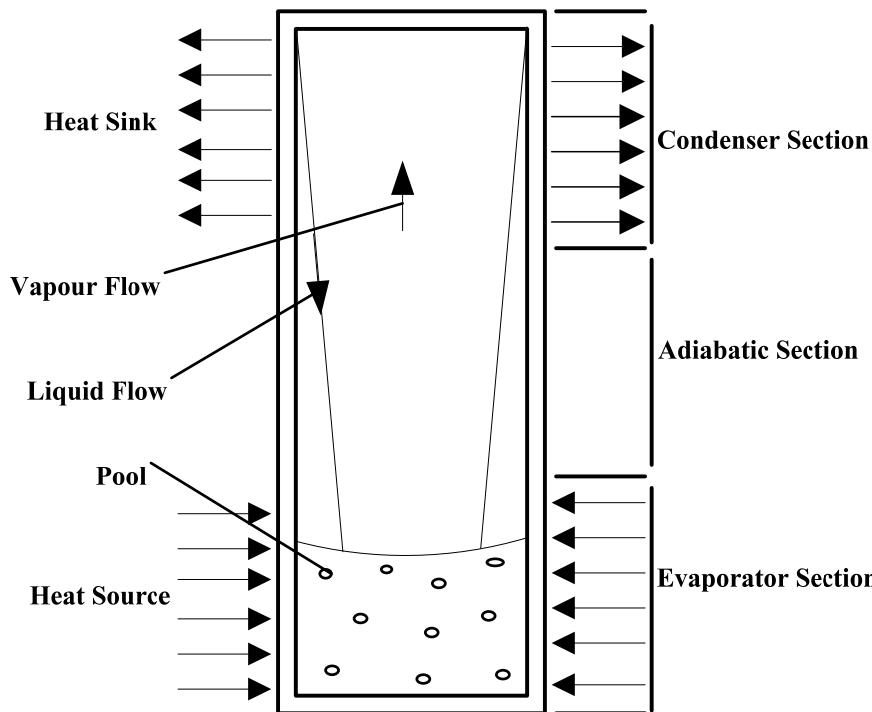
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129

TPCT is actually a wickless, gravity-assisted heat pipe with a small amount of working fluid which is in equilibrium with its own vapor sealed inside container (pipe wall and end caps). The length of the TPCT (similar to heat pipe) is divided into three parts: evaporator section, adiabatic section and condenser section, as shown in Figure 6.

130



131

132 Figure 6 Schematic of the TPCT [2]

133 **2. Heat transfer characteristics**

134 The heat transfer performance ($Q_{\text{Theoretical}}$) of a TPCT can be calculated from the ratio of
 135 temperature difference between the evaporator and condenser (ΔT) and the total thermal resistance
 136 (Z_{Total}) as shown in equation (1) [14].

137

$$Q_{\text{Theoretical}} = \frac{\Delta T}{Z_{\text{Total}}} \quad (1)$$

138 Thus ΔT define by (2);

139

$$\Delta T_{\ln} = \frac{(T_{h,\text{in}} - T_{c,\text{out}}) - (T_{h,\text{out}} - T_{c,\text{in}})}{\ln \frac{(T_{h,\text{in}} - T_{c,\text{out}})}{(T_{h,\text{out}} - T_{c,\text{in}})}} \quad (2)$$

140 The total thermal resistance can be explained based on the many factors which depend on
141 the function or properties of the operation. This is shown as the total resistance of the thermal model
142 by Engineering Sciences Data Unit Item No 80023, ESDU81038 [15].

143 **2.1. Thermal resistance of thermosyphon**

144 The TPCT is a simple heat transfer device allowing its performance to be clearly evaluated.
145 The principle method to calculate performance is the total resistance calculation method ESDU81038
146 [15] and can be explained as follows;

147 The resistance of a TPCT device can be split into 10 valves and divided into 3 groups. The
148 external resistance is the resistance from the outside wall of the TPCT at both the evaporator and
149 condenser sections. The internal resistance is the resistance from the phase change, pool boiling or
150 film boiling, when the vapour pressure drops along the pipe. The resistance from the material property
151 is dependent on the type of material used. Resistance is explained as follows in Figure 7 [2].

152 **2.2. External resistance (Z_1 and Z_9)**

153 Z_1 and Z_9 represent the resistance from the external convection of the pipe:

154
$$Z_1 = \frac{1}{h_{eo} A_{eo}} \quad (3)$$

155
$$Z_9 = \frac{1}{h_{co} A_{co}} \quad (4)$$

156 **2.3. Resistance from material property (Z_2 and Z_8)**

157 Z_2 and Z_8 represent the resistance from the thermal conductivity if the material is:

158
$$Z_2 = \frac{\ln(D_o / D_i)}{2\pi L_e k_x} \quad (5)$$

159
$$Z_8 = \frac{\ln(D_o/D_i)}{2\pi L_c k_x} \quad (6)$$

160 **2.4. Internal resistance ($Z_2, Z_3, Z_4, Z_5, Z_6, Z_7$ and Z_{10})**

161 Z_3 and Z_7 represent the internal resistance due to the working fluid of pool and film boiling
 162 and is devised into:

163 Z_{3p} is resistance from pool boiling; $\frac{1}{\Phi_3 g^{0.2} Q^{0.4} (\pi D_i L_e)^{0.6}}$ (7)

164 Z_{3f} is resistance from film boiling at the evaporator section; $\frac{CQ^{1/3}}{D_i^{4/3} g^{1/3} L_e \Phi_2^{4/3}}$ (8)

165 When $C = \left(\frac{1}{4}\right) \left(\frac{3}{\pi}\right)^{4/3} = 0.325$, Φ_2 is figure of merit (2); $\left(\frac{Lk_l \rho_l^2}{\mu_l}\right)^{0.25}$ and Φ_3 is figure of merit

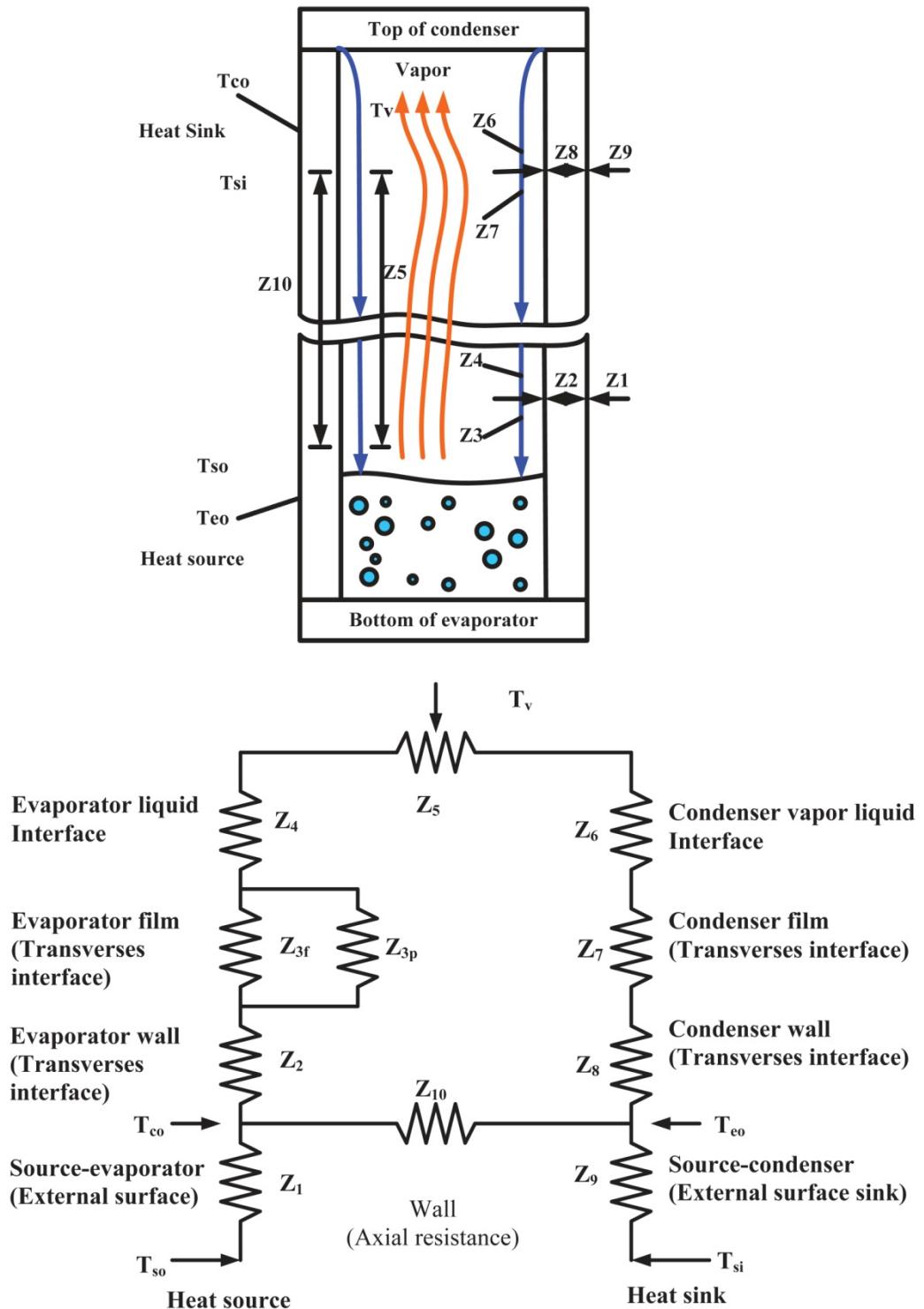
166 $0.325 \times \frac{\rho_l^{0.5} k_l^{0.3} C_{pl}^{0.7}}{\rho_v^{0.25} L^{0.4} \mu_l^{0.1}} \left[\frac{P_v}{P_a} \right]^{0.23}$. The condition using Z_{3p} and Z_{3f} is $Z_3 = Z_{3p}F + Z_{3f}(1-F)$ as

167 equation (9). F is filling ratio that is defined by; $\frac{V_1}{AL_e}$.

168 Z_7 is resistance from film boiling of the working fluid at the condenser section:

169 $Z_7 = \frac{CQ^{1/3}}{D_i^{4/3} g^{1/3} L_c \Phi_2^{4/3}} \quad (10)$

170



171

172

173 Figure 7 Model of total resistance of TPCT, ESDU 81038 [2]

174

175

176 Z_7 and Z_6 is resistance due to the phase change at the evaporator and condenser section
177 respectively. Z_5 is resistance due to pressure drop along the pipe. Z_{10} is resistance due to heat
178 conduction along the axial pipe. Normally, Z_4 , Z_5 , Z_7 and Z_{10} are of small valve and can be
179 negligible.

180 Thus Z_{Total} can be defined as;

$$181 Z_{Total} = Z_1 + \left[(Z_2 + Z_3 + Z_4 + Z_5 + Z_6 + Z_7 + Z_8)^{-1} + (Z_{10})^{-1} \right]^{-1} + Z_9 \quad (11)$$

182 **2.5. Heat transfer coefficients**

183 The heat transfer coefficient ($h_{coefficient}$) or film coefficient, in ALT/CV and in mechanics is
184 the proportionality constant reciprocal the thermal resistance (Z_{Total}) and the thermodynamic driving
185 force for the flow of heat [15-17]. The $h_{coefficient}$ can be evaluated using the following equation;

$$186 h_{coefficient} = \frac{1}{Z_{Total}} \quad (12)$$

187 where, the data on temperature wall distributions, vapour temperature and thermal load were
188 measured. The $h_{coefficient}$ has been referenced by many researchers [10-13]. Thus, this section focuses
189 on pool boiling dynamics in evaporator section of ALT/CV. The $h_{coefficient}$ is an enhancement thermal
190 performance in two phase transfer with nanofluids containing surfactant [18-20] under condition
191 shown in Table 3.

192 **2.6. Heat transfer rate analysis in prototye of oven an advanced loop thermosyphon
193 (OALT/CV)**

194 During the experiments, the variable parameter was controlled in order to calculate the heat
195 transfer rate characteristics of OALT/CV using the calorific method. The following equations were

196 used to calculate the actual heat transfer rate (Q_{act}) in equation (13), with maximum heat transfer rate
 197 (Q_{max}) in equation (14), and then for error analysis recently compiled by Paramatthanuwat et al., [3]
 198 as follows:

199
$$Q_{act} = \dot{m}C_p(T_{in} - T_{out}) \quad (13)$$

200 and ;

201
$$Q_{max} = \dot{m}C_p(T_e - T_c) \quad (14)$$

202 thus ;

203
$$Q = f\left(\dot{m}, T_e, T_c, T_{in}, T_{out}\right) \quad (15)$$

204 However, the ALT/CV and OALT/CV with a fin does not have a theoretical/mathematicical
 205 model to predict heat transfer rate at this point in time. Thus, the Q_{max} in equation (14) was compared
 206 with the theory about the heat rate of convection from the fin in equation (16) from Dewitt [21] as
 207 follows:

208
$$Q_{pre} = h \left[N\eta_f A_f + (A_t - N\eta_f A_f) \right] \theta_b = hA_t \left[1 - \frac{NA_f}{A_t} (1 - \eta_f) \right] \theta_b \quad (16)$$

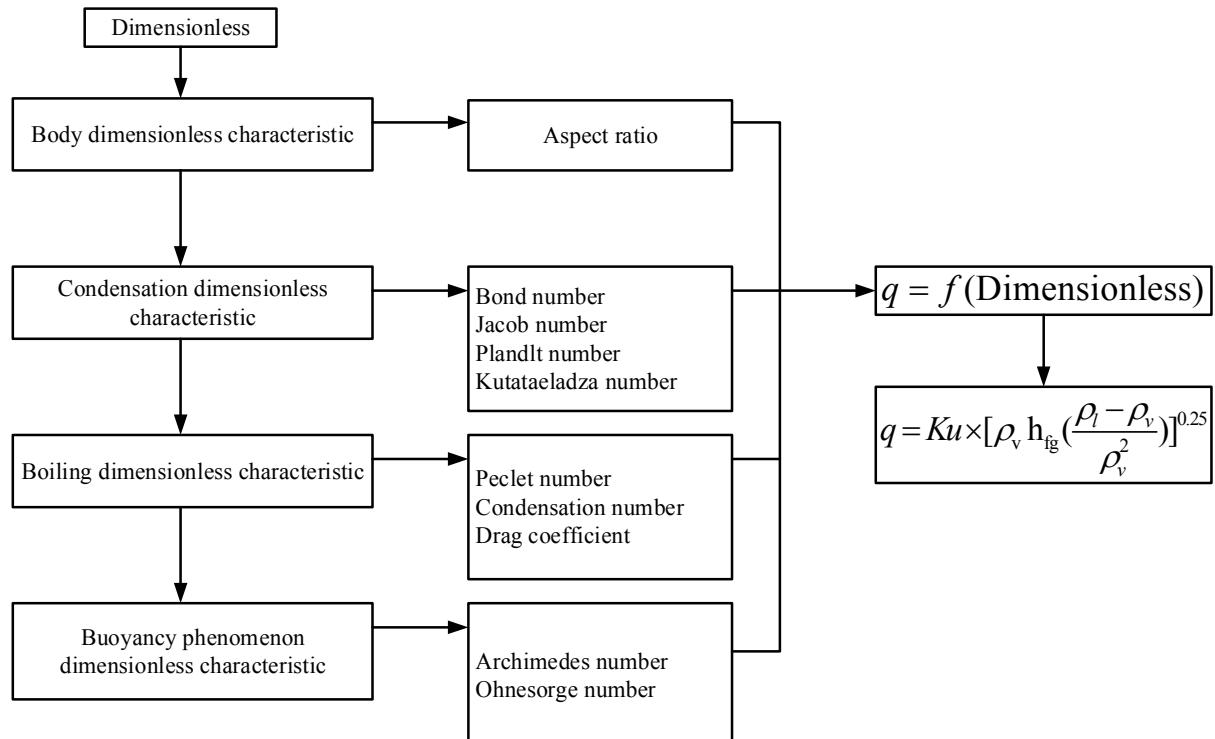
209 Every temperature inside the oven is associated with the convection transfer of OALT/CV.
 210 In particular, there is a similarity between the diffusion of heat and the heat transfer coefficient.
 211 Defining a heat transfer coefficient depends on the heat flux, that is heat flow per unit area and the
 212 thermodynamic driving force for the flow of heat convection in equation (17). Also, represented in
 213 equation (16), it is consistent with Newton's law of cooling and also following Incropera FP [22];

214
$$h = \frac{Q}{A} (T_{air} - T) \quad (17)$$

215 whereas the parameter convection coefficient in equation (16) was assumed to be equal to Equation
216 (17) and the efficiency of a single fin shown in Figure 41 (Chapter V). Moreover, the prime of surface
217 (A) in equation (16) was fixed to be equal to equation (17). Due to Equation (16) having a
218 rectangular geometry it is similar to the loop thermosyphons which have convection heat transfer. The
219 theorem equation (14) and equation (16) was similar. Due to the rectangular shape of the source
220 resembling the ALT/CV from Equation (16), it was determined from this theory.

221 **3. Dimensionless [2, 16, 23, 24]**

222 A dimensionless quantity is a quantity without an associated physical dimension. It can be
223 obtained from the relationship between a variable and a focused property for a particular phenomenon
224 which occurs inside a ALT/CV. A dimensionless term in ALT/CV system, in practice, is defined by a
225 fluids' behaviour, for example, flow of phase fluid, boiling point, condensation, exertion of buoyancy
226 force or etc [23, 25]. A dimensionless quantities study can be explained as follows into 4 groups in
227 Figure 8:



228

229 Figure 8 Dimensionless diagram

230 **3.1. Body dimensionless characteristic**

231 Aspect ratio represents the distance of physical motion for a working fluid (liquid and
 232 vapour). When the aspect ratio increased the heat transfer rate rose phenomenally. Therefore, the
 233 larger aspect ratio leads to pool boiling which occurs with the highest heat transfer rate. However, the
 234 phenomenon that approaches the inside of a confined channel has a lower heat transfer rate. The
 235 aspect ratio can be defined as the following equation (18):

236
$$\text{Aspect ratio} = \frac{L_e}{D_i} \quad (18)$$

237 **3.2. Boiling dimensionless characteristic**

238 Bond number (Bo) is the ratio of buoyancy force to the surface tension force. Bo can be
 239 used to explain boiling phenomenon inside an evaporator section and the state of the vapour bubbles
 240 nucleation. If the bond number reaches a threshold level for a particular liquid, then boiling occurs.
 241 Bo can be defined as the equation (19);

$$Bo = D_i [g \frac{\rho_l - \rho_v}{\sigma}]^{0.5} \quad (19)$$

243 Jacob number (Ja) is the ratio of latent heat to sensible heat of the working fluid. It represents
244 the ability of the phase change in the working fluid. When the working fluid's inner tube obtains heat,
245 then heat has been transferred by a condensation dye to the latent high heat: Therefore, it can be
246 transferred from one end of point to another end point with only a little change in temperature. The
247 denominator is the heat of liquid film. Jais defined by the equation (20);

$$248 \quad Ja = \frac{h_{fg}}{C_{pl}T_v} \quad (20)$$

249 Plandlt number (Pr) is the ratio of momentum diffusivity to the thermal diffusivity of liquid.
250 It represents convection heat transfer in a tube that occurs when the vapour bubble moves from the
251 evaporator section to the condenser section. As heat is collected, the vapour's heat transfer and density
252 of heat flux will be enhanced with a high Pr. It can be defined by the equation (21):

$$253 \quad \Pr = \frac{u_l C_{pl}}{k_l} \quad (21)$$

254 Note that if all the Bo, Ja and Pr have their valves lower than 1; there will be no occurrence
255 of phase change.

256 Kutataeladza number (Ku) is the ratio of the heat flux to the critical heat flux of fluid. Kuis
257 usually applied as the coefficient corresponding to the critical pool boiling and is defined as the ratio
258 of the critical heat flux and the properties of the working fluid. The critical pool boiling starts when
259 the boiling nucleation state is reached. This is when the vapour bubbles begin to spread over the
260 heating surface. In flow boiling, the boiling nucleation state and the after states may be also discussed
261 as the important heat transfer mechanisms; it may therefore be justified to apply Ku in an empirical
262 correlation for the flow boiling within a ALT/CV. It can be defined as follows (22):

263
$$Ku = \left[\frac{q}{[\rho_v h_{fg} (\frac{\rho_l - \rho_v}{\rho_v^2})]^{0.25}} \right] \quad (22)$$

264 **3.3. Condensation dimensionless characteristic**

265 Peclet number (Pe) is the ratio of bulk heat transfer rate to conductive heat transfer rate. It is
 266 defined to be the ratio of the rate of advection of a physical quantity of the flow to the rate of diffusion
 267 of the same quantity driven by an appropriate gradient. In other words, it is the transport of heat to
 268 condensation. There is also the extreme limit for the motion according to small and large Pe with the
 269 absence gravity. It was shown that when the Pe of the continuous fluid is small, that of the dispersed
 270 phase is high. The initial non equilibrium temperature distribution results in the thermal boundary
 271 layers that develop on the inner sides of the interfaces of the drops. A temperature variation of Pe
 272 along the interfaces is encountered. The induced thermo-motion is strongly unsteady and it leads to
 273 either the spontaneous coagulation of condensation in a cold fluid or to the evaporator section. Pe can
 274 be defined as follows (23):

275
$$Pe = \frac{LV\rho C_p}{k} \quad (23)$$

276 Condensation number (Co) is the liquid density ratio and hence the gravitational component
 277 and homogeneous theory for the momentum component (heat flux divided by the product of mass flux
 278 and latent heat of vaporization). The higher the value of Co the easier for the condensate to return to
 279 the evaporator section. Co can be defined as follows (24):

280
$$Co = \frac{h}{k} \left[\frac{\mu^2}{g\rho^2} \right]^{\frac{1}{3}} \quad (24)$$

281 Drag coefficient (Cd) is proportional to gravitational and internal forces that predict
 282 momentum heat transfer rates dependent on the physical motion. Cd is a dimensionless quantity that

283 is used to quantify the drag or resistance of an object in a fluid environment such as air or water. It is
284 used in the drag equation, where a lower Cd indicates the object will have less aerodynamic or
285 hydrodynamic drag. The drag coefficient is always associated with a particular surface area. Cd can
286 be defined as follows (25):

287

$$Cd = \frac{g(\rho - \rho_f)L}{\rho V^2} \quad (25)$$

288 **3.4. Buoyancy phenomenon dimensionless characteristic**

289 Archimedes number (Ar) determines the motion of fluid and solids due to density
290 differences. Ar is dependent on dimension. It predicts the boiling phenomenon approach of internal
291 boiling inside. Ar can be defined as follows (26):

292

$$Ar = \frac{g(\rho_s L^3)}{\mu^2} (\rho_s - \rho_l) \quad (26)$$

293 Ohnesorge number (Z) is proportional to viscous force and to inertial force with surface
294 tension. Z is generally used in momentum heat transfer rates and atomization. Z can be defined as
295 follows (27):

296

$$Z = \frac{\mu}{(g \rho L \sigma)^{\frac{1}{3}}} \quad (27)$$

297 The above-stated dimensionless numbers were correlated with Ku in the form of function
298 (28) to calculate the convection heat transfer capacity of one tube.

299

$$q = f(\text{Dimensionless}) \quad (28)$$

300 Thus:

301 q is heat flux, its equivalent Kutataeladza number (Ku) can be correlated in equation (29)

302
$$q = Ku \times [\rho_v h_{fg} \left(\frac{\rho_l - \rho_v}{\rho_v^2} \right)]^{0.25} \quad (29)$$

303 From Equation (28) and (29) the heat flux and the ALT/CV at a vertical position can be
304 evaluated. This will be shown in chapter 4.

305

306 **4. Nanofluids[2]**

307 A nanometer, 10^{-9} m, is about ten times the size of the smallest atom, such as hydrogen and
308 carbon. However, a micron is barely larger than the wavelength of visible light, thus invisible to the
309 human eye. A millimeter, the size of a pinhead, is roughly the smallest size available in present day
310 machines. The range of scales from millimeters to nanometers is one million, which is also
311 approximately the range of scales presently used in mechanical technology. From the largest
312 skyscrapers to the smallest conventional mechanical machine parts.

313 Nanofluids are solid-liquid composite materials consisting of solid nanoparticles or
314 nanofibers with typical sizes of 1-100 nm suspended in liquid. Nanofluids have attracted great interest
315 recently due to reports of greatly enhanced thermal properties. For example, Table 5 to Table 7 show
316 silver nanoparticle properties, a small amount (<1%volume fraction) of Ag nanoparticles or carbon
317 nanotubes dispersed in ethylene glycol or oil, is reported to increase the inherently poor thermal
318 conductivity of the liquid by 40% and 150%, respectively. Conventional particle-liquid suspensions
319 require higher concentrations (>10%) of particles to achieve such enhancement. However, problems
320 of theology and stability are amplified at high concentrations, precluding the widespread use of
321 conventional slurries as heat transfer fluids. In some cases, the observed enhancement in thermal
322 conductivity of a nanofluids in order of magnitude, were larger than predicted by well-established
323 theories. Other perplexing results in this rapidly evolving field include a surprisingly strong
324 temperature dependence of the thermal conductivity and a three-fold higher critical heat flux
325 compared with the base fluids [26].

326

327

328

329 Table 5 Silver nanoparticle properties [2]

General properties	
<u>Name, symbol, atomic number</u>	silver, Ag, 47
<u>Element category</u>	transition metals
<u>Standard atomic weight</u>	107.8682 g·mol ⁻¹
<u>Electron configuration</u>	4d10 5s1
<u>Electrons per shell</u>	2, 8, 18, 18, 1
Physical properties	
<u>Color</u>	silver
<u>Phase</u>	solid
<u>Density (near r.t.)</u>	10.49 g·cm ⁻³
<u>Melting point</u>	1234.93 K
<u>Boiling point</u>	2435 K
<u>Heat of fusion</u>	11.28 kJ·mol ⁻¹
<u>Heat of vaporization</u>	250.58 kJ·mol ⁻¹
<u>Specific heat capacity</u>	25.350 J·mol ⁻¹ ·K ⁻¹ (25°C)

331 Table 6 Vapour pressure of silver nanoparticles [2]

Vapour pressure						
P(Pa)	1	10	100	1 k	10 k	100 k
at T(K)	1283	1413	1575	1782	2055	2433

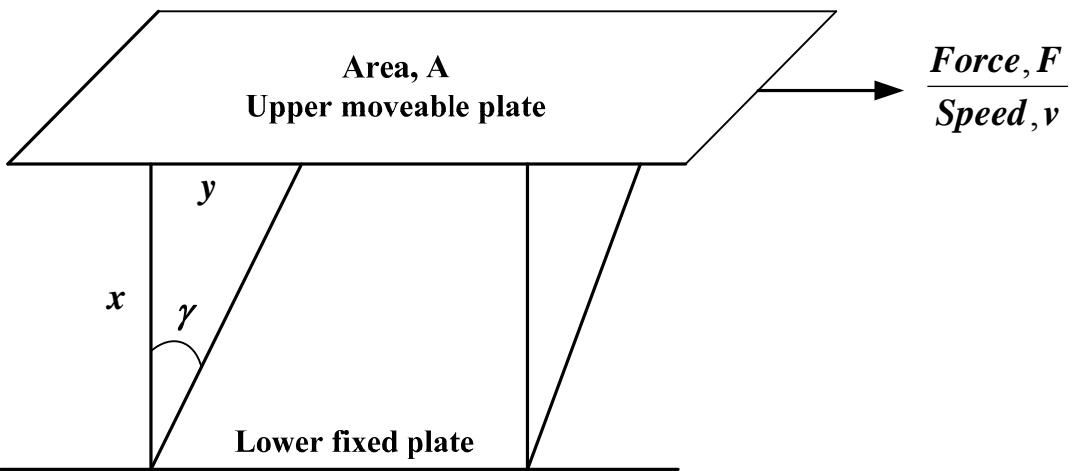
332 Table 7 Silver atomic properties [2]

Atomic properties	
Crystal structure	face-centered cubic
Crystal structure	1, 2, 3 (amphoteric oxide)
Electronegativity	1.93 (Pauling scale)
Ionization energies	1st: 731.0 kJ/mol 2nd: 2070 kJ/mol 3rd: 3361 kJ/mol
Atomic radius	144 pm
Covalent radius	145±5pm
Van der Waals radius	172 pm
Miscellaneous	
Magnetic ordering	diamagnetic
Electrical resistivity	15.87 nΩ·m(20°C)
Thermal conductivity	429 W·m ⁻¹ ·K ⁻¹ (300K)
Thermal diffusivity	174 mm ² /s(300 K)
Thermal expansion	18.9 μm·m ⁻¹ ·K ⁻¹ (25°C)
Speed of sound (thin rod)	2680 m·s ⁻¹ (r.t.)

333 **4.1. Rheological properties of nanofluids [2, 27]**

334 Rheological properties have a very important role in fluids flow. During application,
 335 nanofluids are likely to flow either by forced or natural convection and the flow properties such as
 336 viscosity are therefore, the essential to the study of suspensions containing particles the size of
 337 nanofluids. Rheology is defined as the study of the deformation of flow materials. When force is

338 applied to a liquid, the liquid will flow to relieve the strain from this force. Different systems will
 339 resist this flow more than others, so this resistance is the measurement of the viscosity of the system
 340 [28, 29]. Newton first introduced a basic model to show the flow measurement of the liquid between
 341 two parallel plates as shown in Figure 9



342

343 Figure 9 Rheology diagram analysis [29]

344 Using this model a number of common Rheology terms can be defined as follows:

345 Shear Stress (SS), the force experienced by the liquids is proportional to the area of the
 346 upper plate. SS is defined by the equation below;

347

$$\sigma = \frac{F}{A} \quad (30)$$

348 When: σ is shear stress, Nm^{-2} or Pa . F is force, $m \cdot s^{-1}$. And A is area upper moveable plate, m^2 .

349 Shear Rate (SR), the velocity gradient or the rate of change of velocity at which one layer
 350 passes over an adjacent layer. SR is defined by equation below:

351

$$\gamma = \frac{dv}{dx} \quad (31)$$

352 When: γ is shear rate, s^{-1} . dv is the velocity of the moving plate, $m \cdot s^{-1}$. dx is the distance
353 between the two parallel plates, m .

354 Shear Viscosity (SV) is expressed mathematically as,

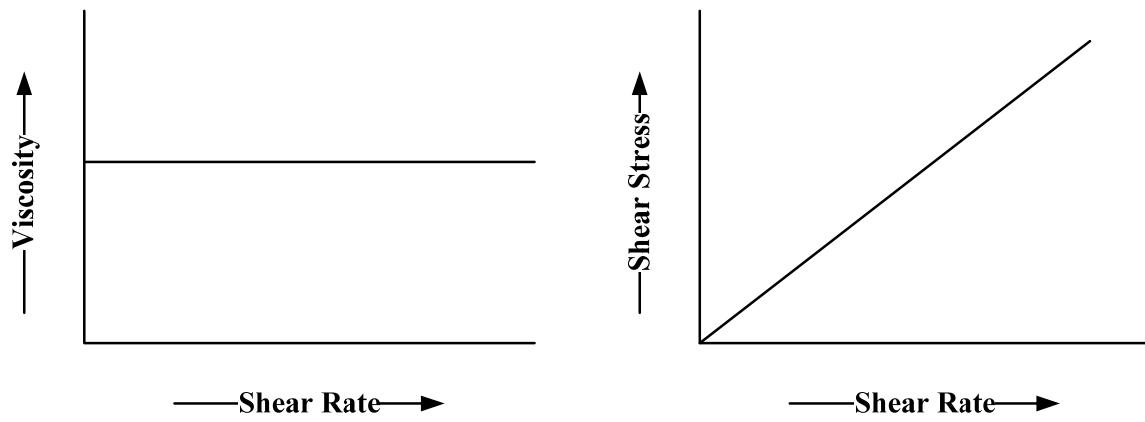
355
$$\eta = \frac{SS}{SV} \quad (32)$$

356 The measurement was the resistance to the flow of liquid. Pascal.second ($Pa.s$) is the basic
357 unit of SV but poise or centerpoise (cp is one hundredth of a Poise) is often used and one cp is
358 equivalent to a millpascal second, $mPa.s$. When quoting SV the SR (measurement method/equipment
359 used) should be stated together with the temperature at which the measurement was taken [28].

360 All materials that show flow behaviour are referred to as fluid. In all fluids, there is a
361 frictional force between the molecules and, therefore display, certain flow resistance which can be
362 measured as a viscosity. The SV is a transporting property which refers to the resistance of material
363 flow. When dealing with nanofluids, we are tempted to consider the dispersed medium under
364 question, as a homogeneous fluid characterized by the properties such as density and viscosity. This in
365 turn, will only require a single set of mass and momentum conservation equation. However, such a
366 simple picture will not provide a useful enough case where the fluid is unsteady and non-uniform
367 [29].

368 **4.2. Newtonian flow behaviour [2, 27]**

369 Isaac Newton found that the shear force acting on a liquid is proportional to the resulting
370 flow velocity. Hence, a fluid is said to be Newtonian if the viscosity remains constant with an increase
371 in SR . Newtonian flow behaviour is observed in flow molecular liquids such as water, mineral oil
372 (without polymer additives) and solvents. However, more complex flow behaviour is expected for
373 fluid containing suspended particles [30]. Newtonian can be showed as in Figure 10.



374

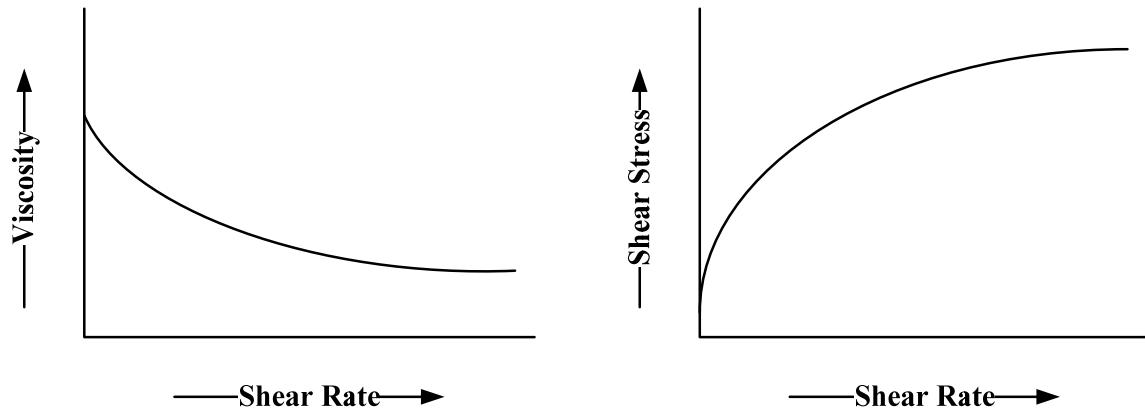
375 Figure 10 Newtonian fluid flow behaviour diagrams[28, 29]

376 **4.3. Non-Newtonian flow behaviour [2, 27]**

377 Fluids, whose SV changes with an increase in SR, are referred to as Non-Newtonian. These
378 fluids could be further classified according to their flow behaviour. Shear-thinning and shear
379 thickening flow behaviour is discussed in the following sections.

380 **4.4. Shear-thinning flow behaviour [2, 27]**

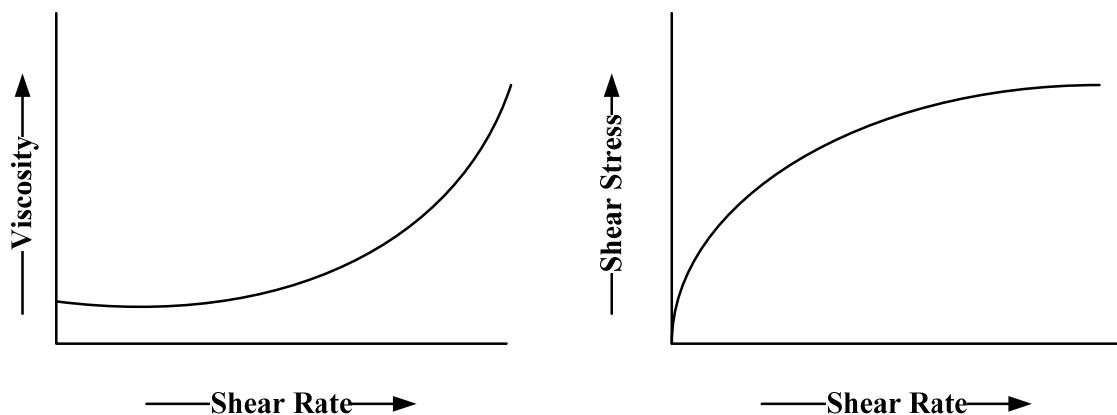
381 Shear-thinning or Pseudoplasticity, for samples that display shear-thinning behaviour, the
382 SV is dependent on the degree of shear load (SL). Thus, the viscosity decreases with an increase in
383 SS. In dispersions, shearing can cause the particles to change the flow direction and also the direction
384 of the flow gradient. This can lead to disintegration of agglomerates or change in particle form. The
385 interaction forces between particles may decrease during the process and cause a lowering in the flow
386 resistance [30]. Shear thinning as shown in the Figure 11.



387
388 Figure 11 Shear-thinning flow behaviour diagrams[28, 29]

389 **4.5. Shear-thickening flow behaviour [2, 27]**

390 Shear-thickening is similar to shear-thinning fluids; the SV of sample displaying shear-
391 thickening behaviour is also dependent on the degree of SL. Thus, the viscosity decreases but
392 increases in SS. With highly concentrated suspension, the probability of particle interaction is much
393 higher and may result in particles becoming wedged together thus increasing flow resistance. The
394 particle shape plays an important role as, during the shearing process, the particle moves and rotates.
395 For example, cube-shaped particles add more volume when rotating than spherical particles. Hence
396 less free space is available for the liquid between the particles [28, 29]. Shear-thickening as shown in
397 Figure 12.



398
399 Figure 12 Shear-thickening diagram[28, 29]

400 **4.6. Yield point [2, 27]**

401 The yield point or yield stress refers to an external force required before a material will start
402 to flow. A typical example is toothpaste; a certain amount of force must be applied before the
403 toothpaste starts to flow [29]. The materials with yield point tend to flow in homogeneously.

404

405 **4.7. The rheological properties analysis [2, 27]**

406 This section describes the analysis model function; Newtonian and Non-Newtonian.

407 Newtonian behaviour flow behaviour is described formally using Newton's law and is
408 defined as follows [29, 30]:

409
$$\sigma = \gamma \cdot \eta \quad (33)$$

410 When: σ is shear stress, Nm^{-2} or Pa . γ is shear rate, s^{-1} . η is viscosity, Pa.s .

411 Non-Newtonian behaviour for shear thinning and shear thickening flow behaviour, three
412 model functions are required for flow without a yield point. The curved model function can be defined
413 as follows [28, 30]:

414
$$\sigma = c \cdot \eta^p \quad (34)$$

415 When: σ is shear stress, Nm^{-2} or Pa . c is the flow coefficient (or power law index) Pa.s . p
416 is the exponent, the following applies:

417 $p < 1$ for shear thinning.

418 $p > 1$ for shear thickening.

419 $p = 1$ for Newtonian behaviour.

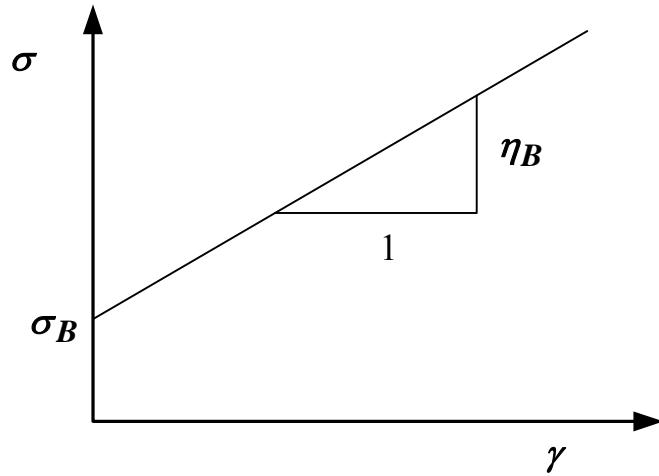
420 **4.8. The yield point flow curve analysis [2, 27]**

421 The Bingham flow model can be defined as follows:

422
$$\sigma = \sigma_B + \eta_B \cdot \gamma \quad (35)$$

423 When σ_B is the Bingham yield point. η_B is the Bingham flow coefficient as shown below in

424 Figure 13



425

426 Figure 13 Flow curve according to Bingham

427 The viscosity of nanofluids can also be estimated with well known formulae. Nevertheless, a
428 more reliable and direct way to calculate the viscosity is through experimental investigation [31].

429 **4.9. Thermal conductivity of nanofluids [2, 27]**

430 To use the Fourier's law, thermal conductivity of the material must be known. This
431 property, which is referred to as a transport property, provides an indication of the rate at which
432 energy is transferred by a diffusion process. This is dependent on the physical structure of: matter,
433 atomic and molecular and is considered to be the state of the matter. This section considers various

434 forms of matter identifying important aspects of behaviour and applying them to nanofluid thermal
435 property valves. Fourier's Law can be defined as [32]:

436
$$k = -\frac{\dot{q}_x}{\partial T / \partial x} \quad (36)$$

437 When: k is thermal conductivity, $\text{W} \cdot (\text{m} \cdot \text{K})^{-1}$. \dot{q}_x is heat flux, $\text{W} \cdot \text{m}^{-2}$. T is temperature, K .

438 Thermal conductivity shows a key role in the enhancement of the heat transfer performance
439 of a heat transfer fluid. Since the thermal conductivity metals are much higher than that of fluids, the
440 suspension containing ultra-fine metal particles are expected to show improved heat transfer
441 properties. It was found that the thermal conductivity of a nanofluid affected the volume fraction, the
442 size and the shape of the nanoparticle suspension in the liquid, as well as the distribution of the
443 dispersed particle [23, 33].

444 **4.10.Theory of nanofluids thermal conductivity [2, 27]**

445 As the thermal conductivity of nanofluids theory is non-existent, scientists have used an
446 existing model for estimating this theory. The Maxwell model was developed to explain the heat
447 transfer characteristics of larger particles in nanofluids research. This Maxwell model has served as a
448 foundation in the development and explanation of a much higher conductivity increase observed in
449 nanofluids. The effective thermal conductivity (k_{eff}) can be defined by [34]:

450
$$k_{\text{eff}} = \frac{k_p + 2k_l + 2(k_p - k_l)\phi}{k_p + 2k_l - (k_p - k_l)} k_l \quad (37)$$

451 When: k_p is thermal conductivity of particle. k_l is thermal conductivity of liquid. ϕ is volume
452 fraction of suspension.

453 Thus, the ϕ can be defined by;

454
$$\phi = \frac{v_p}{v_p + v_l} \quad (38)$$

455 When: v_p is volume of nanoparticles in fluid. v_l is defined as volume of base fluid.

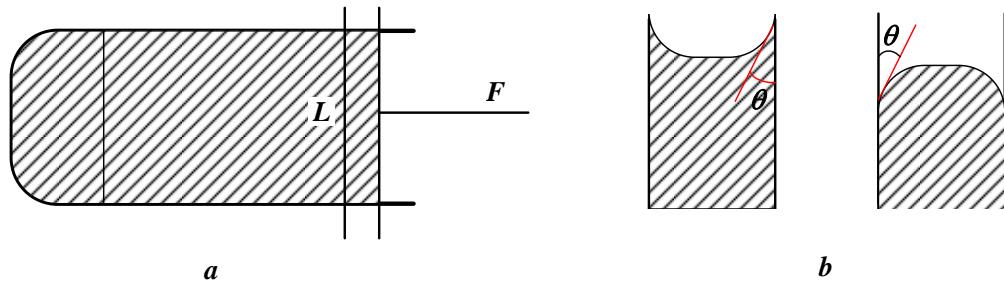
456 The thermal conductivity plays a role in the heat transfer enhancement of fluids. It is higher
457 in the solid metals than in liquids. The contained nanoparticles in suspension are expected to improve
458 the thermal properties. Consequently, the Maxwell model shows that the effective thermal
459 conductivity of fluids containing ultra-fine particles, depend on the thermal conductivity of the
460 spherical particles, the base liquid and the volume fraction of the solid particles [23, 34].

461 **4.11.The contact angle [2, 27]**

462 The contact angle is the angle of liquid or vapour which meets the interface on a solid
463 surface. It is this specific characteristic of most systems to determine the interactions across the three
464 interfaces. The contact angle of a small liquid droplet which is resting on a flat horizontal solid
465 surface shows the role of a boundary condition. It is measured by using a contact angle goniometry. It
466 does not have a high boundary limitation on a liquid or a vapour interface, thus it can be equally
467 applied between two liquids or two vapours. In the case of a molecule at or near the surface of a
468 liquid, the attracting forces will no longer balance out and the molecule will experience a resultant
469 force inwards. Due to this effect, the liquid will tend to take up a shape with a minimum surface area
470 [14]. In the case of a free-falling drop in a vacuum, this would become a sphere. Due to this
471 spontaneous tendency to contract, a liquid surface behaves like a rubber membrane under tension. In
472 order to increase the surface area, this work must be done on the liquid. The energy associated with
473 this work is known as the free surface energy. The corresponding free surface energy per unit surface
474 area is given the symbol, σ ; Figure 14 which shows the surface tension and the pressure difference
475 across a curved surface. In Figure 14a, the area is increased by moving one side a distance dx and the
476 work done is equal to Fdx , hence the increased energy is $2\sigma_l / dx$.

477 Factor 2 arises since the film has two free surfaces. Hence, if T is the force per unit length
478 for each of the two surfaces $2T_l dx = 2\sigma_l / dx$ or $T = \sigma_l$ This force per unit length is known as the
479 surface tension. It is numerically equal to the surface energy per unit area measured in any consistent
480 set of units, e.g. N/m. Since the latent heat of vapourization, L , is a measurement of the forces of
481 attraction between the molecules of a liquid we might expect surface energy or surface tension σ_l to
482 be related to L . This is found to be the case. Solids will also have a free surface energy and, in the
483 magnitude, it is found to be similar to the value for the same material in the molten state.

484 Figure 14b shows that a liquid is in contact with a solid surface. Molecules in the liquid
485 adjacent to the solid will experience forces from the molecules of the solid, in addition to the forces
486 from other molecules in the liquid. Depending on whether these solid/liquid forces are attractive or
487 repulsive, the liquid/solid surface will curve upwards or downwards.



488

489 Figure 14 Surface tension and pressure difference across a curved surface [14]

490 For example, two attractive and repulsive forces are water and mercury. The attractive
491 forces are the forces when the liquid is wetter than the solid. The angle of contact made by the liquid
492 surface with the solid is known as a contact angle, θ . For wetting, θ will lie between 0 and $\frac{\pi}{2}$ and for
493 non-wetting liquid, $0 > \frac{\pi}{2}$, as indicated in Figure 15 Wetting and non-wetting contact angle [14]

494 The wetting condition occurs when total surface energy is reduced by wetting;

495 $\sigma_{sl} + \sigma_{lv} < \sigma_{sv}$ (39)

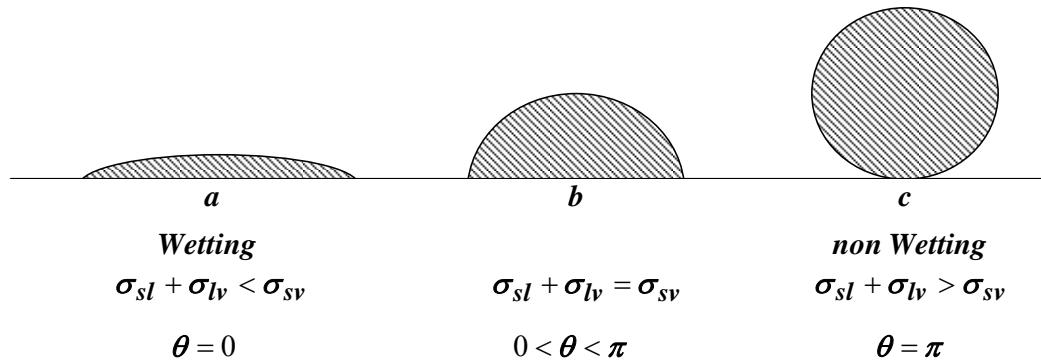
496 The non-wetting will not occur is defined as;

497 $\sigma_{sl} + \sigma_{lv} > \sigma_{sv}$ (40)

498 The intermediate condition of partial wetting is defined as;

499 $\sigma_{sl} + \sigma_{lv} = \sigma_{sv}$ (41)

500 When: s is solid, l is liquid. And v is vapour.

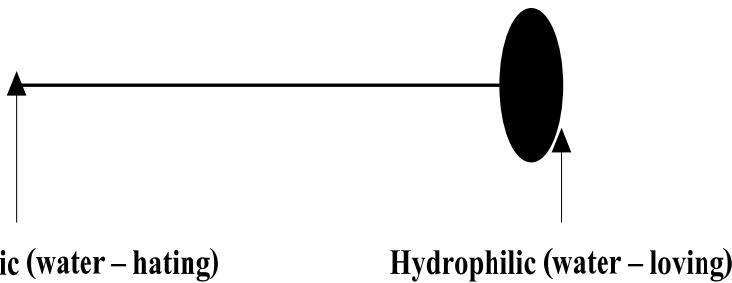


502 Figure 15 Wetting and non-wetting contact angle [14]

503

504 **5. Surfactant [2]**

505 Etymology - The surfactant term is a blend of surface active agent. Surfactants are usually
 506 organic compounds that are amphiphilic, meaning they contain both hydrophobic groups (their "tails")
 507 and hydrophilic groups (their "heads"). Therefore, they are soluble in both organic solvents and water.
 508 The surfactant term was coined by Antara products in 1950. In Index Medicus and the United States
 509 National Library of Medicine, "surfactant" is reserved for the meaning pulmonary surfactant. For the
 510 more general meaning, "surface active agent" is the heading as shown in Figure 16.

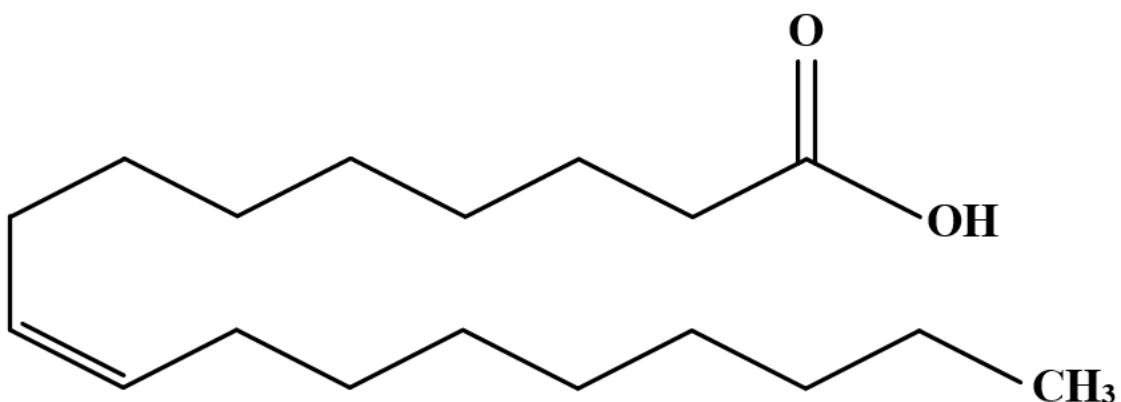


511

512 Figure 16 Surfactant[2]

513 Properties - Surfactants reduce the surface tension of water by adsorbing at the liquid-gas
514 interface. They also reduce the interfacial tension between oil and water by adsorbing at the liquid-
515 liquid interface. Many surfactants can also assemble in the bulk solution into aggregates. Examples of
516 such aggregates are vesicles and micelles. The concentration at which surfactants begin to form
517 micelles, is known as the critical micelle concentration or CMC. When micelles form in water, their
518 tails form a core that can encapsulate an oil droplet and, their (ionic/polar) heads form an outer shell
519 that maintains favorable contact with water. When surfactants assemble in oil, the aggregate is
520 referred to as a reverse micelle. In a reverse micelle, the heads are in the core and the tails maintain
521 favorable contact with the oil. Surfactants often classified into four primary groups; anionic, cationic,
522 non-ionic and zwitterionic (dual charge). Thermodynamics of the surfactant systems are of great
523 importance, theoretically and practically. This is because surfactant systems represent systems
524 between ordered and disordered states of matter. Surfactant solutions may contain an ordered phase
525 (micelles) and a disordered phase (free surfactant molecules and/or ions in the solution). Ordinary
526 washing up (dish washing) detergent for example, will promote water penetration in soil, but the
527 effect would only last a few days (many standard laundry detergent powders contain levels of
528 chemicals such as sodium and boron, which can be damaging to plants and should not be applied to
529 soils). Commercial soil wetting agents will continue to work for a considerable period, but they will
530 eventually be degraded by soil micro-organisms. Some can however, interfere with the life-cycles of
531 some aquatic organisms, so care should be taken to prevent run-off of these products into streams and
532 excess products should not be washed down.

533 Oleic acid - Oleic acid (Figure 17 and Table 8) is a mono-unsaturated omega - 9 fatty acid
534 found in various animal and vegetable sources. It has the formula $\text{CH}_3(\text{CH}_2)^7\text{CH}=\text{CH}$
535 $(\text{CH}_2)^7\text{COOH}$. The trans-isomer of oleic acid is called elaidic acid. The term oleic means related to,
536 or derived from, oil or olive.



537
538 Figure 17 Chain of Oleic acids [2]

539 Table 8 Other names of oleic properties [2]

Oleic properties	
(9Z)-Octadecenoic acid	
(Z)-Octadec-9-enoic acid	
cis-9-Octadecenoic acid	
cis-Δ9-Octadecenoic acid	
Oleic acid	18:1cis-9
Molecular formula	C ₁₈ H ₃₄ O ₂
Molar mass	282.4614 g/mol
Density	0.895 g/mL
Melting point	13-14°C (286 K)
Boiling point	360°C (633 K) (760 mmHg)
Solubility in water	Insoluble
Solubility in methanol	Soluble

540

CHAPTER III SILVER NANOFLOIDS PROPERTIES

541 This chapter describes the results and discussions of silver nanofluids properties when
542 preparing for filled as working fluids in ALT/CV. It is divided into a number of sections to determine
543 the thermal properties of silver nanofluids containing surfactant, as follows:

544 **1. Literature review [35]**

545 Cooling is one of the most important challenges facing numerous industrial sectors. Despite
546 the considerable amount of research and development focusing on industrial heat transfer
547 requirements, major improvements in cooling capability are still insufficient because conventional
548 heat transfer fluids possess poor heat transfer properties. Nanofluids, which are engineered by
549 suspending ultrafine metallic or non-metallic particles of nanometer dimensions in traditional cooling
550 fluids, have shown great enhancement in thermal conductivity and convective heat transfer
551 coefficient [2, 11, 36, 37].

552 This section also contains the literature review for thermal properties. Many researchers
553 have discussed the thermal properties points on which this study is based, together with the
554 background of the research and explanations of the problems faced. This study highlights the theories
555 and experiment for investigating the characteristics of thermal properties. Points of importance will be
556 emphasized, with significance given to the properties of nanofluids and surfactants and their use in
557 this experiment. Also included is the explanation of the characteristics of nanofluid behaviour in silver
558 nanofluids containing surfactant. Thus, the researchers used different methods depending on the base
559 fluids, nanofluid/nanoparticle type, etc. Recently, Thermal conductivity of 0.1 to 0.4% volume
560 concentration silver (Ag) nanoparticles in water were investigated. The nanofluids were formulated
561 using the ultrasonic vibration method for 3 hours and thermal conductivity enhancement showed 10%
562 at 0.4% of concentration [38]. Using a different method, the synthesis of silver nanofluids was
563 performed using high-pressure homogenization with a volume fraction 0.1 to 0.3% in water. The
564 highest thermal conductivity of the nanofluids showed an 18% increase at the concentration of 0.3%

565 [39]. Moreover, regarding the difference in base fluid, Ag nanofluids in toluene have shown 9%
566 thermal conductivity enhancement with a very low loading of 1.10-3 vol% [40-43]. Consequently,
567 nanofluids show better cooling capacity with respect to water in conventional heat pipes since
568 nanoparticles can flatten the temperature gradient of the fluids and reduce the boiling limit [36, 40,
569 41]. In addition, the concentration of nanofluids may affect the enhancement of thermal conductivity.
570 The studied silver nanofluids in ethylene glycol (EG) with 10,000 ppm concentrations showed 18%
571 thermal conductivity enhancement [44]. Then, the investigated carbon black (CB) in deionized water
572 with sodium dodecylsulfate (SDS) as well as Ag nanoparticles in silicon oil with oleic acid (OA), and
573 with the maximum enhancement of thermal conductivity compared to the base liquid, was 9% for the
574 wt% of the carbon black (CB) nanofluids and the wt% of the Ag nanofluids respectively [45]. The .1
575 wt% copper (Cu) aqueous nanofluids with 0.14 wt% of sodium dodecylbenzene sulfonate (SDBS) as
576 surfactant can generate maximum thermal conductivity enhancement up to 10.7% [46].

577 The rheological behaviour of nanofluids is essential in establishing adequate application and
578 design of processing. The 8 wt% titania nanoparticles in the EG showed Newtonian behaviour at a
579 low shear rate, and the shear viscosity was strongly dependent on the temperature and concentration
580 of the nanoparticles [47]. Then, the studied 1 vol% silver NP in ethanol with polyvinylpyrrolidone
581 (PVP) was stabilized [48]. The rheological results suggest that the PVP helped to decrease the
582 nanoparticle's size, resulting in low fluid viscosity and Newtonian fluid behaviour but remarkably
583 high thermal conductivity. Meanwhile, the 4.38 vol% silver nanofluids in the diethylene glycol (DEG)
584 showed Newtonian behavior at high viscosity [49]. However, different literature data have shown that
585 nanofluids have non-Newtonian behavior, particularly at a low shear rates. The most important
586 influence could be the effective particle concentration, the range of shear rate, and the viscosity of the
587 base liquid [48]. Then, it was found that the TiO₂ nanoparticle in the EG exhibited shear thinning
588 behavior when the particle concentration was higher than ~2% [47]. The investigated shear thinning
589 behaviour was 3% γ -Al₂O₃ and 10% TiO₂ in water [50]. Another main reason for the non-Newtonian
590 behavior could be the aggregation of nanoparticles in the nanofluids. Lu reported that physical
591 properties may change when the surfactant affects surface tension and viscosity. For instance, Al₂O₃

592 in water, at a 1:10 weight ratio with ammonium poly (PMAA-NH4), has demonstrated shear thinning
593 behaviour (a decrease in viscosity with an increased shear stress rate), which yields a good dispersion
594 rate when using PMAA suspension up to 47.5 %vol [51]. Then it was found that the 4 vol% of γ -
595 Al_2O_3 , TiO_2 , and CuO nanofluids with 0.5 wt% of carboxymethyl cellulose (CMC) in deionized water
596 containing up to 4 vol% of particle concentration showed non-Newtonian behaviour with shear
597 thinning [52].

598 In this paper, 0.5 wt% silver nanoparticle-based aqueous nanofluids with oleic acid (OA)
599 and potassium oleate surfactant (OAK^+) as surfactant were prepared by sonicating in water bath with
600 a cooling technique for a period time of 12 hours. The effect of the additive concentration on the
601 thermal properties was studied experimentally (Thermal conductivity, specific heat, density, viscosity,
602 contact angle, and application of thermal enhancement), and the rheological behaviour (the correlation
603 between shear stress and shear rate) was investigated experimentally and theoretically. Moreover, the
604 heat enhancement cooling of the fluid (HEC) was investigated experimentally and it was confirmed
605 that nanofluids/nanofluids containing surfactant could be used in the application of heat transfer. The
606 methods of the experiment are briefly explained in section 2. Section 3 shows the experimental results
607 and offers a discussion. The conclusions to the study are in section 4.

608 **2. Materials and methods [35]**

609 **2.1. Nanofluids and thermal property study**

610 Figure 18 shows a schematic diagram of the preparation the nanofluids. Water-based silver
611 nanofluids were formulated with dry silver nanoparticles (Sigma-Aldrich, USA), OA, and OAK^+
612 (Sigma-Aldrich, USA) by using a two-step method [45, 53]. The 0.5, 1, and 1.5 wt% of OA and OAK^+
613 were added to the 0.5 wt% silver nanofluids, which showed controlled and variable parameters as
614 seen in Table 1. After sonicating for 12 hours with a cooling technique, the particle size was measured
615 using a nano-size particle analyzer (ZEN 3600 MALVERN, USA) in the range between 0.6 nm and
616 6.0 μm . The thermal properties of the nanofluids were measured using the hot-wire method (PSL
617 Systemtechnik GmbH) from 20°C to 80°C. The rheological characteristics of the NF were analyzed

618 using a Rheo-microscope Physica MCR301 (Anton Paar GmbH). The measurements were based on
 619 the controlled shear stress model with the stress ranging from 0.05 to 5 Pa. The maximum uncertainty
 620 was found to be 1.7% [2, 53, 54].

621 The rheological behaviour of the NF containing OAK^+ can be expressed with the power
 622 model in Eq. (42) with the viscosity as following the power law model indices less than $n \leq 1$.

623
$$\eta = K \gamma^{n-1} \quad (42)$$

624 In Eq. (42), η is the apparent viscosity, γ is the shear rate, K is the consistency index, and n
 625 is the power law index. The power law index of the nanofluids decreases with increasing
 626 nanoparticles concentration, and increases with increasing temperature [52]. Apparently, the viscosity
 627 of the NF decreases as the shear rate increases. Thermal conductivity

628 The idea of thermal conductivity is non-existent in nanofluids theory scientists have used an
 629 existing model for estimated. The Maxwell model was developed to explain the heat transfer
 630 characteristics of larger particles in nanofluids research. This model has served as a foundation in the
 631 development and explanation of the much higher conductivity increase observed in nanofluids. The
 632 effective thermal conductivity (k_{eff}) can be defined by the following [34, 55]:

633
$$k_{eff} = \frac{k_p + 2k_l + 2(k_p - k_l)\phi}{k_p + 2k_l - (k_p - k_l)\phi} k_l \quad (43)$$

634 Thus, the ϕ can be defined by:

635
$$\phi = \frac{V_p}{V_p + V_l} \quad (44)$$

636 **2.2. Contact angle instrument**

637 In order to measure the contact angle of the sample fluids, the valves are required to be at
638 room temperature. The temperature was controlled with a precision of $\pm 1^\circ\text{C}$. In this study, the drops
639 of fluids were measured using a Contact Angle Meter Model: DM-CE1; Kyowa Interface Science.
640 The accuracy was $\pm 0.5^\circ$ (repeatability described in standard deviation). The following liquids were
641 used in the experiment: a copper plate with a diameter of 60 mm and a thickness of 0.3 mm were used
642 as a test surface. A droplet of nanofluids was generated at a very low rate (1 $\mu\text{l/s}$) and detached from
643 the syringe needle tip as soon as it touched the copper plate. Consecutive photographs were used to
644 measure the contact angles. The spatial resolution was estimated to be about 50 μm on the basis of the
645 focused area and camera pixel size. A video was taken while the droplet was spreading over the
646 copper plate from initial contact to equilibrium position. The temporal resolution was estimated based
647 on the frame speed of the CCD camera at 30 fps. For each concentration, three experiments were
648 performed and the average was ascertained. The measurement settings were then adjusted and the
649 software was initialized [16, 56].

650 **2.3. Specific heat**

651 The dynamic of specific heat was applied in the experiment. This was according to
652 Rajabpour et al., [57] regarding the application of the theory model to nanofluids.

653
$$C_{p,\text{nf}} = \phi C_{p,n} + (1 - \phi) C_{p,bf} \quad (45)$$

654 This second model has served as a foundation in the development and explanation of the
655 much higher specific heat observed in nanofluids from nanoparticles. The effective specific heat
656 ($C_{p,\text{eff}}$) can be defined as follows [58]:

657
$$C_{p,\text{eff}} = \frac{\phi(\rho C_p)_n + (1 - \phi)(\rho C_p)_{bf}}{\phi\rho_n + (1 - \phi)_{bf}} \quad (46)$$

658 The measurement uncertainty for the specific heat was calculated by propagating the
659 precision uncertainties of all the individual measurements required to determine the specific heat in
660 Equation (46) and can be defined as follows [58]:

661

$$\mu_{total} = \sqrt{\sum_i^n \left(\frac{\partial C_{p,sample}}{\partial x_i} \mu_i \right)^2} \quad (47)$$

662 Equation (46) and (47) should be noted that for Newtonian nanofluids. For the non-
663 Newtonian fluid, the variation of the rheology does not depend on direct models but on the volume
664 fractions of the nanoparticles [59, 60].

665 **2.4. The heat transfer enhancement**

666 Figure 19 shows a schematic diagram of the experimental apparatus, which consists of the
667 heat enhancement cooling of the fluid (HEC) and peripheral devices. The heat enhancement cylinder
668 was made from Stan less steel (AISI 304) with a diameter and height of 1,500 mm and 3,000 mm. The
669 HEC was the heat source from the Stan less heater (2,000 Watt) with a diameter and height of 500
670 mm and 2,000 mm. The heat was supplied by circulating the Stan less through to HEC through to
671 20%, 40%, 60%, 80%, and 100% respectively of the heatsource. The cooling and pre-cooling section
672 was heat sink from a cold bath. The cooling fluids are shown in

673 Table 2. Eighteen thermocouples were connected through a data logger (Yokogawa DX200 with
 674 $\pm 0.1^\circ\text{C}$ accuracy, 20 channel input, and -200°C to $1,100^\circ\text{C}$ measurement temperature range). Type K
 675 thermocouples (OMEGA with $\pm 0.1^\circ\text{C}$ accuracy) were attached to the inlet, the outlet, and the surface
 676 of the heating and cooling as the HEC. The inlet temperature of the cooling fluids was maintained at
 677 20°C and a floating Rota meter (PLATON PTF2 ASS-C with a volumetric flow rate of 0.2 Liters/min -
 678 1.5 liters/min) was used to control the flow rate of the cooling fluid during the experiments. During
 679 the experiment, the volumetric flow rate was set at 0.25 liters/min in order to calculate the heat
 680 transfer enhancement of the cooling fluid using the calorific method. The following equations (48,49
 681 and 50) were used for calculating one of the heat-transfer rates and for error analysis [61].

682
$$Q = m C_p (T_{out} - T_{in}) \quad (48)$$

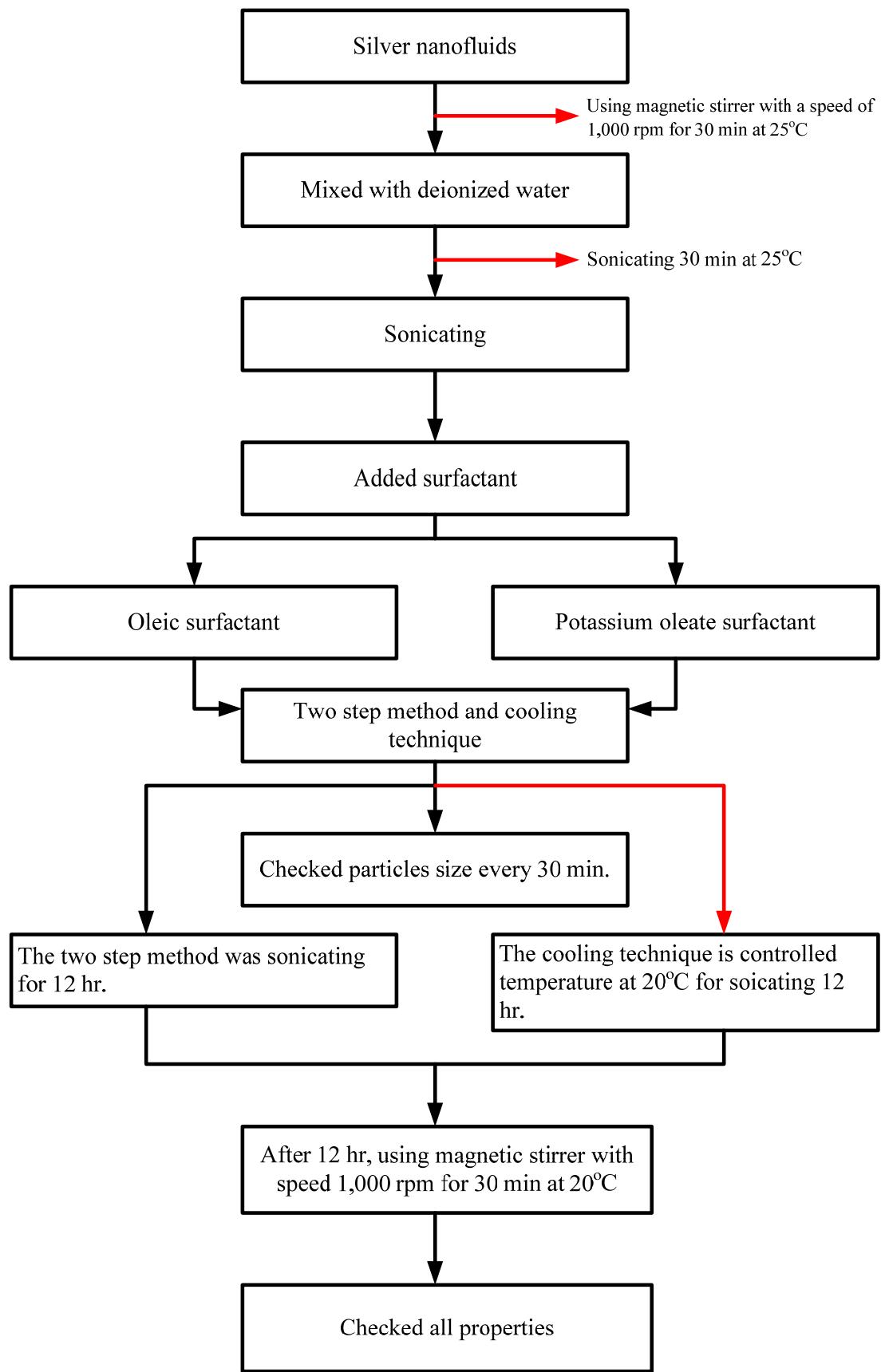
683 Thus:

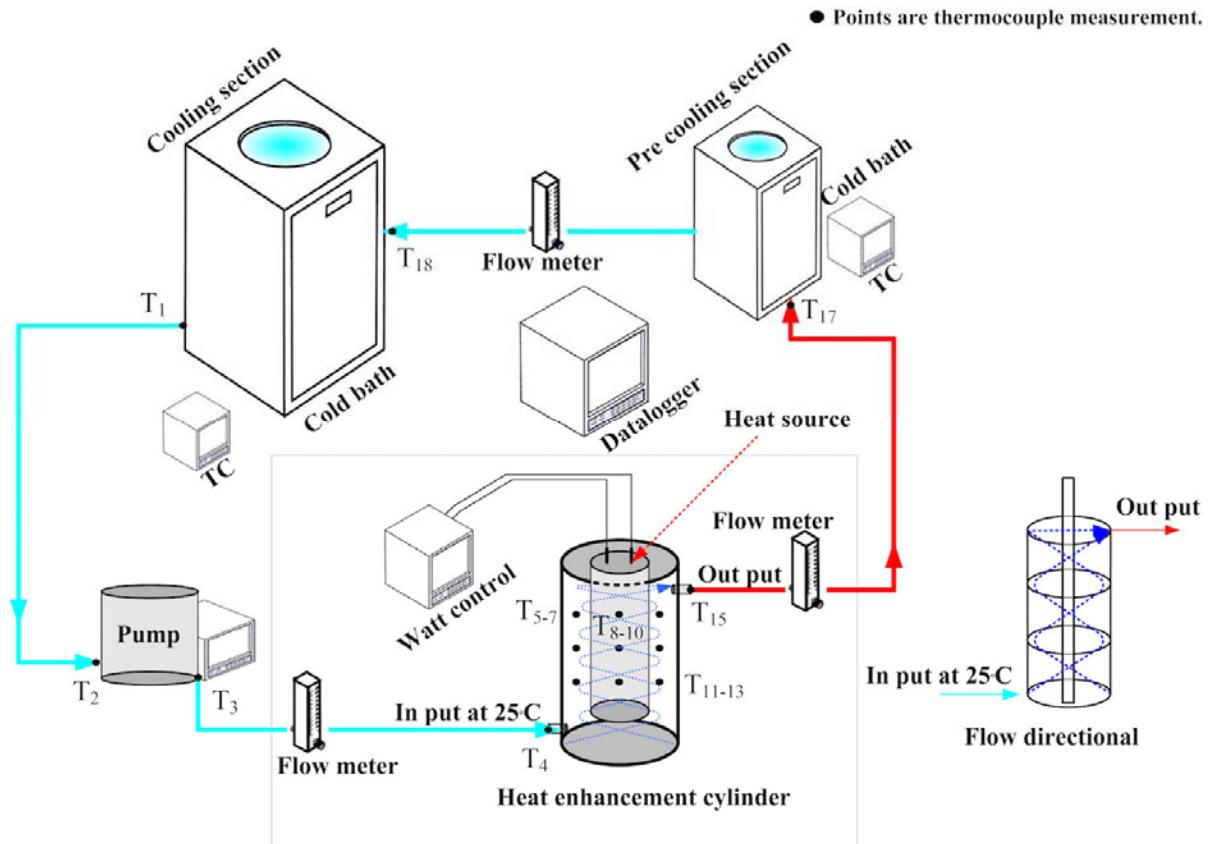
684
$$Q = f(m, T_{out}, T_{in}) \quad (49)$$

685 The error analysis of the heat transfer can be obtained from [23]:

686
$$Q_{Error} = \left[\left(\frac{\partial Q}{\partial m} \times m \right)^2 + \left(\frac{\partial Q}{\partial T_{out}} \times T_{out} \right)^2 + \left(\frac{\partial Q}{\partial T_{in}} \times T_{in} \right)^2 \right]^{0.5} \quad (50)$$

687 In order to experiment with a wide range of aspect ratios, the following parameters were
 688 corresponding set, as shown in Table 2, to formulate the heat transfer characteristics of the HEC [24].





691

692 Figure 19 Schematic diagrams of the HEC experimental apparatus

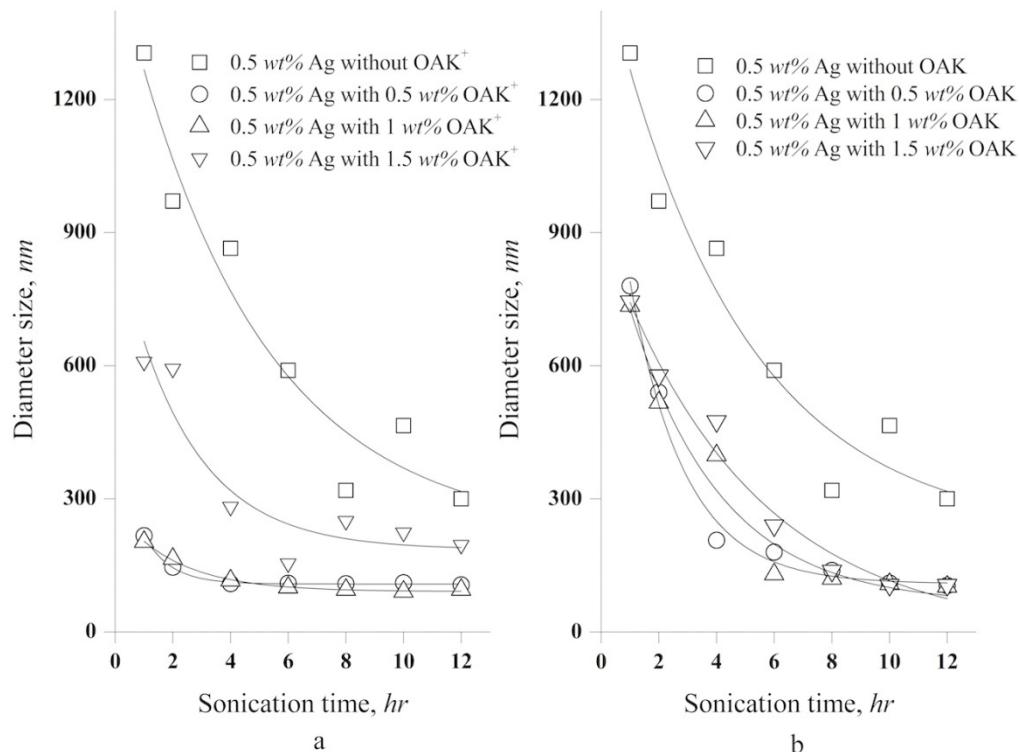
693 **3. Results and discussion**

694 **3.1. The nanoparticles size**

695 Figure 20 shows the average particle size as a function of sonicating time. Zero point 5 wt%silver
 696 nanoparticles based nanofluids (NF) with surfactants as a stabilizer have an average particle size of
 697 ~100 and ~95 nm respectively. It can be seen that the average particle size decreases as sonicating
 698 time increases. Moreover, the red ellipse in

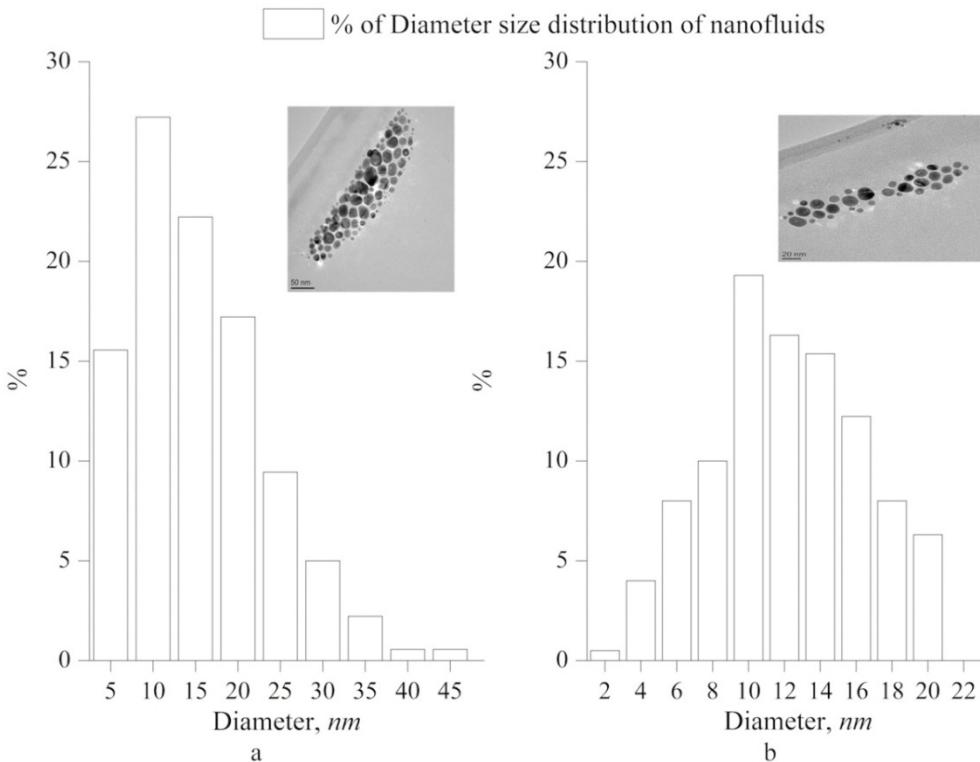
699 Figure 20a, NF+0.5OAK+ and NF+1OAK+ was seen to cause smaller particle size for a continuous
 700 period of 12 hours when compared with Figure 3b. This indicated that the two-step method and
 701 cooling technique did not break the agglomerate into primary particles [47, 62]. After sonicating for
 702 12 hours, the sample was put into the TEM (Oxford Instruments) to check for average the particle
 703 size, as shown in

704 Figure 21a, suggesting that the size distribution of particles NF+1OAK⁺ was between 5 – 25 nm. The
705 TEM image also shows that the long chains of potassium oleate combined with outside nanoparticles
706 and prevented them from aggregating together [63, 64], which means that the viscosity and surface
707 tension of the surfactant provided enough support to stabilize the dispersion of NP in deionized water
708 [2].



709

710 Figure 20 Relationship between sonication times and particle size



711

712 Figure 21 Relationship between Diameters with percentage of dispersed size and TEM micrograph at
713 silver nanofluids containing surfactant at concentration 1 wt%

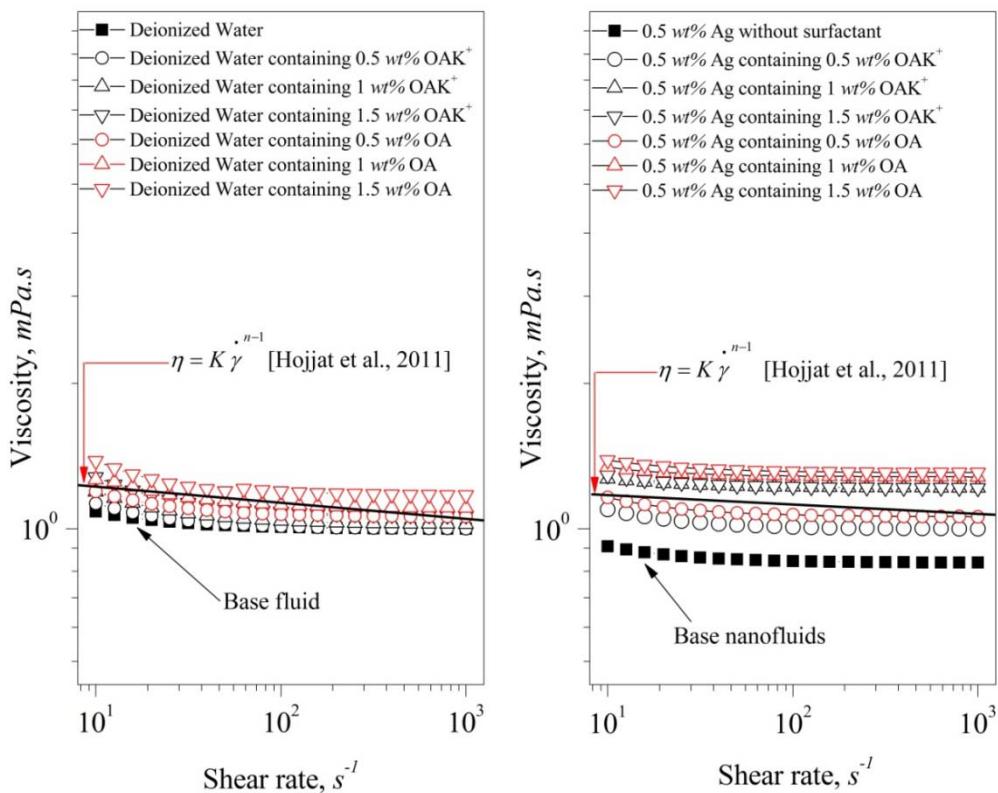
714 **3.2. Rheological properties of nanofluids**

715 The shear rate and shear stress had an effect on the rheological properties, such as viscosity.
716 The rheological properties of the nanofluids containing surfactant are important to its thermo physical
717 property. In this study, a surfactant was used to employ the NF's heat transfer rate. The OA and
718 OAK⁺ are known for their ability to decrease viscosity and surface tension due to the organic and
719 hydrocarbon interaction with oxygen which exists in deionized water [64, 65]. However, the case was
720 compared with the same group of surfactant (OA and OAK⁺) but with a difference in potassium salt
721 (K⁺). The potassium salt was helpful in balancing the pH value of the liquid and dissolving the
722 solutions of fat oil catalysis interaction with hydrogen. Thus, the OAK⁺ has the ability to span the
723 NP's random motion throughout the deionized water [2, 53, 66-68].

724 The rheology measurement results are shown in Figure 23 and the shear rates with the
725 viscosity of silver nanofluids were measured at 30°C according to Paramethanuwat et al., [2]. As
726 shown in Figure 22, the viscosity of all samples decreased in the first 101s-1 to 103s-1 intervals and
727 the NF showed Newtonian behaviour. This behaviour could have been caused by the change of
728 concentration in the OA and OAK^+ , which was 0.5, 1 and 1.5 wt%. The most preferable OAK^+
729 concentration was 1 wt%, which was sufficient to distribute the NP with the lowest and most stable
730 viscosity. The long chain nature of the OAK^+ molecular structure helped to decrease the NF's surface
731 tension. It was concluded that there was an apparent change in the viscosity of the NF and deionized
732 water; however, the NF's viscosity was still larger than that of the deionized water when the shear rate
733 rose [69] according to Equation (42) [52]. Thus, the NP existing in the deionized water containing
734 OAK^+ , which affected the flowing behaviour of the nanofluids, was the main cause of the decrease in
735 viscosity [70, 71]. The potassium salt which produces an effect properly surfactant was helpful in
736 balancing the physical properties of the base liquid. As can be seen in Figure 23, along with the
737 increasing shear rate, the nanofluids with OAK^+ concentration smaller or larger than 1 wt% possessed
738 larger shear stress. From the cross point onward, all of the results of the OAK^+ were of almost the
739 same valve and the R^2 was close to 1. This was in line with the study of Hojjat et al., [52], who stated
740 that the transition metal in the same group with silver nanofluids containing OAK^+ produced the
741 Newtonian fluid [2].

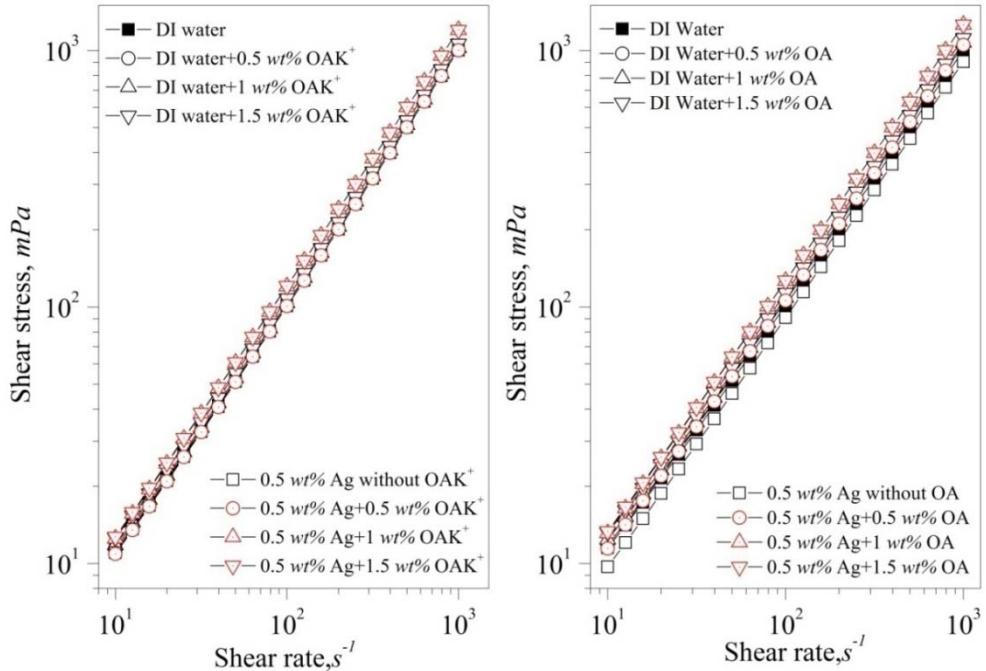
742 Figure 24 shows the viscosity of NF+1OA compared with 0.5 wt%NF+1 OAK^+ as a function
743 of shear rates. It was observed that for all operating temperatures and shear rate larger than 10¹s-1, the
744 viscosity became stable and NF showed Newtonian behaviour. Thus, it was well established that for
745 all operating temperatures, the system's rheological behaviour exhibited a similar trend. Figure 25
746 shows the relationship of the shear rates and shear stress in accordance with Figure 24. It could be
747 explained that the higher temperature increased the intermolecular distances, which decreased the
748 interaction between the molecular structures of deionized water and OAK^+ , resulting in decreased
749 viscosity and surface tension [47]. Obviously, the OAK^+ could help decrease the physical properties
750 more than the OA. The surfactant behaved like an interfacial shell between the nanoparticles and base

751 fluids and modified the surface tension of the nanofluids. The surface tension decreased when the
 752 concentration of surfactant increased [2, 72]. The OAK⁺ exhibited good adsorption of the silver
 753 particles and the particles uniformly had a direct effect on the shear stress. The optimization of the
 754 chain length in group OA organic compounds was C18, which was effective for particle dispersing
 755 stabilization. The optimized length also improved the colloidal stability and increased the non-
 756 precipitation period for the nanoparticles to be uniformly dispersed [45, 63, 73]. Moreover, OAK⁺
 757 achieved more stable suspension than the pure NP did in the deionized water. This might be related to
 758 the Newtonian property of the nanofluids observed in this study when the rheological properties were
 759 observed at operating temperature [2, 53, 74, 75].



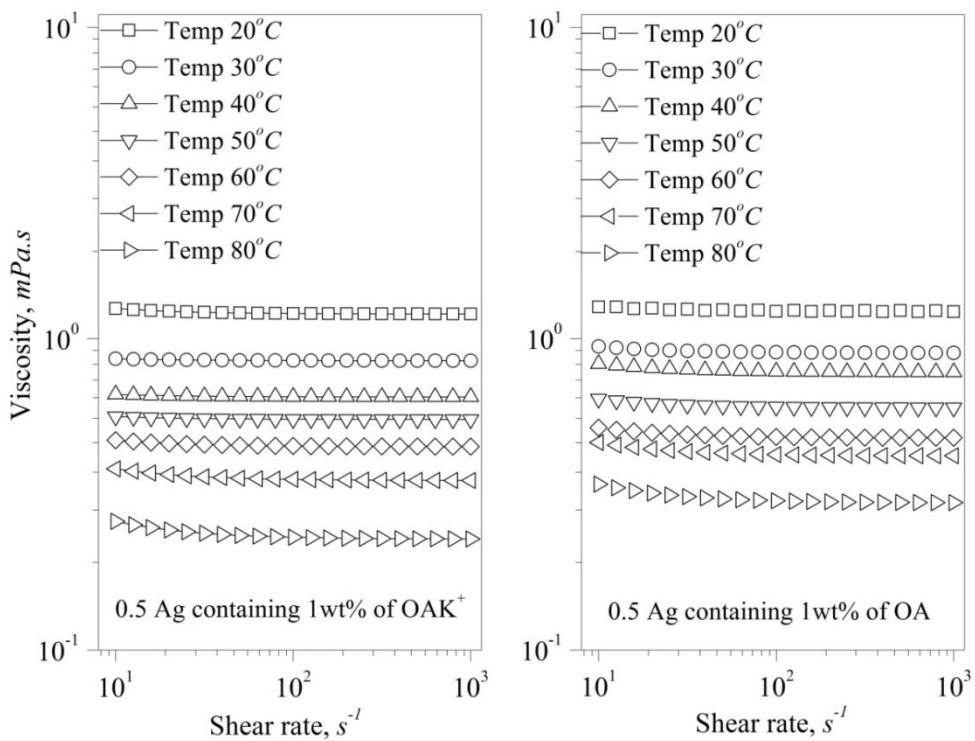
760

761 Figure 22 Relationship between shear rates with viscosity of silver nanofluids at operating temperate
 762 30°C



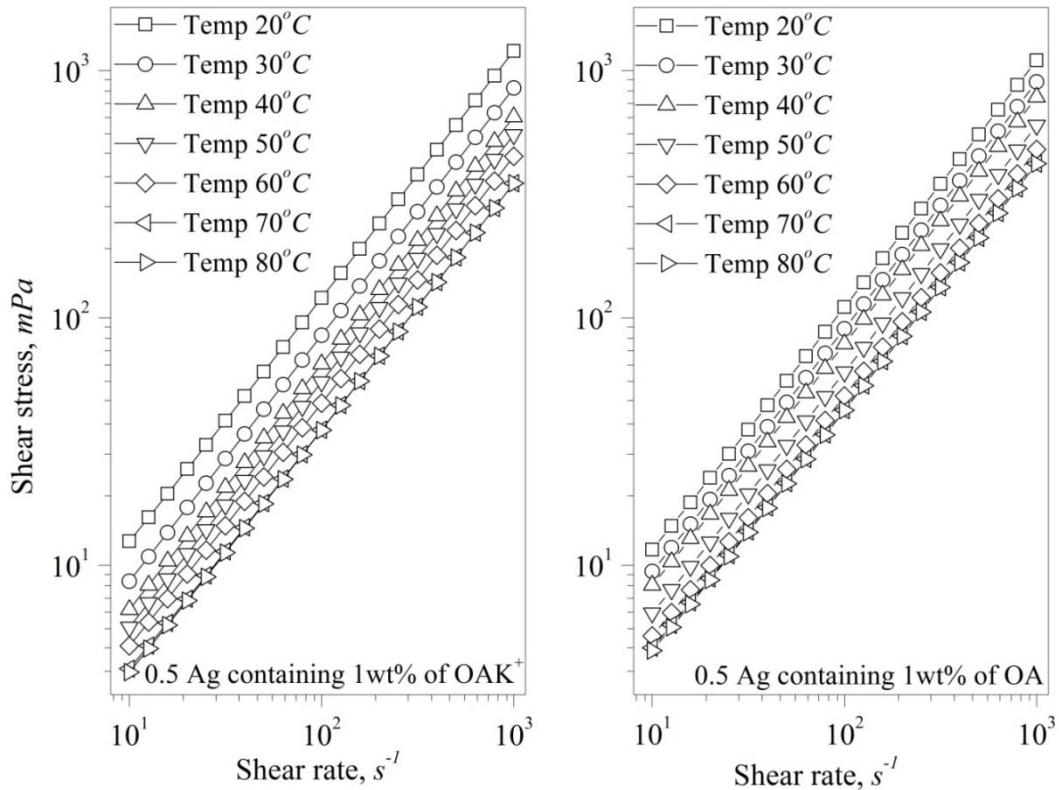
763

764 Figure 23 Relationship between shear rates with shear stress of silver nanofluids at operating
765 temperate 30°C



766

767 Figure 24 Relationship between shear rates with viscosity



768

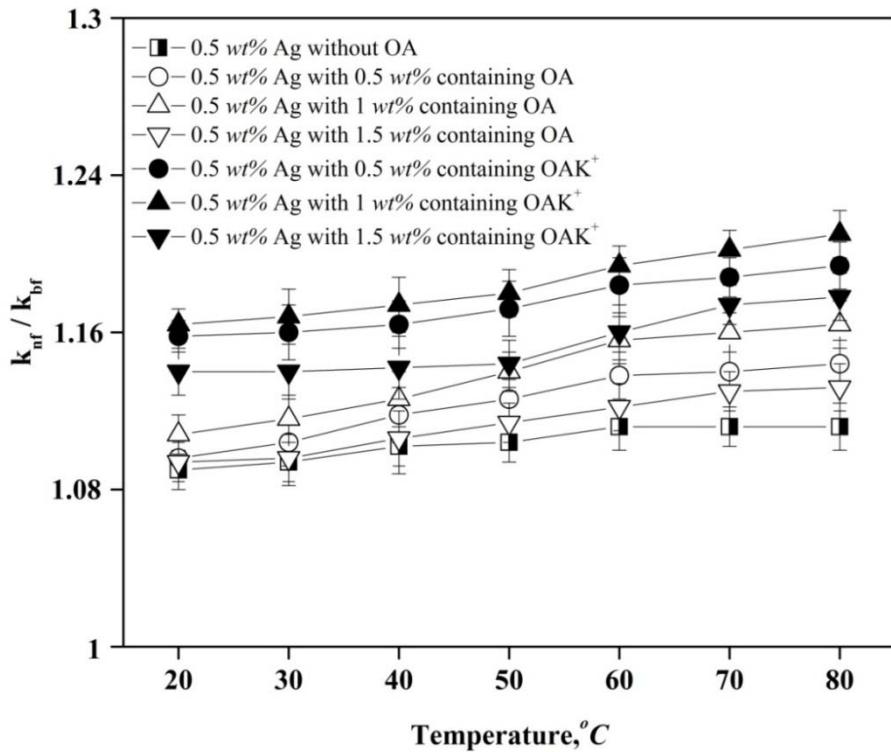
769 Figure 25 Relationship between shear rates and shear stress

770 **3.3. Thermal conductivity of nanofluids**

771 The thermal conductivity effective of the silver nanofluids containing potassium oleate
 772 ($\text{NF}+\text{OAK}^+$) as a function of temperature is shown in Figure 26. It can be seen that the thermal
 773 conductivity of the nanofluids depend on the linearity of the temperature, and the enhancement of the
 774 thermal conductivity of $\text{NF}+\text{OAK}^+$ was different when the surfactant concentration was 0.5, 1, and 1.5
 775 wt%. In all cases the $\text{NF}+\text{OAK}^+$ showed superior performance to that of the water. One wt% OAK^+
 776 showed the lower and highest increase in the thermal conductivity of 15% at 20°C and 28% at 80°C
 777 throughout all samples, indicating that the thermal conductivity increases independently on surfactant
 778 concentration [47]. The nanoparticles dispersed in the liquid increased the surface area for the heat
 779 absorption. In the case of $\text{NF}+\text{OAK}^+$, the OAK^+ decreased the surface tension of the nanoparticles,
 780 stabilized the nanoparticles by uniformly distributing them, and increased the interface area of the
 781 nanofluids with the deionized water [63]. The surface tension had a significant influence on the

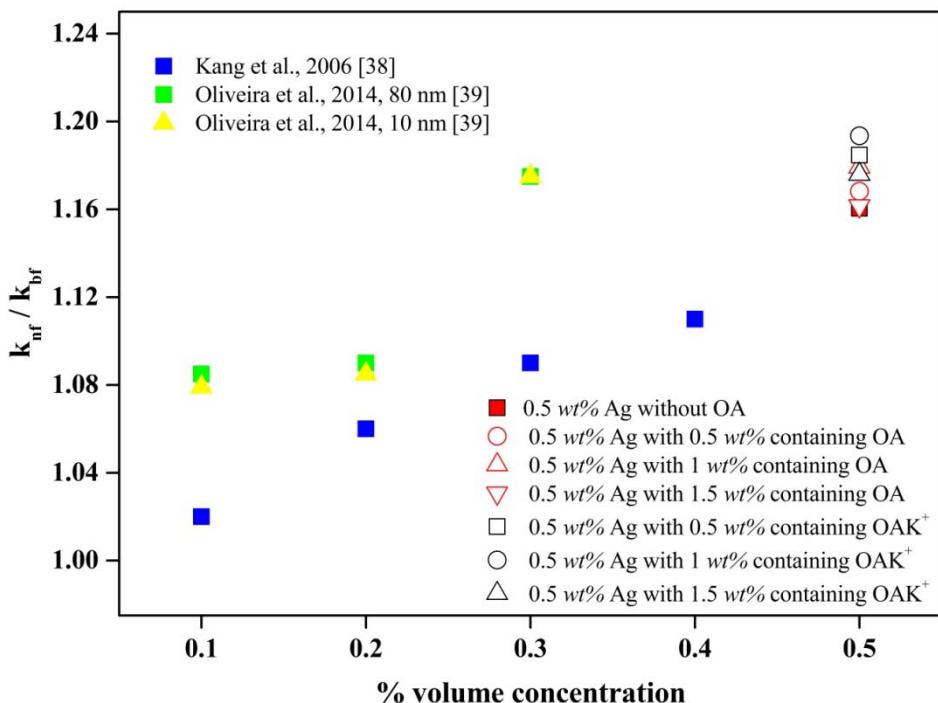
782 thermal process since the property departure and interfacial equilibrium depend on it [23]. The high
783 OAK⁺ concentration appeared to hinder the aggregation and entanglement of the NP, which was
784 observed at the bottom of the liquid [44, 46]. According to our experimental results, 1 wt% OAK⁺ was
785 enough to homogeneously disperse the NP and produce efficient thermal transfer between the
786 particles and deionized water, and consequently resulted in the highest thermal conductivity
787 enhancement [43, 76-79].

788 However, the current experimental results contrast with those of Kang et al., [38] and
789 Oliveira et al., [39] as shown in Figure 27. In regards to this study, the results were achieved using the
790 same silver nanoparticles however with a difference in concentration and surfactant. The result from
791 Kang et al., demonstrated an increase of relative thermal conductivity. Thus, it is important to note the
792 vast differences among different experimental conditions, especially in regard to the method of
793 preparation and nanoparticles concentration. For example, Kang et al.,[38] showed a maximum
794 k_{nf} / k_{bf} of ~1.11 at 0.4% volume concentration with nanoparticles diameter of 8-15 nm, whilst
795 Oliveira et al., [39] showed a maximum k_{nf} / k_{bf} of ~1.17 at 0.3% volume concentration with
796 nanoparticles diameter of 10 and 80 nm. This contrasts significantly with the results achieved in this
797 study of approximately 1.19 with nanoparticles of diameter 5-25 nm at NF+1OAK⁺. Of particular
798 note, the research performed by Kang et al., in 2006 and Oliveira et al., in 2014, did not specify the
799 temperature under which the experiment was conducted. Furthermore, relative thermal conductivity
800 was analyzed as a function of nanoparticles concentration and size. Thus, it can be determined that
801 relative thermal conductivity is dependent upon the nanofluid's method of preparation, as well as the
802 nanoparticles concentration and size.



803

804 Figure 26 Relationship between temperature with k_{nf} / k_{bf}



805

806 Figure 27 Comparison different experimental result

807 **3.4. Dynamic of specific heat capacity (DSC)**

808 The heat energy absorbed ratio is a substance to increase in temperature. Table 9 shows the
809 dynamic of specific heat capacity: nanofluids. The DSC was used to measure the deionized water
810 deionized water containing surfactant (OA and OAK^+) and the NF containing surfactant (OA and
811 OAK^+) at 1 wt% respectively, which was then compared with Equation (47) according to Rajabpour
812 et al., [57] and O'Hanley et al., [58]. The results indicated a trend similar to that found in Rajabpour et
813 al., [57] and O'Hanley et al., [58]. It was found that, the good resulted appeared when using OAK^+ as
814 a surfactant. The silver nanofluids containing OAK^+ surfactant was subjected to repulsion forces
815 between the positively-charged hydroxyl groups (OA) of the functionalized nanoparticles and the
816 potassium salt hydroxyl groups on the silver. However, the potassium salt of the acidic group
817 functionalizing the nanoparticles became polarized in the water solution. On the other hand, when the
818 positivity-charged potassium cation (K^+) groups were attached to the nanoparticles, the free ends (the
819 carboxylic groups) became negatively charged. It is possible that the particle-fluid interactions and
820 long-range electrostatic interactions between the nanoparticles may have affected the capillary
821 properties of the nanofluids. Therefore, the negative forces between the solid and the nanofluids
822 induced a specific heat capacity. Both parameters were dependent on particle size and surfactant
823 concentration.

824 **3.5. The contact angle and surface tension**

825 In this study, the drop contact angle of working fluids was based on the room operating
826 temperature. Furthermore, in order to investigate the effect of adding surfactant to the nanofluids,
827 surface wettability, the static contact angles of the sessile droplets, and the surface tension of the
828 pendant drop (Contact Angle Meter Model: DM-CE1 ; Kyowa Interface Science) were measured on
829 flat copper plates at room temperature as shown in Figure 28. This method was based on Khandekar
830 et al., [36] and Rahimi et al., [80]. When the surfactant was added it caused reduced adhesion between
831 the working fluids dropping and the metal surface and made the total surface free energy. These are
832 the manifestations of the interaction of the different molecular forces. The effects on the bulk thermo
833 physical properties need to be addressed and surfactant dealt with nanofluids thus according to

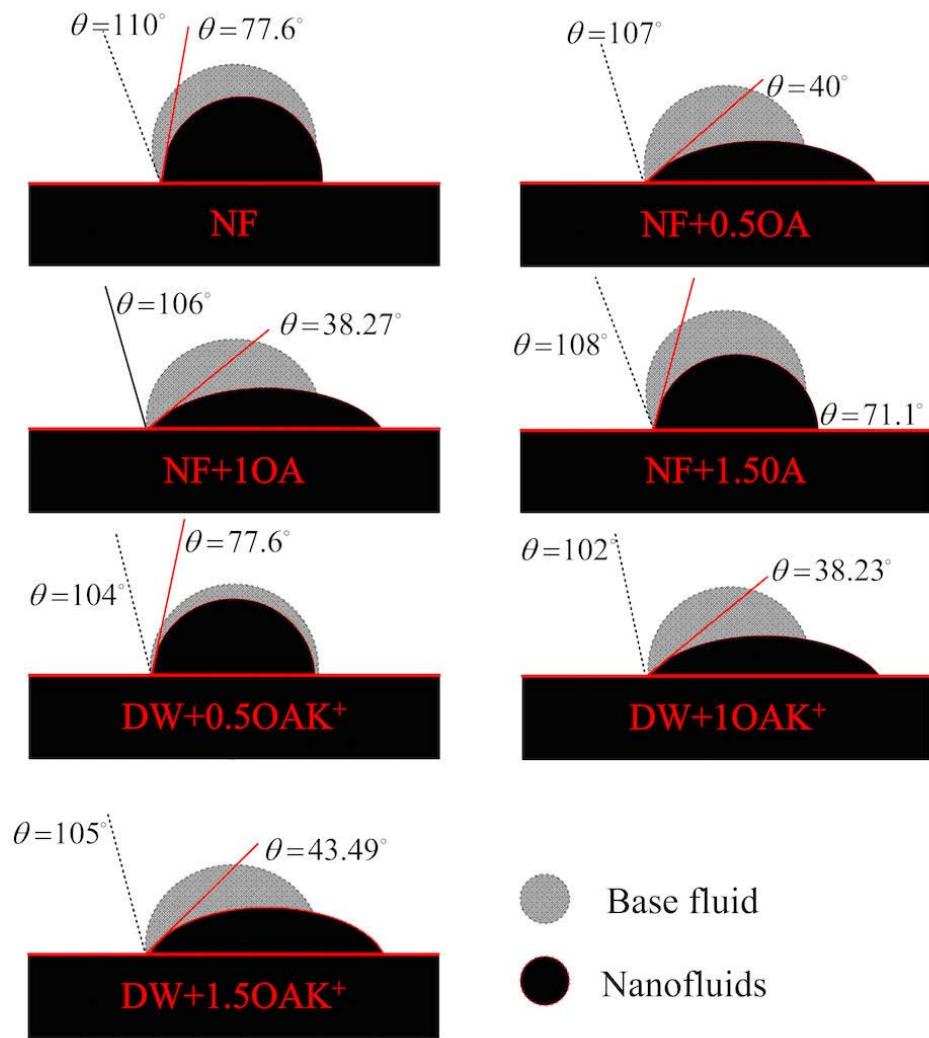
834 Radiom et al., [81]. It was further found that NF+1 OAK⁺ decreased the water drop contact angle and
835 surface tension to 38.23° and 28.69 mN/m, respectively (compared with pure water having 110° and
836 72.8 mN/m). Thus, the contact angle and surface tension depended on the operating temperature and
837 time. They have an effect on the boiling phenomenon in the heat transfer application. The transfer
838 rate improves and decreases the surface tension with wet ability and contact angle [2, 82].

839 **3.6. Heat transfer enhancement**

840 The experimental results clearly showed the effect of the percentage of heat supplied on the
841 percentage of thermal heat enhancement, as shown in

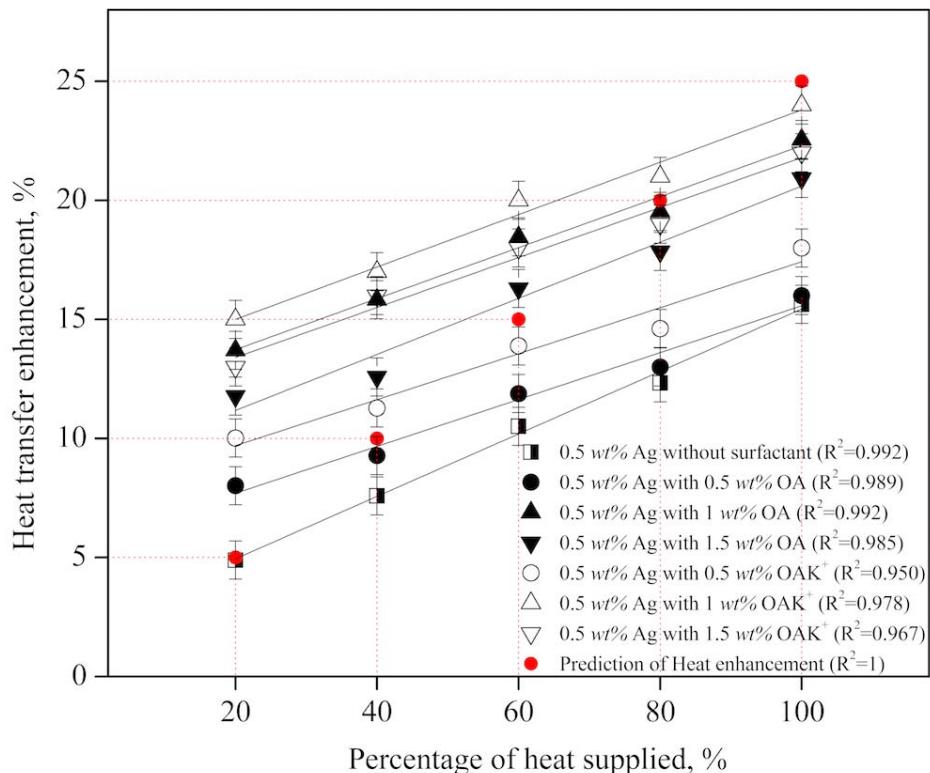
842 Figure 29. When comparing the percentage of heat supplied using the different working fluids, it was
843 seen that the NF+1OAK⁺ showed that the percentage of thermal enhancement was higher than with
844 the other working fluids. Considering the case where the working fluid was NF+1OAK⁺ with a heat
845 input at 100% of heat supplied, the thermal enhancement reached 24.75±0.08%. The increase of
846 maximum percentage of thermal enhancement with an increase in the percentage of heat supplied can
847 be attributed to the increase in different temperatures (ΔT). In addition, the thermal motion of the
848 nanoparticles enhanced the thermal properties of the nanofluids. The OA group helped with the
849 homogeneous dispersion of the nanoparticles in the nanofluids. Moreover, the potassium cation (K⁺)
850 significantly contributed to accruing the thermal property increase, which had effects on the heat
851 transfer rate mechanism more efficiently than the thermal diffusion in the fluid [34, 45, 83, 84].

Temperature (°C)	Deionized water containing 1 wt% OA			Deionized water containing 1 wt% OAK ⁺			0.5 Ag containing 1 wt% OA			0.5 Ag containing 1 wt% OAK ⁺		
	Theoretical C_p (kJ/kg·K)	Measured C_p (kJ/kg·K)	Theoretical C_p (kJ/kg·K)	Measured C_p (kJ/kg·K)	Theoretical C_p (kJ/kg·K)	Measured C_p (kJ/kg·K)	Theoretical C_p (kJ/kg·K)	Measured C_p (kJ/kg·K)	Theoretical C_p (kJ/kg·K)	Measured C_p (kJ/kg·K)	Theoretical C_p (kJ/kg·K)	Measured C_p (kJ/kg·K)
20	-	4.182±0.070	-	4.189±0.056	-	4.226	4.227±0.002	4.226	4.225±0.055	4.225±0.055	4.269±0.02	4.225±0.055
30	-	4.183±0.068	-	4.189±0.036	-	4.228	4.227±0.045	4.228	4.225±0.103	4.225±0.103	4.269±0.120	4.225±0.103
40	-	4.182±0.068	-	4.190±0.032	-	4.229	4.228±0.020	4.229	4.225±0.003	4.225±0.003	4.270±0.025	4.225±0.003
50	-	4.182±0.068	-	4.204±0.048	-	4.231	4.242±0.025	4.231	4.226±0.028	4.226±0.028	4.284±0.008	4.226±0.028
60	-	4.183±0.068	-	4.204±0.060	-	4.237	4.242±0.012	4.237	4.227±0.040	4.227±0.040	4.284±0.023	4.227±0.040
70	-	4.187±0.068	-	4.214±0.089	-	4.241	4.245±0.121	4.241	4.229±0.104	4.229±0.104	4.295±0.023	4.229±0.104
80	-	4.189±0.068	-	4.224±0.010	-	4.252	4.262±0.121	4.252	4.236±0.112	4.236±0.112	4.305±0.039	4.236±0.112



853

854 Figure 28 Wettability at room operating temperature



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856

857 Figure 29 Thermal enhancement of working fluids

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865 **4. Conclusion**

866 Study on NF containing surfactant is leaded tonanoparticles size, the rheological properties
867 of the nanofluids, the thermal conductivity of nanofluids, the dynamic of specific heat capacity
868 (DSC), wet ability (contact angle and surface tension) and heat enhancement. The important details
869 are discussed below:

870 - The silver nanofluids containing OA and OAK^+ were conducted on thermal conductivity and
871 rheological properties at various concentrations of OA and OAK^+ and operating temperatures.
872 It was found that the NF containing 1 wt% of OAK^+ yielded better particle size ~ 95 nm.

873 - At a shear rate range of 101s⁻¹ to 103s⁻¹, the samples showed Newtonian behaviour, which
874 showed Newtonian behaviour, suggesting that the shear stress and viscosity decreased the
875 high solid loading. For a given surfactant concentration, the consistency viscosity and shear
876 stress of the base fluid and all of the nanofluids decreased with an increase in temperature,
877 which confirmed that temperature had a strong effect on the shear stress and viscosity of the
878 nanofluids. The rheological produced Newtonian.

879 - The NF containing 1 wt% of OAK^+ gave the highest thermal conductivity. It can be seen that
880 the thermal conductivity enhancement was from 11% at 20°C to 28 % at 80°C when compared
881 with the base fluids. The DSC was increased with respect to operating temperature increase.
882 Explicitly, the thermal conductivity of the NF containing the surfactant was in respect to the
883 operating temperature, showing increments at all concentrations.

884 - The specific heat of NF containing OA and OAK^+ was superior in specific heat capacity, over
885 water studied in all experimental conditions. The presence of surfactant had clearly
886 contributed to the rise in specific heat capacity.

887 - It was concluded that the static contact angles of the OA group surfactant used, have better
888 wettability characteristics, dependent on the surfactant concentration. Moreover, the NF

889 containing 1 wt% of OAK+ could be good at reducing the wettability and the OA group
890 improved the colloidal stability which potassium cation (K+) increased the non-precipitation
891 period for nanoparticles to be uniformly dispersed in the base fluid. The nanofluids containing
892 1wt% of OAK+ produced a good contact angle of 38.23°.

893 It could be concluded that the amount of the thermal enhancement of the nanofluids
894 containing surfactant contributed to the greater rise of the thermal performance over the base
895 fluid/nanofluids by approximately 80%.

896 **CHAPTER IV THERMAL PERFORMANCE OF LOOP**
897 **THERMOSYPHON**

898 This chapter is described on the heat transfer rate behaviour in of an advanced loop
899 thermosyphon with check valve (ALT/CV) which is filled with silver nanofluids, and containing OA
900 and OAK^+ . Traditional working fluids such as water, ethanol, etc., have poor physical properties
901 which makes it difficult to change phase [10, 11]. Silver nanoparticles were chosen to be added in the
902 traditional working fluid because it has the highest thermal conductivity in the transition metal group
903 and could be a good stabilizer in a base fluid compared to most solids [11]. The particles could
904 improve thermal performance. The problem with nanofluids is agglomeration with un-dispersed
905 nanoparticles due to the decrease in the lead time period for nanoparticles to be uniformly
906 dispersed[85]. Consequently, this article focuses on improving working fluid properties which is
907 achieved by adding surfactant. The OA and OAK^+ could decrease the surface tension and increase the
908 homogeneity of the dispersed nanoparticles [48, 85]. Silver nanoparticles containing OA and OAK^+
909 could increase thermal conductivity and heat transfer coefficient. The purpose of the silver
910 nanoparticles containing OA and OAK^+ in the working fluid is for filling of the TPCT was to find out
911 the optimal OA and OAK^+ surfactant concentration in order to maximize heat transfer rate. The new
912 working fluid has a unique characteristic where the heat transfer rate is slightly higher than the
913 traditional working fluid, as follows:

914 This chapter describes the heat transfer rate behaviour of an advanced loop thermosyphon
915 with check valve (ALT/CV) which is filled with silver nanofluids and contains OA and OAK^+ .
916 Traditional working fluids such as water, ethanol, etc., have poor physical properties, which makes it
917 difficult to change the phase [10, 11]. Silver nanoparticles were chosen to be added to the traditional
918 working fluid because it has the highest thermal conductivity in the transition metal group and can be a
919 good stabilizer in a base fluid compared to most solids [11]. Additionally, the particles can improve
920 thermal performance. The problem with nanofluids is agglomeration with un-dispersed nanoparticles
921 due to the decrease in the lead time period for nanoparticles to be uniformly dispersed [85].
922 Consequently, this article focuses on improving working fluid properties, which is achieved by adding

923 surfactant. OA and OAK^+ can decrease the surface tension and increase the homogeneity of the
924 dispersed nanoparticles [44, 81]. Silver nanoparticles containing OA and OAK^+ can increase thermal
925 conductivity and the heat transfer coefficient. The purpose of the silver nanoparticles containing OA
926 and OAK^+ in base fluid is filling in the ALT/CV, was to find out the optimal OA and OAK^+ surfactant
927 concentration in order to maximize the heat transfer rate. The new working fluid has a unique
928 characteristic, where the heat transfer rate is slightly higher than the traditional working fluid, as
929 discussed below.

930 **1. Literature review**

931 The objective of this study is based on the ALT/CV using silver particles containing OA and
932 OAK^+ as the working fluid. Moreover, it has proven beneficial to improve the working fluid
933 properties as a way to allow for smaller nanoparticles which ultimately reduce the thermal resistance
934 contained in the base fluid as the working fluid in the ALT/CV. The effects of the operating
935 percentage of the heat supplied (heat input), the working fluids, the loop size of the thermosyphon and
936 thermal behaviour (thermal resistance, heat transfer coefficient, relative of thermal efficiency) under
937 normal operating conditions were studied.

938 This literature review consists of a number of points to be reviewed and discussed. The
939 requirements for the ALT/CV and the nanofluids will be summarized, together with the existing
940 characterization measurement and application of the online methods, which will also be reviewed.
941 Thus, the addition of nanoparticles to a base fluid forms so-called nanofluids, as the nanoparticles can
942 improve the heat transfer rate, which can be used in the ALT/CV or heat exchanger applications.
943 Then, nanofluids are used as working fluids in the heat pipes. The points to be considered include the
944 method using the nanofluids and the effect this has on the ALT/CV. Thus, the heat transfer rate
945 behaviour of the ALT/CV is essential in establishing an adequate application and design of the
946 processing. Payakaruk et al., [25] studied the heat transfer rate of a two phase closed thermosyphons
947 (TPCT) made from copper tubes with diameter as 7.5, 11.1 and 25.4 mmID. Five working fluids were
948 chosen: water, ethanol, R-22, R 123, and R-134a. It was concluded that the working fluid affected the

heat transfer rate at inclination angles of 20 to 70°. The filling ratios had no effect on the ratio of the heat transfer at any angle; however, the properties of the working fluids were affected. Ristoiu et al.,(2003) [86] experimentally studied the effect of inclination angles on the heat transfer of a wickless solar heat pipe (Thermosyphon). The thermosyphon used acetone, methanol, and water as working fluids. The copper heat pipe investigated here exhibited the highest heat transport rate at inclinations of around 40° - 45°. Then, the study of nanofluids was conducted by suspending ultra-fine metallic or non-metallic particles from the nanometer dimension group in base fluids such as water, oil, and ethylene glycol. Noie (2005) [87] reported that the heat transfer performance of the TPCT with aspect ratios was 7.45, 9.8 and 11.8. The parameter had input heat transfer rates of $100 < Q < 900$ W with filled the working fluid ratios of $30\% \leq FR \leq 90\%$. It was found that, with the filling ratio at 60%, a higher heat transfer rate occurred and was lower at 90%. Moreover, a drop in temperature was expected due to the internal resistance of boiling and condensation. After the heat transfer coefficient measurement, it was found that the heat transfer of all the filling ratios were reasonable for finding the empirical correlation between the experimental results and predicted values. Then, Noie et al., (2007) [4] studied the effect of the inclination angle from 5° to 90° with different filling ratios (Distilled water as the working fluid) of 15%, 22%, and 30%. The thermal performance of a TPCT(copper tube diameter of 14.5 mmID, length 1,000 mm) was investigated experimentally under normal operating conditions. It was found that the TPCT had the highest thermal performance in the inclination angle range of 15° to 60°. The interesting phenomenon of geyser boiling occurred in our experiments for filling ratios equal to or greater than 30%. The geyser boiling puts no limitation on the thermal performance of the thermosyphon; however, it should be avoided because it damages the condenser end cap due to the slug striking. Another study was applied to nanofluids in 2009 [18]. This investigation, studying Al_2O_3 in the water, showed that the efficiency of the TPCT was enhanced up to 14.7% at 3 vol%. It was also found that the temperature distribution on the TPCT was at a lower level when nanofluids were used as opposed to pure water. The thermal resistance of the TPCT was less. When nanofluids were changed with the higher thermal performance of the TPCT loaded with nanofluids proved its potential as a substitute for a convectional one with pure water [20]. In 2011, Khazaee et al., [88] presented a study of the heat transfer characteristics of the TPCT made from a

977 copper tube of 15 mmOD and a length of 2,000 mm with iron oxide-nanofluids (Averagediameter of
978 4–5 nm) in water as the working fluid. It was found that the effects of the TPCT inclination angle
979 wasbetter at 90°. However, this depends on the operating temperature and nanoparticles concentration
980 levels of the heat transfer characteristics of the TPCT. The thermal resistance of the TPCT with
981 nanoparticles solution waslower than that with pure water. This shows that thermal resistance
982 decreases as the volume concentration increases.However, this depends on the operating temperature
983 and nanoparticles concentration levels of the heat transfer characteristics of the TPCT. The thermal
984 resistance of the TPCT with nanoparticles solution is lower than that with pure water. This shows that
985 thermal resistance decreases as the volume concentration increases. The different types of nanofluids
986 have been investigated by Khandekar et al., [20]. This used Al_2O_3 , CuO and laponite clay mean
987 diameter < 100 nm in water. The TPCT was made from copper tubing with a filling ratio was of
988 100%. It was found that nanofluids show inferior thermal performance than pure water. The wet
989 ability of all nanofluids on copper substrate, having the same average roughness as that of the TPCT,
990 is better than pure water. A scaling analysis in an evaporator side Peclect number found this
991 eventually leads to poor thermal performance. The condenser and evaporator were an important factor
992 on overall performance. When studying the TPCT, the physical effects are very important.

993 Paramatthanuwat et al., [3] studied the heat transfer of the thermosyphon made from copper
994 tubes with 7.5, 11.1, and 25.4 mmID, and the aspect ratios (Le/di) chosen for the study were 5, 10,
995 and 20. The filling ratios chosen for the study were 30%, 50%, and 80% with respect to the
996 evaporator length. Pure water and silver nanofluids were used in thermosyphon and compared with
997 three operating temperatures of 40, 50 and 60°C. It was found that the best filling ratio was at 50% for
998 the temperature to affect the heat transfer rate of the thermosyphon. Kang et al., (2009) [77]
999 investigated silver nanofluids in water applied as a working fluid in a heat pipe. The silver nanofluid
1000 showed better cooling with respect to water because the nanoparticles could flatten the temperature
1001 gradient of the fluid and reduce the boiling limit because it increased theeffective liquid conductance
1002 in the heat pipe. Hence, the nanofluids aremore interesting as cooling fluids for devices because they
1003 have a higher energy density. Kim et al., (2007) [89] studied the pool boiling of dilute dispersions of

1004 alumina, zirconia, and silica in water. It was found that the nanofluids improved the layer and were
1005 important to the surface wettability, as shown by a reduction in the static contact angle on the
1006 nanofluid-boiled surfaces compared with the pure boiled water surfaces.

1007 Thus this section focuses on the physical effects of substance based fluid functions and
1008 applications of nanofluids. However, few studies have been carried out to improve physical
1009 properties, for example decreasing surface tension or changing the structure or decreasing the
1010 viscosity of the base fluid. Hwang et al.,(2008) [45] investigated silver nanofluids in silicon oil with
1011 oleic acid (OA) produced by a one-step method (Magnetron sputtering),which showed the largest
1012 improvement in thermal conductivity over that of silicon oil. It was observed that silver nanoparticles
1013 were homogeneously dispersed and were stable for a long period of time in thesilicon oil. Li et
1014 al.,(2008) [46] investigated the thermal conductivity dependence with respect to the appropriate
1015 concentrations of an added surfactant. The work focused on the effect of pH, and the nanofluids
1016 chosen were Cu and sodium dodecylbenzenesulfonate (SDBS) as the surfactant. It was found that the
1017 surfactant was recommended to correct the thermal conductivity for practical applications of
1018 nanofluids. The maximum thermal conductivity could be enhanced by up to 10.7%. Hojjat et al.,
1019 (2011) [52] reported on the γ Al₂O₃, TiO₂ and CuOnanofluids with 0.5 wt% of carboxymethyl
1020 cellulose (CMC) in de-ionized water containing up to a 4 vol% of particle concentration. They
1021 concluded that the apparent viscosity of the base fluid and nanofluids decreasedwhen the shear rate
1022 increased. Rahimi et al., (210) [80] focused on thecontact angle and used a 1 m height TPCT with
1023 water as the working fluid with pressure operating at 0.75 and 160 mbar. It was found that the
1024 evaporator had more than the hydrophilic (4°) and the condenser had more than the hydrophobic
1025 (120°), so it was possible to raise the thermal performance to 15.27% and to decrease the thermal
1026 resistance to $\times 2.35$ when compared with plain water. Qi et al., (2001) [90] and Hwang et al., (2008)
1027 [45] concluded that the surfactant could reduce the surface tension of the working fluid, which would
1028 improve the colloidal stability and would increase the lead time period for thenanoparticles to be
1029 uniformly dispersed. This could increase the surface area for silver nanofluids to absorb heat; thus this
1030 was found to be an enhancement for boiling fluid.

1031 After that many researcher had trying to improve the performance of TPCT which was
1032 advanced the TPCT such as Fumito et al., (2003) investigated the loop closed thermosyphon, and
1033 showed that the heat transfer rate was better than the single tube thermosyphon[91]. Khodabandeh et
1034 al., (2010) [92] found that the maximum heat transfer flux of the loop thermosyphon was 20-44
1035 W/cm² which is larger than the single tube, due to the non-countercurrent flow and non-vapour lock
1036 limit. In any application, e.g. the plate heat source, one kind of heat pipe is always applied. Then,
1037 Chang et al., (2010) investigated the thermal performance of a two-phase closed-loop thermosyphon
1038 with a thermal resistance model for electronic cooling. Results indicated that the evaporator and
1039 condenser thermal resistance decreased about 15.5% [93]. However, the heat was dissipated to the
1040 evaporator section, the saturated working fluid was vaporized to the evaporator section and then the
1041 heat was released to the condensing area. The loop thermosyphon performance was limited in
1042 receiving a heat source in the evaporator section; due to the separated tube construction. For these
1043 reasons, Jengsooksawat et al., (2008) [94] designed the loop thermosyphon with chamber (LTVC) to
1044 solve the weak points in the limitation of heat being received in the evaporator area and the limitation
1045 of the condensing area. The TPCT made adjustments to for the problems which used the LTVC. Then,
1046 Jengsooksawat et al.,(2014) [95] reported about loop thermosyphon with vapour chamber. It was
1047 found that, the data showed that the LTVC yielded the valve of the relative of thermal efficiency of
1048 about 1 at R-11, a filling ratio of 60%, a velocity of 0.5 m/s, and an aspect ratio of 2.5 in the study
1049 conditions. It was further found that the larger vapour chamber was superior in the rating of the heat
1050 transfer over other vapour chambers in all experimental conditions in this study. Moreover, It was
1051 solved the weak points in the limitation of heat being received in the evaporator area and the
1052 limitation of the condensing area with chamber.

1053 Previous research studies on TPCT and ALT/CV have rarely focused on nanofluids
1054 containing surfactant and application of check valve. Check valve can solve problems of reverse flow
1055 in a tube. It can control direction of flow and separately manage flow of liquid and vapour. Moreover,
1056 nanofluids containing surfactant can resolve agglomeration issues and increase thermal performance
1057 of the ALT/CV.

1058 **2. Experimental apparatus and procedure**

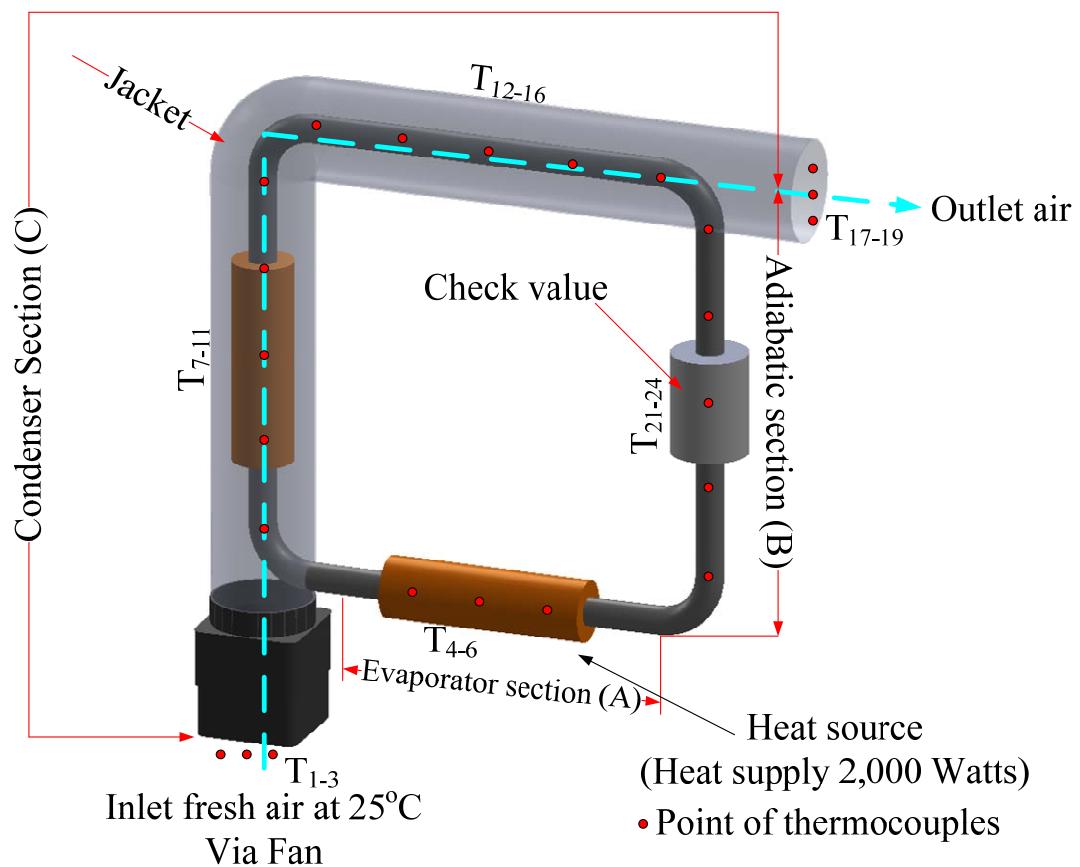
1059 This section describes the experimental set up, theparameters of the study, and theprocedure
1060 as shown in Table 1. Thus, the total number of variables used in the experiment was 2,430
1061 (5×3×3×3×3×6). The experiment had repeated threefold.

1062 Nanofluidsareproduced using themethod discussed in Chapter III andare suspended in a base
1063 fluid such as water. The nanofluid preparation involves many steps, such as changing the pH valve of
1064 the suspension, and using surfactant activators and ultrasonic vibration. The nanoparticles suspended
1065 in the base fluid are stable for long periods of time. In this research, thenanofluidswevereobtained using
1066 theultrasonic vibration method anda sonicator(bath type, operating frequency, and power source of the
1067 sonicator were 43 kHz and AC100 ~120V/AC220~240V 50/60 Hz, respectively) for 12 hours. The
1068 nanoparticles (Silver nanopowder is <100 nm particle size, 99.9% metals basis), oleic acid (OA), and
1069 potassium oleate(OAK⁺)were purchased from SIGMA-ALDRICH, Inc. in the USA. The silver
1070 nanoparticles were suspended in deionized water with a concentration of 0.5 wt% (NP). After that, the
1071 NP contained OA and OAK⁺ at concentrations of 0.5, 1 and 1.5 wt%, respectively.

1072 Figure 30 shows a schematic diagram of the experimental apparatus, which consisted of an
1073 ALT/CV. Section A, as the evaporator, is the heat source with a heat supply of 2,000 Watts: section B
1074 is the adiabatic; and section C, as the condenser, is the heat sink with fresh air (controlled at 25°C).
1075 The TPCT was made from copper tubes with an internal diameter of 12.70 mm. The ALT/CV had
1076 three sections, an evaporator, adiabatic, and condenser, of equal loop size of 30 cm, 40 cm, and 50
1077 cm, with an installation fin of 8 FPI in thecondenser section.Twenty-four temperatures were
1078 controlled and measured in the adiabatic section, which was constant at $\pm 4^{\circ}\text{C}$ points of thermocouple
1079 Type K (OMEGA with $\pm 0.1^{\circ}\text{C}$ accuracy). The thermocouplewere installed along with a data logger
1080 attached to the ALT/CV (Yokogawa DX200 with $\pm 0.1^{\circ}\text{C}$ accuracy, 24 channel input, and a -200°C to
1081 1100°C measurement temperaturerange) with the circle symbol point shown in Figure 30. The jacket
1082 was covered with insulation. Fresh air flow and velocity werecontrolled by the use of an AC motor.

1083 The controlled and variable parameters are shown in Table 3. During the experiment, the air velocity
1084 was set at 0.6 m/s (25°C) in order to calculate the heat transfer characteristics of the ALT/CV using
1085 the calorific method. The following equations were used for calculating of the heat transfer rates and
1086 for error analysis [58].

1087 The following equations were used to calculate the heat transfer rate and for error analysis
1088 recently compiled by Paramatthanuwat et al., (2010) [3].



1089

1090 Figure 30 Schematic diagram of experimental apparatus

1091 **3. Results and discussion**

1092 **3.1. Temperature distribution of loop thermosyphon**

1093 Figure 31 show the adiabatic temperature at the upper part and the side of the ALT/CV at
1094 different times, where the temperature showed a positive trend. The maximum temperature difference
1095 of both was $\pm 5^{\circ}\text{C}$ at T_{inlet} and T_{outlet} with NP containing 1 wt% of OAK⁺ as a working fluid and heat
1096 input at OPHS of 100%. T_{inlet} was higher than T_{outlet} and vice versa, causing higher thermal
1097 performance. The adiabatic temperature of the ALT/CV was able to reach a temperature higher than
1098 the ambient temperature, so heat transfer. The working fluid and the position of heat source were
1099 important, as they led to thermal performance [96]. In addition, the start up to 400 sec
1100 showed transient transfer and then at 400 sec to the end showed a steady state. Accordingly, the
1101 transient transfer that occurred from the unequally-filled working fluid flow as vapour + liquid in the
1102 loop and the passed check valve of the thermosyphon resulted in the nonuniform distribution of the
1103 temperature. The point of steady state, the ball moving through a saturated vapour and superheat
1104 experiences a force in direction opposite to its motion. The terminal velocity is achieved when the
1105 drag force is equal in magnitude but opposite in direction to the force propelling the object. The effect
1106 of ball buoyancy in ALT/CV, the pressure acting motion ball is in upward direction, according to Eq.
1107 (51) then the ball acting motion is downward direction according to (52). The force direction
1108 due to the pressure difference (ΔP) that flow was separated vapour with liquid. This relation was
1109 observed and defined in the heat transfer mechanism to generate pressure at evaporator section. In this
1110 case, the $P_{upward} \geq F_{ball\ total}$ caused ball buoyancy affect to T_{inlet} and T_{outlet} had a difference (ΔT).
1111 Consequently, the ALT/CV helped to regulate and control the $\Delta P \propto \Delta T$ for ALT and check valve
1112 operation, as shown in Figure 32.

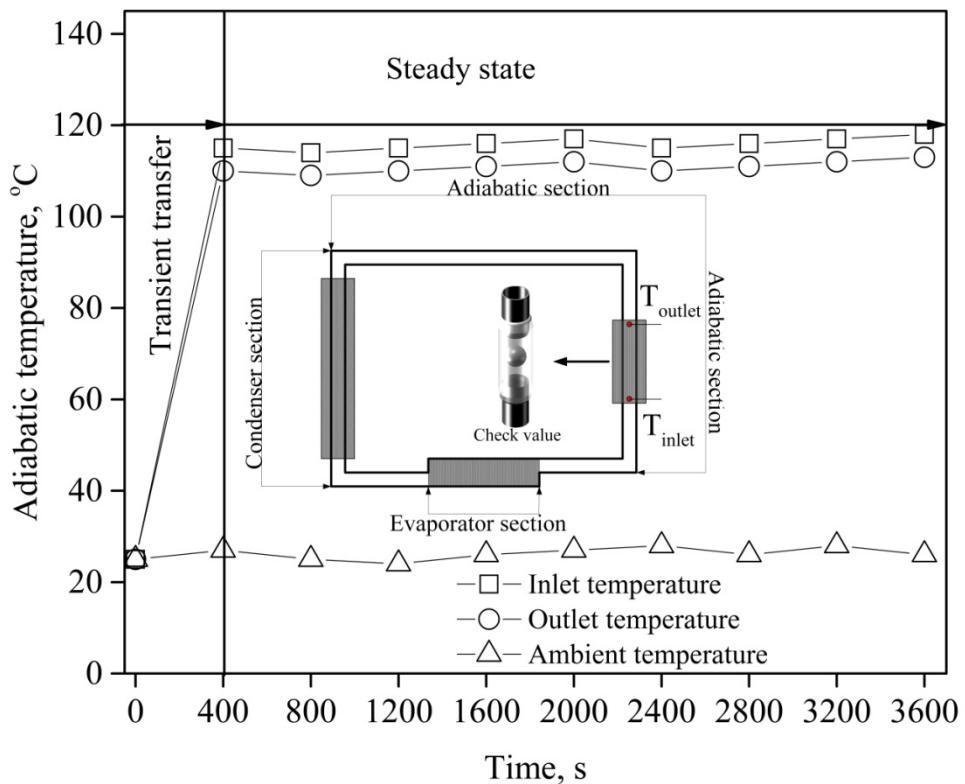
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$$\frac{F_{upward}}{A_{ball}} > \frac{(-F_\tau - F_\sigma - F_g + F_d)_{ball}}{A_{ball}} \quad (51)$$

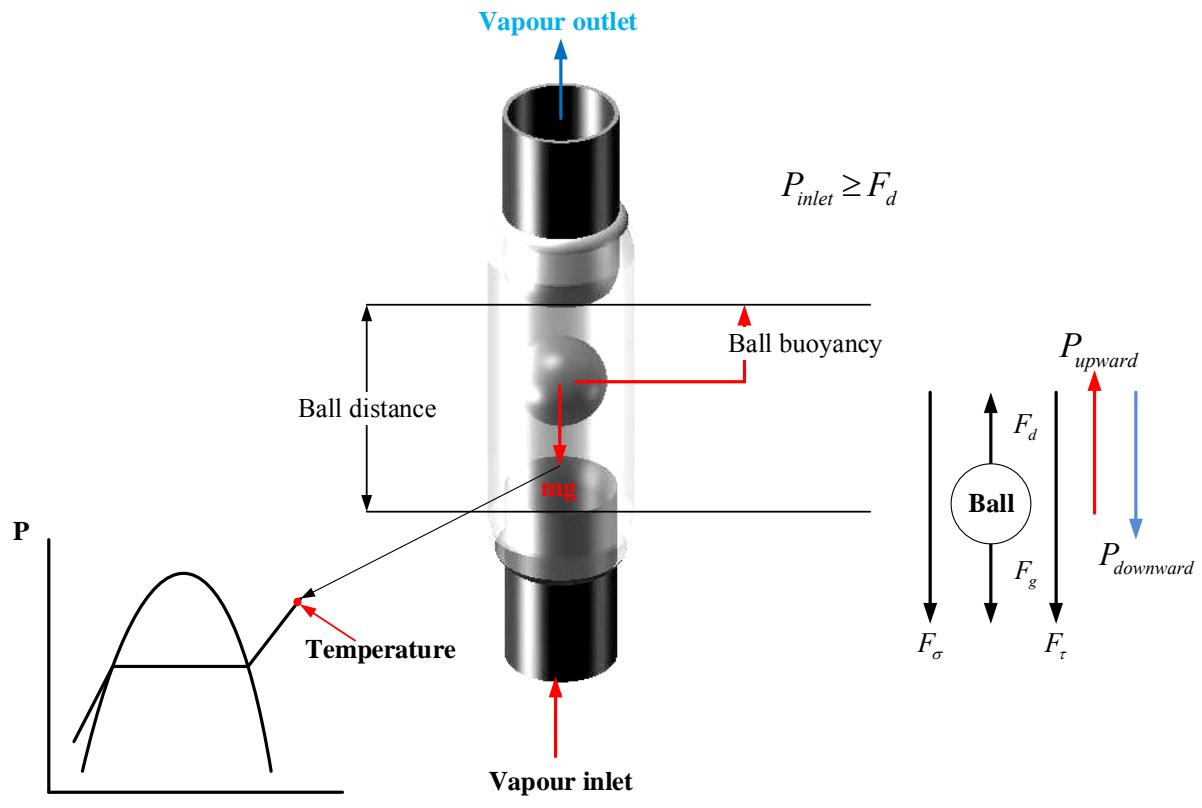
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$$\frac{F_{downward}}{A_{ball}} < \frac{(F_\tau + F_\sigma + F_g - F_d)_{ball}}{A_{ball}} \quad (52)$$



1116

1117 Figure 31 Temperature distribution of loop thermosyphon in the adiabatic section



1118

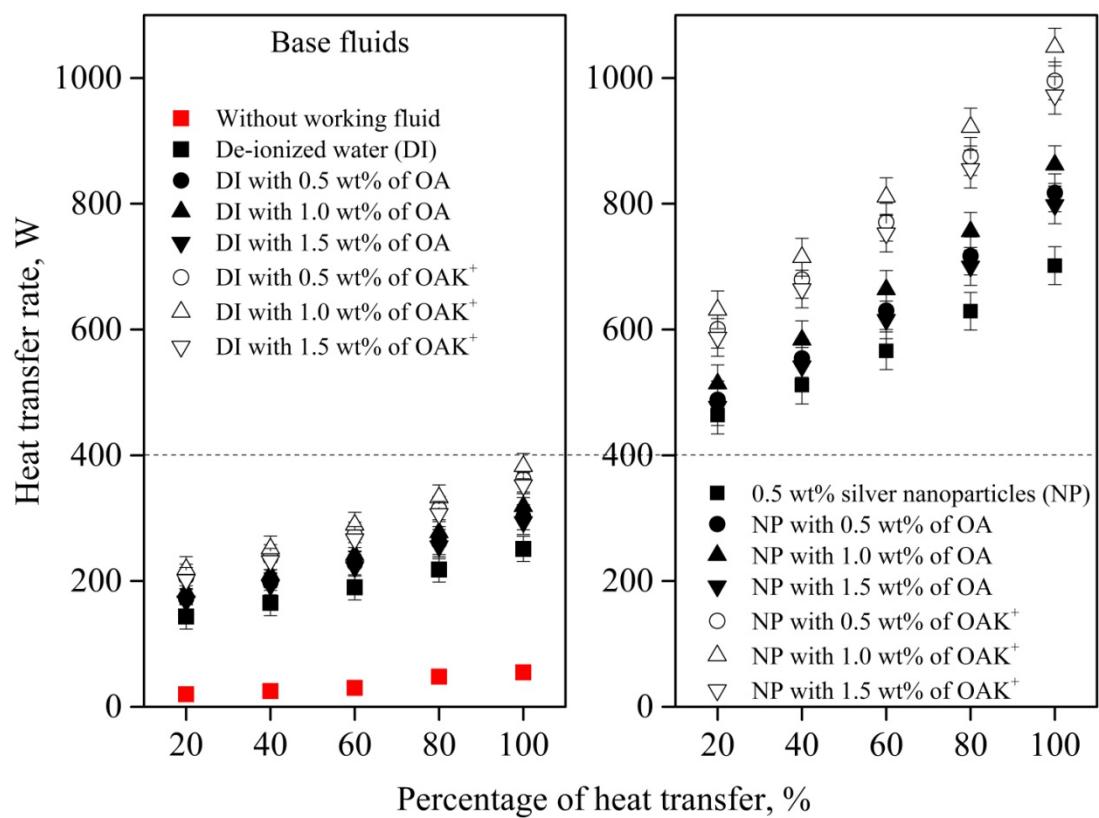
1119 Figure 32 Check valve operation

1120 **3.2. Effect of operating percentage of heat supplied (Heat source)**

1121

1122 Figure 33 shows the heat transfer of the ALT/CV at a loop size of 50 cm as a function of the operating
 1123 percentage of the heat supplied (OPHS), filled with fourteen working fluids. The data shown
 1124 correspond to the filling ratio of each working fluid of 50% of evaporator volume. The
 1125 results indicate that the NP contained 1 wt% of OAK⁺ which produced a heat transfer rate valve of
 1126 1,150W at OPHS of 100%. Then, all of experimental data were compared with the heat transfer rate
 1127 of the ALT/CV without the working fluid or base fluid. In all cases the heat transfer rate of the NP
 1128 contained 1 wt% of OAK⁺ and showed superior performance over other parameters throughout all
 1129 experimental conditions in this study. It can be observed that for a given working fluid, the heat
 1130 transfer rate of the base fluid and all of the NP and NP containing surfactant increased along with

1131 anincrease of the OPHS, which confirms that the OPHS had a strong effect on the heat transfer rate
 1132 and the working fluid properties. It was therefore determined that the peak heat transfer rate occurs
 1133 when the percentage of heat supplied is 100%. Moreover, the viscosity and surface tension were
 1134 found to decrease after the introduction of OA and K^+ [13, 45]. The viscosity and surface tension of
 1135 the working fluids decreased as a result ofthe increase in operating temperature that theworking
 1136 fluidis easier to boil. Furthermore, the OA would allow the particles to disperse uniformly within the
 1137 base fluid, but high OA concentration appearedto hinder the aggregation ofthe NP, which was
 1138 observed at the bottom of the liquid [44, 46]. The outcome of this experiment by raising the heat
 1139 transfer rate is according to Hwang et al.,(2008) [45], Sharma et al., (2011) [44] and Li et al., (2008)
 1140 [46]. In this study, the optimal concentration of OAK^+ added to the NP was 1 wt%, having the highest
 1141 heat transfer rate throughout all conditions.

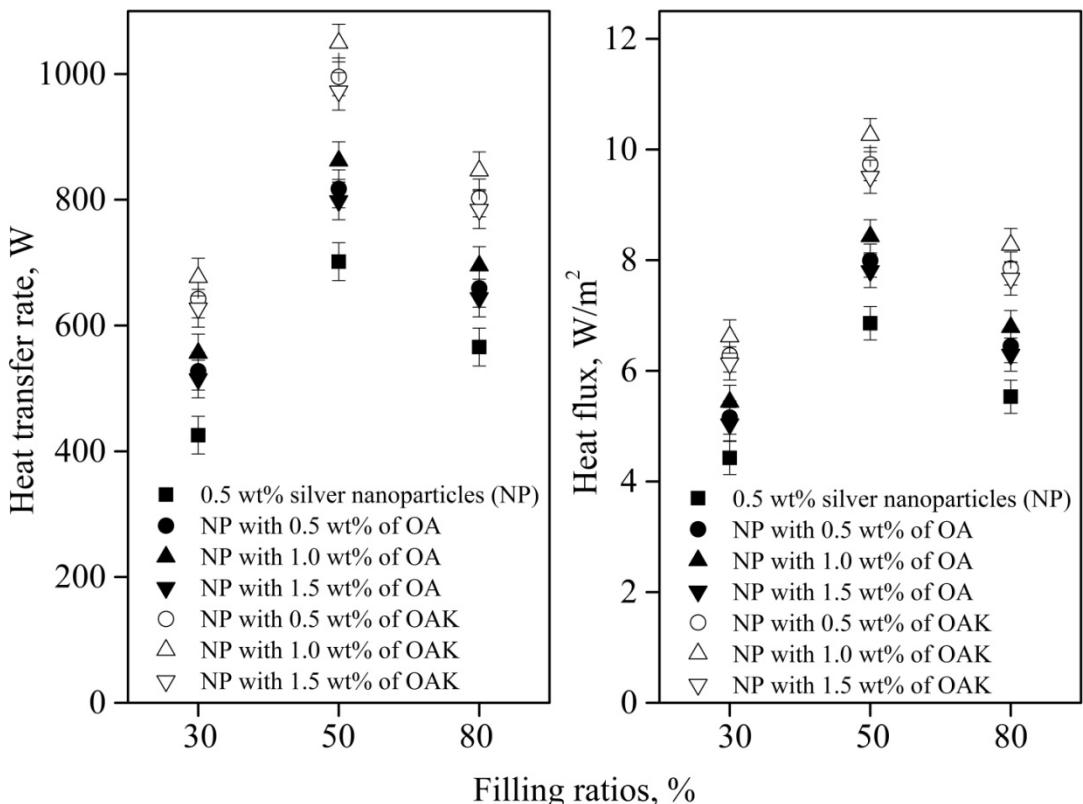


1142

1143 Figure 33 Effect of operating percentage of heat supplied (Heat source)

1144 **3.3. Effect of working fluid ratios**

1145 Figure 34 shows the heat transfer rate and heat flux of the filling ratio. The comparative heat
1146 transfer rate and heat flux among the 3 filling ratios were defined by 30, 50, and 80% with the
1147 ALT/CV at a loop size of 50 cm. The result of the heat transfer rate and heat flux showed a maximum
1148 value of 1,150 W and 10.5 W/m² at the filling ratio of 50%, OPHS of 100%, and NP containing 1 wt%
1149 of OAK⁺. It can be observed that the heat transfer rate and heat flux experienced a peak as a function
1150 of the filling. The optimum filling ratio for the addition of the working fluid in the ALT/CV was 50%,
1151 where critical film thickness occurred and the highest heat transfer rate was achieved [4, 16, 87, 97].
1152 Moreover, the NP could be helpful for increasing thermal performance because of the particles that are
1153 increased in the base fluids. The suspension of the nanoparticles could increase the surface areas of
1154 the working fluid and heat capacity of the base fluid. Furthermore, the optimization of the chain
1155 length of the oleic acid was 18 with K⁺, which was effective for particle dispersing stabilization. The
1156 optimized chain length also improved the colloidal stability and increased the non-precipitation period
1157 so that the nanoparticles could be uniformly dispersed [4, 35, 45]. The NP containing oleic acid was
1158 superior in its thermal performance over the deionized water of approximately 80% in all the
1159 experimental conditions in this study.



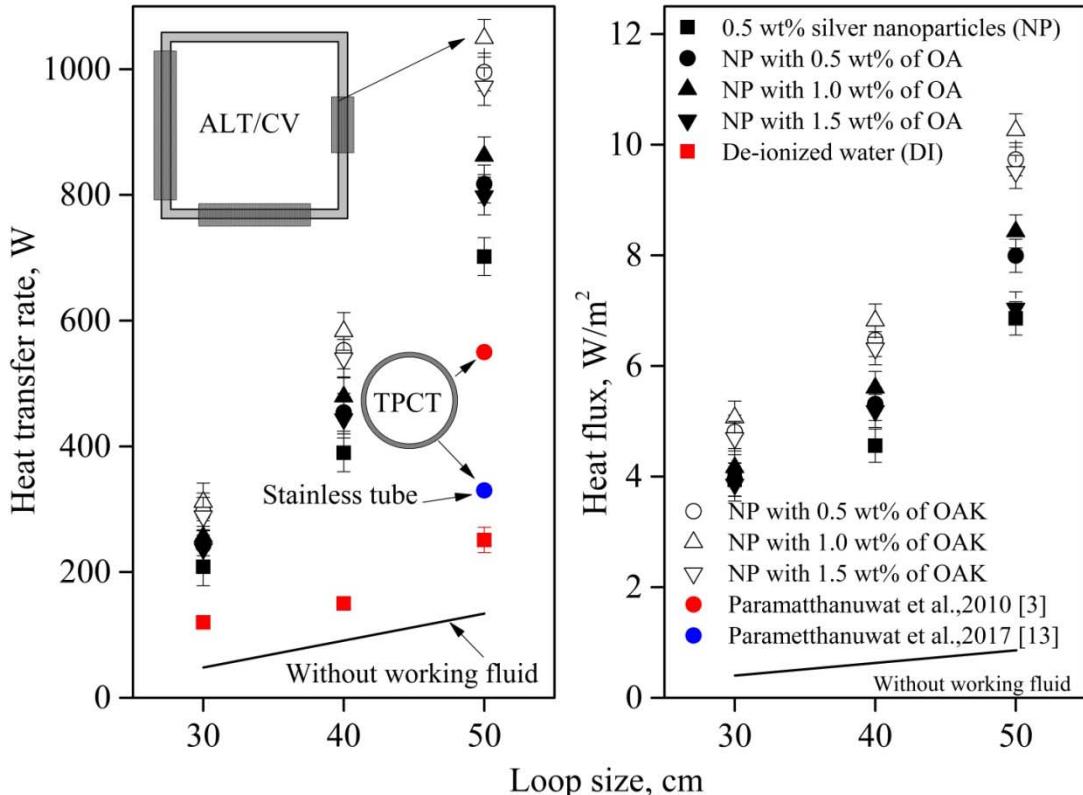
1160

1161 Figure 34 Effect of working fluid ratios

1162 **3.4. Effect of the loop size of the thermosyphon**

1163 Figure 35 shows the heat transfer rate and heat flux of the loop size of the thermosyphon. The
 1164 experimental results clearly indicated the aspect ratios with the use of NP containing 1 wt% of OAK⁺,
 1165 producing the maximum heat transfer rate values of 1,150 W with a loop size of 50 cmOPHS of
 1166 100%. Furthermore, the 1 wt% of OAK⁺ had the highest heat transfer rate more so than the TPCT used
 1167 by Paramatthanuwat et al., (2010) with NP, $\frac{L_e}{d_i} = 20$ ID 25.4 mm at $T_v = 50^\circ\text{C}$ [3] and
 1168 Paramatthanuwat et al., (2017) with RTPCT from the stainless tube with NP containing 1 wt% of OA
 1169 at OPHS of 100% [13]. However, the current experimental results contrasted with those of
 1170 Paramatthanuwat et al., (2010) and Paramatthanuwat et al., (2017). Thus, it is important to note the
 1171 vast differences among different experimental conditions, especially in regard to the type of heat pipe

1172 and thermosypon. Accordingly, it can be seen that while the loop size increased from 30 cm to 50 cm,
 1173 the heat transfer rate behaviour only slightly increased. Thus, the increase in the loop size is
 1174 equivalent to the increase in the evaporator section size of ALT/CV, which will result in increased
 1175 heat transfer rate behaviour. The larger evaporator section led to pool boiling, thus increasing the heat
 1176 transfer rate [3, 16, 25]. On the other hand, the small aspect ratio led to the boiling in the confined
 1177 channel, thus lowering the heat transfer rate [16]. Thus, the loop size and evaporator section was
 1178 changed in terms of the ALT/CV's length, the filling ratio was changed, thus causing the thermal
 1179 received valve to change that depend on the proportion of ALT/CV's length [20]. In addition, the
 1180 silver nanoparticles are very small, so when NP contained oleic acid and K^+ , the particles were
 1181 dispersed uniformly in the fluid. This increased the surface area in the bubble diameter for absorbing
 1182 heat [98, 99], thus enhancing the boiling of the fluid. Therefore, it could be predicted that the amount
 1183 of condensate was so high that it could return to the evaporator section, thereby ensuring an ample
 1184 amount of working fluid for boiling and phase transition [34, 85, 100].

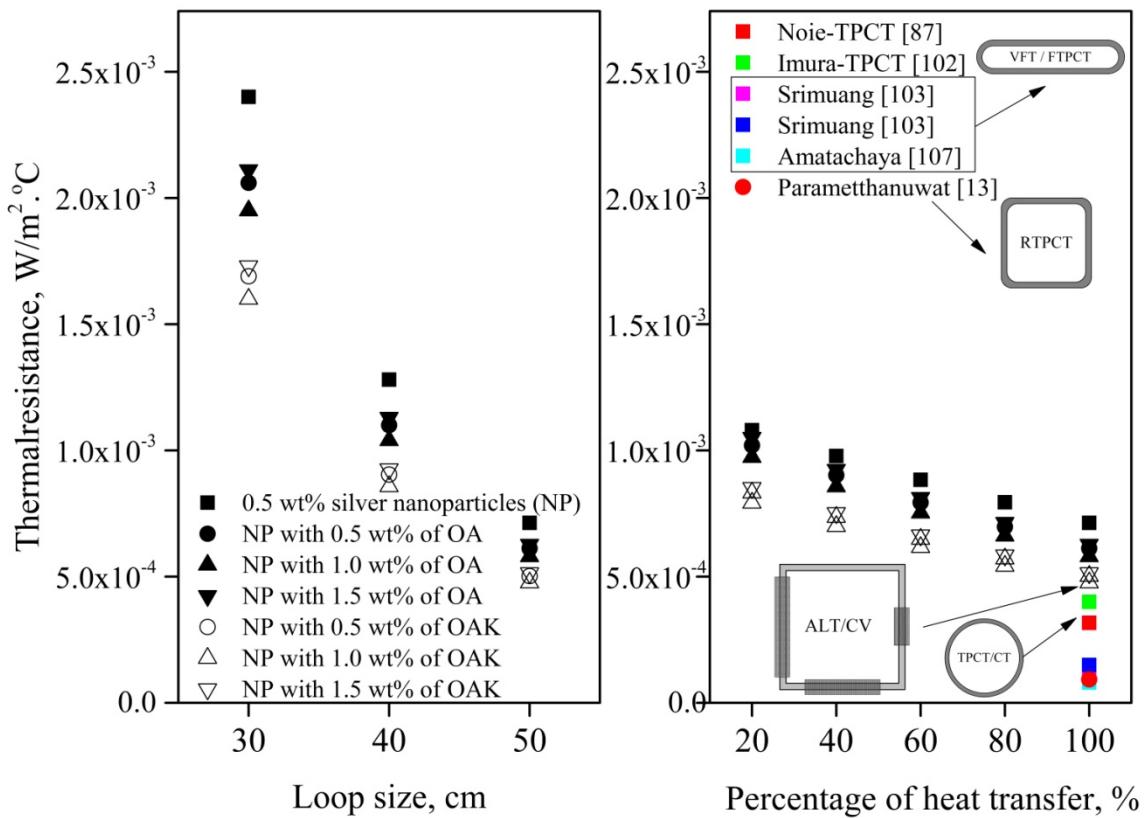


1185

1186 Figure 35 Effect of the loop size of the thermosyphon

1187 **3.5. Effect of Thermal resistance**

1188 The ALT/CV influenced the thermal transport to the evaporator section then boiling
1189 phenomenonoccurred. The thermal resistance contoured the pool boiling/film boiling of
1190 thermosyphoninthe evaporator section. The thermal resistance can be estimated from a number of
1191 ESDU81038 [15]. Thus, the nanoparticles existed in the base fluid containing surfactant, which
1192 affected the flow of the nanofluids, causing the surface tension and wettability to decrease but it was
1193 found thatthenanofluids containing surfactant produced high thermal property valves [35, 101]. Thus,
1194 the boiling phenomenon can be expressed according to thethermal/physical properties of the working
1195 fluids and physical of the thermosyphon, which was important for thermal resistance. Accordingly,
1196 when the cross sectional was changed it caused reduced friction loss and affected the phase transfer
1197 sophenomenon film boiling decreased [13]. However, the current experimental results contrasted with
1198 those of Noie (2005-TPCT) [87], Imura et al.,(1997-TPCT) [102], Srimuang et al., (2009-VFT and
1199 CT) [103], Amatachaya et al., (2010-FTPT) [104] and Parametthanuwat et al., (2017) [13], as shown
1200 in Figure 36, with the type of thermosyphon. In regard to this study, the results were achieved using a
1201 different heat source, working fluid, and condition. It was found that the minimum thermal resistance
1202 that occurred was $4.76 \times 10^{-4} \text{C/W}$ of a loop thermosyphon size of 50 cm, the NP containing 1 wt% of
1203 OAK+. It is obvious that while the loop size of thethermosyphon increased, the thermal resistance only
1204 decreased. The heat source was varied by changing the percent of heat supplied. It was further found
1205 that the thermal resistance of the ALT/CV washigher when increasing the heat source because the
1206 ALT/CV is a single loopand has a higher total area. Studies cited for this result werecompared with
1207 Noie (2005-TPCT) [87], Imura et al.,(1997-TPCT) [102], Srimuang et al., (2009-VFT and CT) [103],
1208 Amatachaya et al., (2010-FTPT) [104], and Parametthanuwat et al., (2017) [13] concerning thetype of
1209 thermosyphon (TPCT, CT, and VFT) according to themaximum value of each those conditions at
1210 maximum heat source. Moreover, the type of thermosypon was approached boiling phenomenon. The
1211 pool boiling phenomenon only occurred when the lower thermal resistance which was high thermal
1212 performance was reached. On the other hand, the pool boiling phenomenonwas approached inside the
1213 confined channel, which had higher thermal resistance.



1214

1215 Figure 36 Effect of Thermal resistance

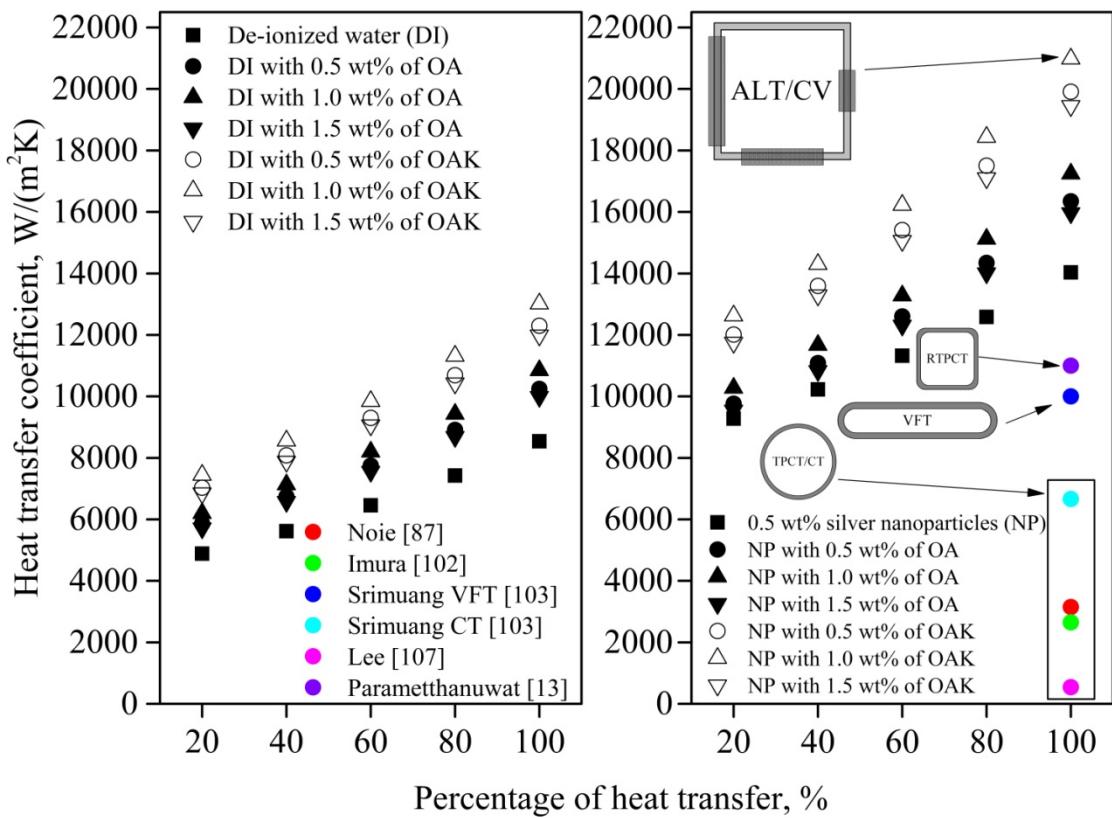
1216 **3.6. Effect of the heat transfer coefficient**

1217 Figure 37 shows the heat transfer coefficient (HTC). The ALT/CV was presented with a
 1218 loop size 50 cm, and an OPHS of 100%. It was observed that 50% of the filling ratio of the samples
 1219 showed similar positive trends. Furthermore, the HTC of the ALT/CV depend on filling ratios and NP
 1220 containing OAK⁺ was different when the surfactant concentration was 0, 0.5, 1 and 1.5 wt%. In all
 1221 cases the filling ratio of 50% with NP containing 1 wt% of OAK⁺ showed superior performance than
 1222 the other concentrations. The NP that contained 1 wt% of OAK⁺ had the highest HTC values of
 1223 20986.029 W/m²K. On the other hand, the HTC seemed to have an inverse effect on the thermal
 1224 resistance [10, 16]. Generally, it can be explained that the filling ratio and surfactant concentration
 1225 increased when the viscosity of the working fluids changed, resulting in the movement of the
 1226 nanoparticles. The Brownian motion of the nanoparticles inside the NP had a greater increase in

1227 dispersal, which in turn increased the convection transfer. Consequently, the convection-like effects
1228 remarkably increased, which lead to an increased HTC [105, 106].

1229 All of the heat transfer coefficient data increased with an increased percentage of heat
1230 supplied. The experimental results of this current research can be compared with the work of
1231 Parametthanuwat et al., (2017) [13], Noie (2005) [87], Imura et al., (1997) [102], Srimuang et al.,
1232 (2009) [103], and Lee et al., (2003) [107]. It can be seen that the heat transfer coefficients of the
1233 ALT/CV depended linearly on the percentage of heat supplied throughout all the samples, indicating
1234 that the heat transfer coefficients increased independently on the surfactant concentration. The
1235 enhancement of the heat transfer coefficients of the NP containing surfactant was different when the
1236 surfactant concentrations were 0.5, 1 and 1.5 wt%. In all cases, the DI water containing surfactant and
1237 NP containing OAK⁺ showed superior performance than the base fluids. However, in all cases, the
1238 ALT/CV showed superior performance than the TPCT, CT, and VFT. The results between the five
1239 studies cited show an agreement between the difference in the thermosyphon type and cross-sectional
1240 and physical conditions. However, the heat transfer coefficients were higher than in the five studies
1241 cited because of the ALT/CV was installed check valve for controlling the liquid and vapour, which
1242 was made those for separating. This may have been due to the improvement pool boiling phenomenon
1243 in the evaporator section and the condensation phenomenon in the condenser section. Hence the heat
1244 transfer coefficients depend on the phase change transfer. Moreover, the nanoparticles (NP) dispersed
1245 in the liquid can increase the surface area for heat absorption. In the case of the NF containing OA
1246 and K⁺, the OA and K⁺ will decrease the surface tension of the NP, and stabilize the NP by uniformly
1247 distributing NP and increase the interface area of the nanofluids with deionized water [39]. The surface
1248 tension has a significant influence on the thermal process since it depends on property and interfacial
1249 equilibrium [108]. A high OA and K⁺ concentration appeared to hinder aggregation, and entanglement
1250 of the NP was observed at the bottom of the liquid [44, 109]. According to our experimental results, 1
1251 wt% OAK⁺ was enough to homogeneously disperse the NP and produce efficient thermal transfer
1252 between the particles and de-ionized water. This consequently resulted in the highest thermal

1253 properties enhancements [43, 76, 77, 110]. NP and was dominated by the pool boiling phenomenon due
 1254 to the large number of heat transport inside the ALT/CV.

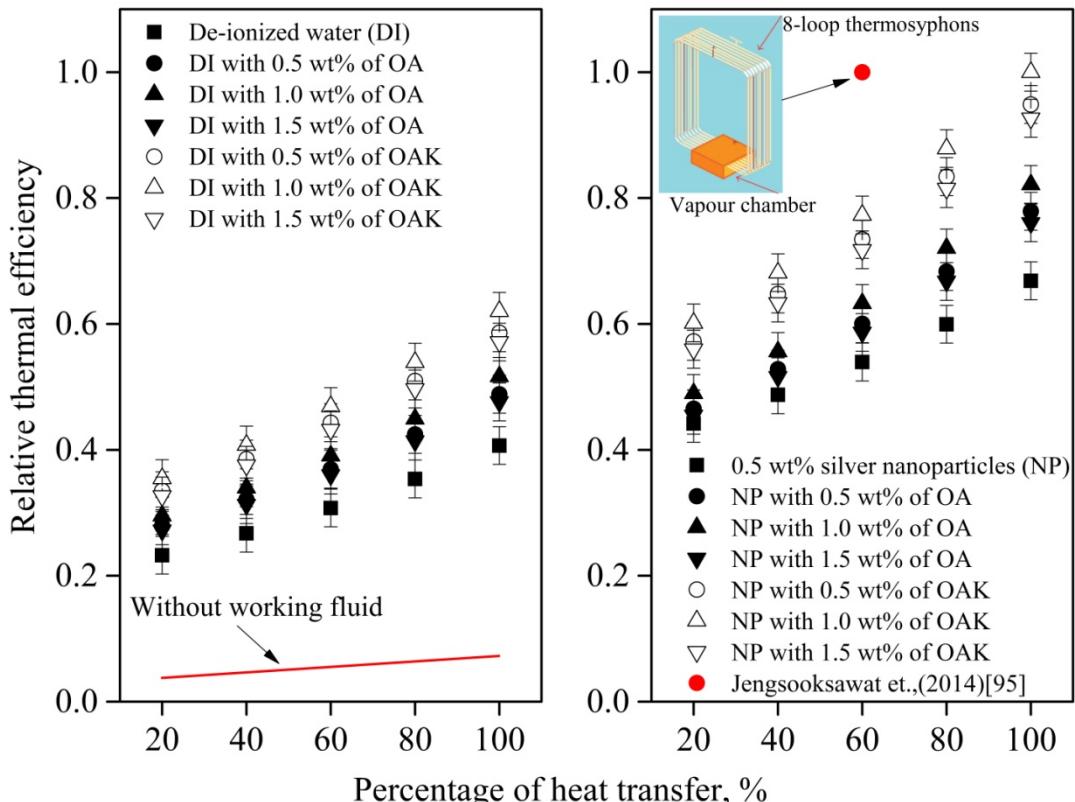


1255

1256 Figure 37 Effect of the heat transfer coefficient

1257

3.7. Effect of the relative of thermal efficiency



1258

1259 Figure 38 Effect of the relative of thermal efficiency

1260 Figure 38 shows the relative thermal efficiency (RTE) of all working fluidsfiled. The RTE
 1261 was presented with a loop size of 50 cm at a filling ratio of 50% and an OPHS of 100%. According to
 1262 our experiment, the 1 wt% of OAK⁺ seemed to produce the maximum of RTE. The check valve of the
 1263 loop thermosyphonwas helping the highest heat transfer behaviour as this check valve could separate
 1264 and control the flow of the liquid and vapour[111-113]. However, the current experimental results
 1265 contrasted with the results of Jengsooksawat et al., (2014) [95]. Studies cited for this result
 1266 were recompared with Jengsooksawat et al., (2014) [95], was used 8 loop thermosyphons with OPHS of
 1267 1,200 W, an air velocity of 0.5 m/s, and a filling ratio of 60% of ethanol as a working fluid. Thus, it is
 1268 important to note the vast differences among different experimental conditions, especially in regard to
 1269 the type of heat pipe and thermosypon. In addition, the thermal motion of the nanoparticles enhanced

1270 the thermal conductivity of the nanofluids. The OA and K^+ helped the homogeneous dispersion of the
1271 nanoparticles in the nanofluids. Moreover, Brownian motion significantly contributed to the rise in the
1272 thermal conductivity of the nanofluids. Thus, the thermal conductivity was superior, it had the heat
1273 transfer rate mechanism efficiently over thermal diffusion in the fluid [34, 43].

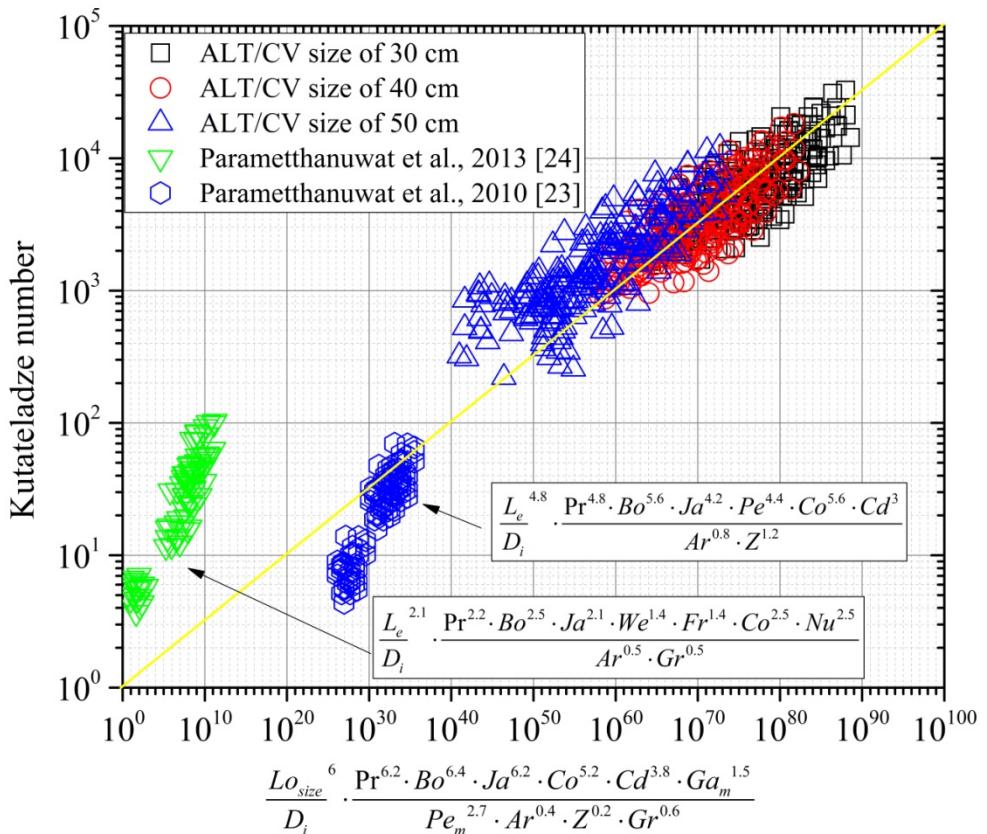
1274 **3.8. Dimensionless**

1275 **3.8.1. The effect of the dimensionless group on Kutateladze number (Ku)**

1276 The dimensionless parameters on heat transfer characteristics of the ALT/CV; thus, $\frac{Lo_{size}}{D_i}$,

1277 $Pr, Bo, Ja, Co, Cd, Ga_m, Pe_m, Ar, Gr$ and Z values can be used to formulate a correlation to
1278 predict the heat transfer rates of the ALT/CV. The standard least square curve fitting technique was
1279 adopted. The power function was found to be the best representation to formulate a correlation.
1280 The correlation of the Ku number to predict the heat transferred from the ALT/CV was:

$$1281 Ku = 3.24 \left[\frac{Lo_{size}}{D_i}^6 \cdot \frac{Pr^{6.2} \cdot Bo^{6.4} \cdot Ja^{6.2} \cdot Co^{5.2} \cdot Cd^{3.8} \cdot Ga_m^{1.5}}{Pe_m^{2.7} \cdot Ar^{0.4} \cdot Z^{0.2} \cdot Gr^{0.6}} \right]^{1.57} \quad (53)$$



1282

1283 Figure 39 Relationship between the dimensionless group and Kutateladze number

1284 The results are shown in Figure 39. It was further found that the filling ratio had no effect on
 1285 the ratio of the heat transfer rates in the vertical position, but the properties of the working fluid
 1286 affected the heat transfer rate. Furthermore, when the dimensionless groups were compared with the
 1287 results of Parametthanuwat et al., (2013) [24] and Parametthanuwat et al., (2010) [23], it was found
 1288 that the dimensionless group of the ALT/CV was higher than determined of them. In regards to this
 1289 study, the results were achieved using the silver nanoparticles; however, there was a difference in the
 1290 surfactant and the type of thermosyphon. Thus, it is important to note the vast differences in
 1291 experiment conditions, especially in regards to the method of heat transfer rate. In addition, a
 1292 correlation for predicting the heat flux for the ALT/CV in the vertical position has been established.
 1293 Moreover, the coefficient of determination (R^2) of this equation was 0.8.

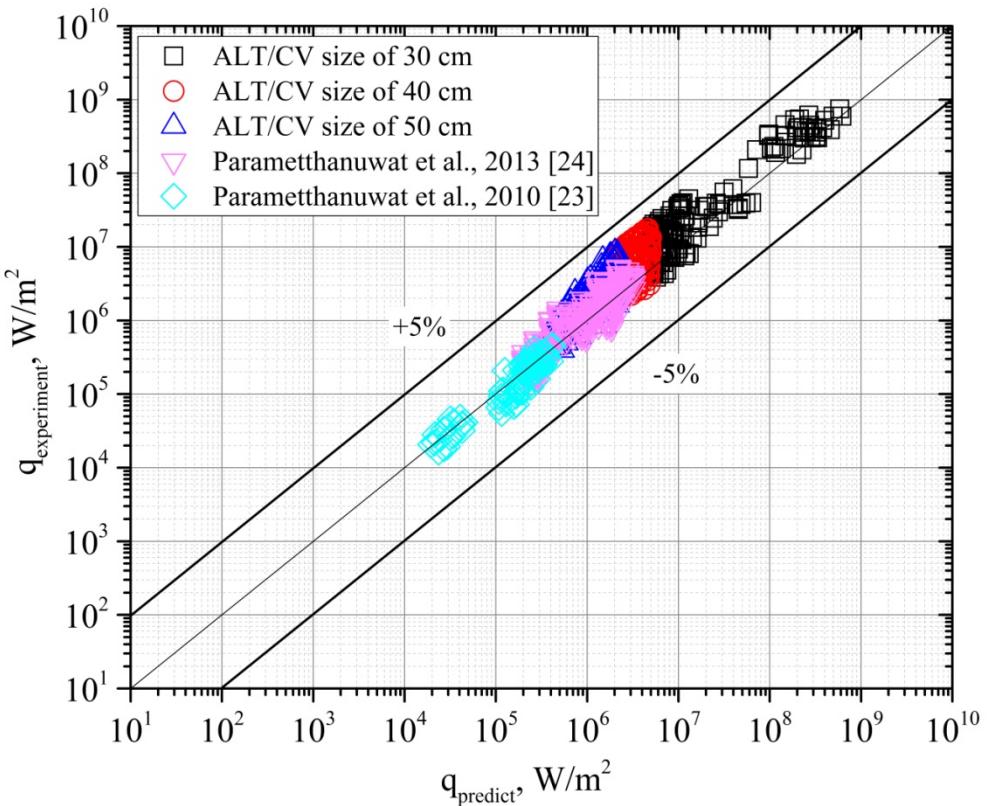
1294

3.8.2. Correlation equation

1295 From the results obtained, the heat flux of the ALT/CV positioned vertically can be
 1296 evaluated from equation (54). As can be seen in Figure 40. Parametthanuwat et al., (2010) [23] used
 1297 TPCT without surfactant and Parametthanuwat et al.,(2013) [24] used TPCT with OA, the heat flux
 1298 was closed. Then, ALT/CV used NP containing OAK^+ , had a heat flux similar trend with TPCT. The
 1299 standard deviation (STD) of the experimental heat flux and the predicted heat flux using equation (29)
 1300 was $\pm 5\%$. Therefore, it can beconcluded that this equation can be used to predict the heat flux of the
 1301 RTPCT and TPCT, as shown in Figure 40.

1302

$$q = 3.24 \left[\frac{Lo_{size}^6 \cdot Pr^{6.2} \cdot Bo^{6.4} \cdot Ja^{6.2} \cdot Co^{5.2} \cdot Cd^{3.8} \cdot Ga_m^{1.5}}{Pe_m^{2.7} \cdot Ar^{0.4} \cdot Z^{0.2} \cdot Gr^{0.6}} \right]^{1.57} \times \left[\rho_v h_{fg} \left(\frac{\rho_v - \rho_l}{\rho_v^2} \right) \right]^{0.25} \quad (54)$$



1303

1304 Figure 40 Comparison between $q_{predict}$ and $q_{experiment}$

1305 **4. Conclusion**

1306 This chapter examined the thermal behaviour in an ALT/CV and in particular emphasized a
1307 new “loop type of thermosyphon installed check valve” design. It was divided into a number of
1308 sections to demonstrate the thermal behavior, as follows:

1309 - The ALT/CV at a loop size of 50 cm yielded better heat transfer behaviour with a filling ratio
1310 of 50% with respect to the evaporator volume when the NP with 1 wt% of OAK⁺ was
1311 used as a working fluid and the percentage of the heat supplied was 100%. It was further
1312 found that the loop size of 50 cm and a filling ratio of 50% with respect to the evaporator
1313 volume were superior in thermal behaviour over other parameters throughout all
1314 experimental conditions in this study, with a relative thermal efficiency of about 1.

1315 - In the dimensionless analysis, it was found that the filling ratio did not affect the heat
1316 transfer rate; however, the properties of the working fluid did [23, 24]. Therefore, these
1317 properties dominated the dimensionless parameters and the Ku correlation and showed
1318 that the standard deviation for predicting heat flux was $\pm 5\%$.

1319 Finally, in conclusion, the ALT/CV led to an enhancement and produced a better
1320 performance than the normal type of thermosyphon. The loop type of thermosyphon installed check
1321 valve is a new type of thermosyphon, which is appropriate for use in heat transfer engineering. The
1322 presence of OA and K⁺ clearly contributed to the rise in the heat transfer rate and the improvement of
1323 nanofluid properties.

1324

1325

CHAPTER V APPLICATION OF LOOP THERMOSYPHON

1. Introduction and Literature review

The dimensionless is formulated from Chapter IV to correlate and predict the heat transfer characteristic of the ALT/CV to Kutateladze number (Ku). The silver nanofluids containing OAK⁺ surfactant in a prototype of oven an advanced loop thermosyphon with check valve (OALT/CV) are then applied. The OALT/CV used in this study was a special type which uses nanofluids in the loop thermosyphon to transfer heat from the evaporator to condenser without external energy requirement. The primary objective of this study is to design and test the OALT/CV so that it will increase the heat transfer to water. The heat will be helpful to distribution the temperature profile in the OALT/CV which is regularly. This OALT/CV was designed using a correlation of Kutateladze number (Ku). This review can be summarized from the challenges of the on-line characterization of thermal performance, rheological behaviour and OALT/CV.

Da Silva et al., [114] used TPCT in cooking chambers using a real size prototype. The TPCT has 1,200 *mm* of total length with water as working fluid and filling ratio of 60%. It was found that the temperature inside the prototype was shown to be highly uniform due to the large area of the condenser which was responsible for heating of the cooking chamber. Therefore, the incidence of radioactive energy over any product placed in the cooking chamber will be highly uniform, avoiding over/under cooked products. Based on that, the model shows the internal temperature and the prototype can be determined theoretically with good precision. Moreover, Milanez [115] studied TPCT heat enclosure with the dimensions 0.38×0.48×0.61 *m*, eight stainless steel, and added water as the working fluid. It was found that the enclosure heated using the TPCT had more uniform temperature and radioactive heat transfer coefficient distributions compared to the conventional approach. The conventional enclosure tends to present larger convective heat transfer coefficients than the thermosyphon assisted enclosure because of movement of the exhaust gases. The radiation field inside the thermosyphon assisted enclosure is more uniform than the conventional enclosure. A more

uniform radiation field is obviously better when applying the enclosure for cooking purposes as it cooks more evenly. Nanofluid plays an important role in improving TPCT's heat transfer. In 2010, Parametthanuwat et al., [116] applied TPCT for energy conservation containing silver and gold nanofluid at the 0.5 wt% concentration. It was found that the TPCT which used silver nanofluid appeared to have uniform temperature distribution. Consequently, processing time and LPG consumption could be reduced by 10 *min/unit* and 1.8 *kg/unit*, respectively. The original oven was 37.1% more effectiveness after the installation of the TPCT and was 69.8% more effectiveness when using silver nanofluids. Obviously, the installation of the TPCT containing nanofluid was helpful to thermal distribution and improved the overall thermal effectiveness. Then, Kiniman et al., [117] improved the top heat mode closed loop oscillating heat pipe with a check valves air pre-heater (THMCLOHP/CV/AP) for chilli drying. The THMCLOHP/CV/AP showed the highest effectiveness of 0.1 at an operating temperature of 80°C with an air velocity of 0.5 *m/s*. The processing time used for drying chilli was 2 hours and 20 minutes. The quality of the colour measurement of the chilli exceeded the marketplace with the total colour difference (ΔE^*) being significantly different ($p \leq 0.05$). Moreover, Topuz et al., [118] studied the effect of methods and storage on the colour of paprika with the Refractance Window Drying method (RWD). Thus, the colour statistical analyses pointed out were significantly different ($p < 0.05$), according to Kiniman et al., [117]. Finally, the quality of the product such as colour, oven time and energy thrift have been important in the oven process.

2. The ordinary oven chilli

The ordinary chilli oven is usually driven by solar method, fluidized bed and etc. An oven without a heat pipe represents lower operational cost, but an oven with a heat pipe is considered economical and easier to operate. Something to note is that the chilli oven method rejects energy at high temperature, resulting in energy losses which is under moisture standard lower at 13% wet basic. Also, to guarantee reasonable temperature distribution, a ventilator, similar to the one employed to ovens with a heat pipe, is used, according to Kiniman el al., [117]. A diagram of the installation of a ALT/CV is shown in Figure 41 and process oven diagram shown in Figure 48.

3. The experimental apparatus analysis

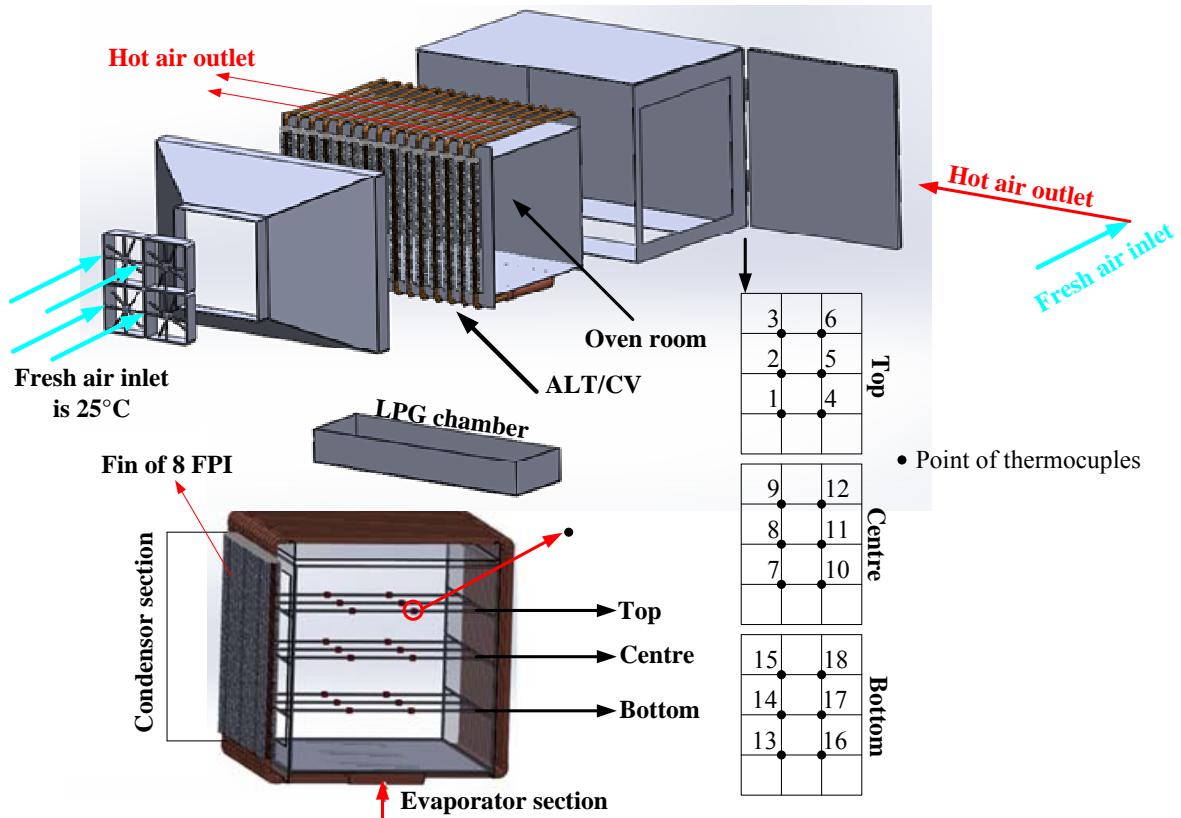


Figure 41 The prototype of oven an advanced loop thermosyphon with check valve (OALT/CV)

2.1. Chilli oven design

The prototype of oven an advanced loop thermosyphon with check valve (OALT/CV) in Figure 41 was designed using the Kutataladza number (Ku). The above-stated dimensionless numbers were correlated with Ku form Equation (54), to calculate the convection heat transfer capacity of one tube. From Equation (53), the heat flux of TPCT at a vertical position can be evaluated from Equation (54). The calculation showed that the number of tubes for OALT/CV is 15 loops.

Figure 41 shows the dimensions of the OALT/CV and Table 4 shows condition study. The loop thermosyphon was combined with a vapour chamber that could be separated from the loop thermosyphon (adiabatic section and condenser section) and the vapour chamber (evaporator section). The fifteen loop thermosyphons with check valve were made from copper tube with an inside diameter of 12.70 mm. The OALT/CV had a dimension size of 600×600×600 mm (W×L×H) with an 15 loop thermosyphon with check valve length size of the adiabatic, condenser and evaporator sections being 1000, 500 and 500 mm respectively. The fin was assembled as 8 FPI of the area of the condenser in order to improve heat exchange. The heat was removed from the condenser sections with convective forced heat transfer that was blown through this section. Air flow and velocity was controlled by the use of an AC motor. The fifteen loop thermosyphons with check valve were connected to a vapour chamber as an evaporator section.

2.2. Temperature distribution measurement in oven room

This section focuses on temperature distribution inside the OALT/CV which uses the conventional approach as shown in Figure 41. Figure 41 presents the temperature reading points used to monitor the temperature distribution inside a baking oven. Eighteen thermocouples were installed for data collection (Yokogawa DX200 with $\pm 0.1^\circ\text{C}$ accuracy, 30 channel input and -200°C to $1,100^\circ\text{C}$ measurement temperature- range) and were used with type K thermocouples (OMEGA with $\pm 0.1^\circ\text{C}$ accuracy) [116]. Then, all the data from Figure 41 inside the oven was used in a computer program to determine the temperature contour.

2.3. Quality of chilli analysis

The colour of chilli analysis was determined using Hunter lab-MiniScan EZ to measure colour in terms of L^* , a^* and b^* . The L^* , a^* , b^* model is an international standard for colour measurement developed by the Commission International d'Eclairage (CIE) in 1976. The luminance or lightness component (L^*) is value, ranging from 0 to 100. The colour component green to red (a^*) and blue to yellow (b^*) is ranging from -120 to +120. Then, all components are calculated to a total colour difference (ΔE^*) in equation (55), which has recently been compiled by Parametthanuwat et al. [116] and Kiniman et al. [117].

$$\Delta E^* = \left[(\Delta L^*)^2 + (\Delta a^*)^2 + (\Delta b^*)^2 \right]^{0.5} \quad (55)$$

The sensory of chilli was tests. All samples of ovened chili were blend into chili powder using blender Vitamix Company: USA as a model Vitamix 5200 at 5,000 rpm for 1 min. Then, the consumer acceptability of size, odor, and colour of the powder chili was evaluated by a 5-score hedonic scale sensory test [119, 120]. The categories for consumer acceptability were 1 = low, 2 = slightly low, 3 = intermediate, 4 = slightly high, and 5 = high. The number of consumers was 20. The experiment was performed in 3 replicates.

The shear force of chilli analysis was evaluated using Texture Analyzer (TA.XT plus, Stable Micro Systems, UK) to imitate the cutting and blending of chilli powder in manufacture industry. The 1 N load cell was equipped with blade set and the speed of the knife was set at 3 mm/s. The chilli sample size was a cylinder length of 30 mm and a diameter of 10 mm. Statistical analyses were based on Duncan's multiple range tests for means separation. Significant differences among the data were observed when $\alpha < 0.05$ [116, 117, 120].

4. Result and discussion

3.1. The temperature distribution

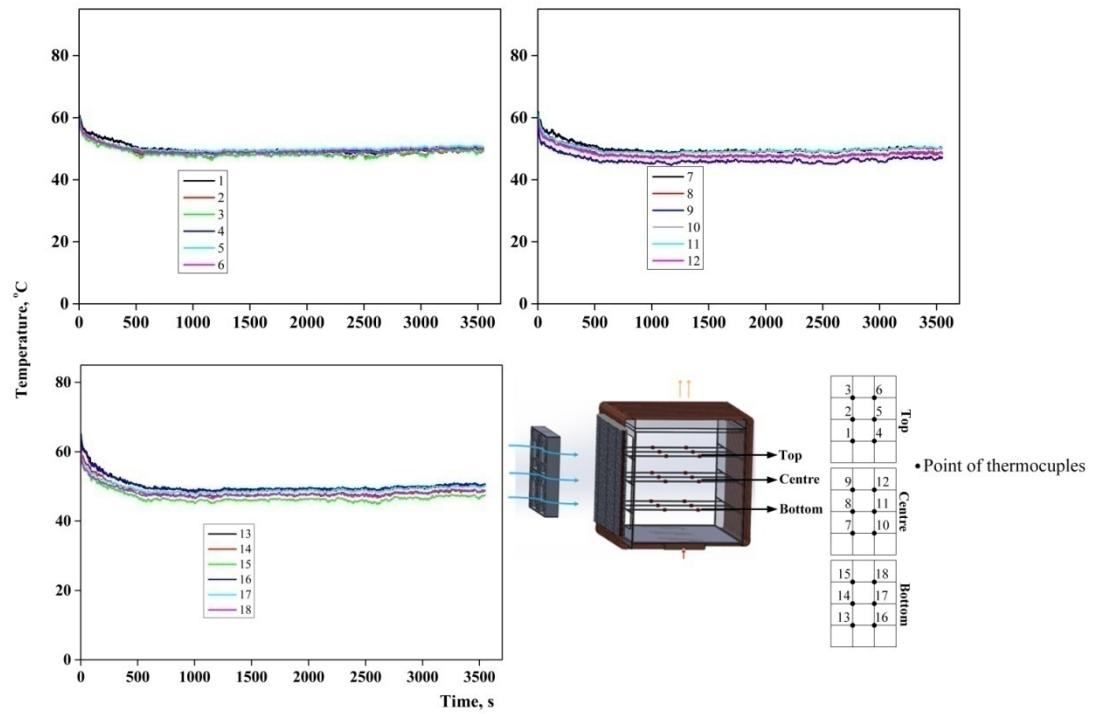


Figure 42 The distribution temperature inside the OALT/CV - Without workingfluids

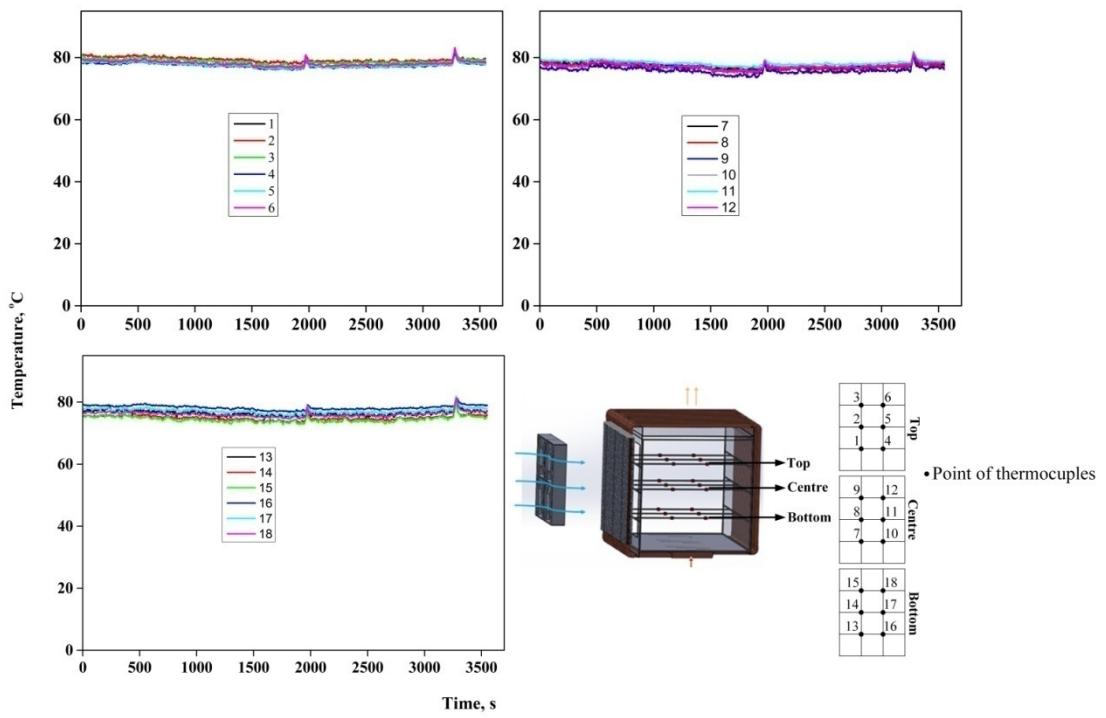


Figure 43 The distribution temperature inside the OALT/CV - Deionized water

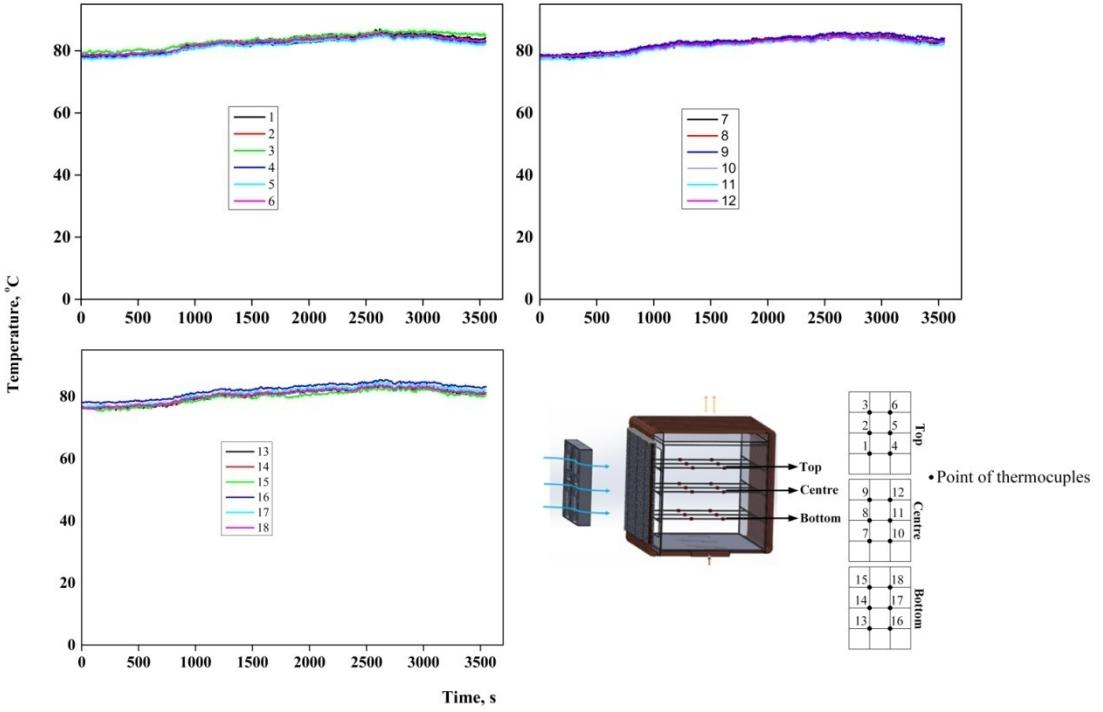


Figure 44 The distribution temperature inside the OALT/CV - 0.5 wt% silver nanoparticles (NP)

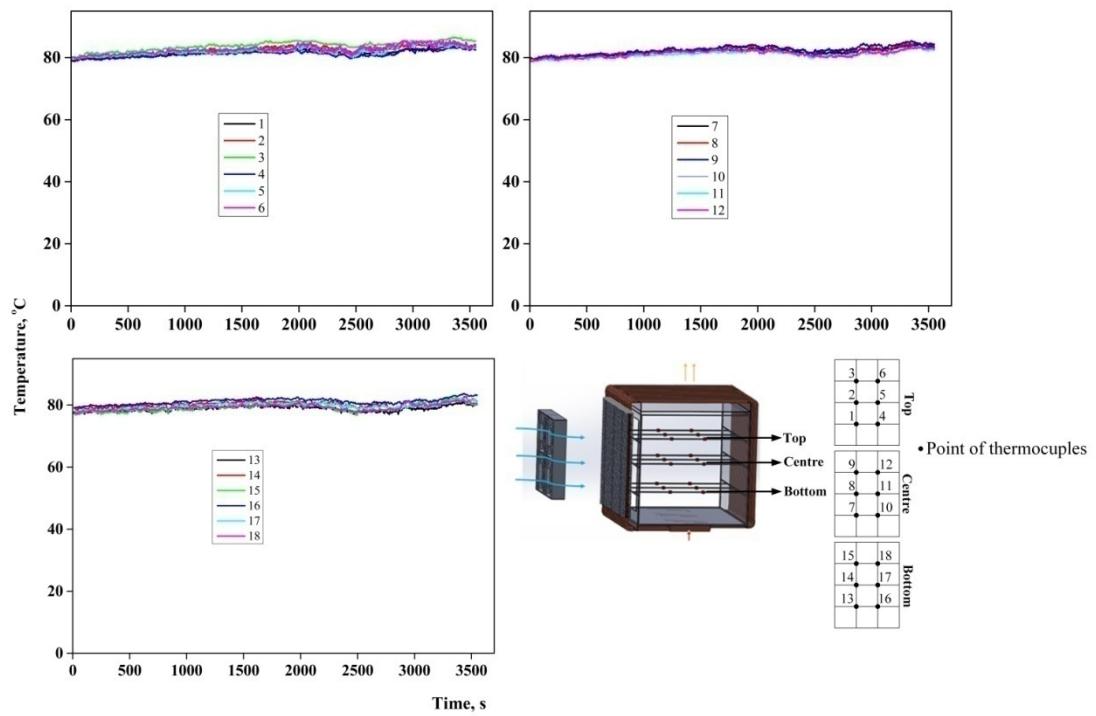


Figure 45 The distribution temperature inside the OALT/CV - NP containing 1.0 wt% of OA

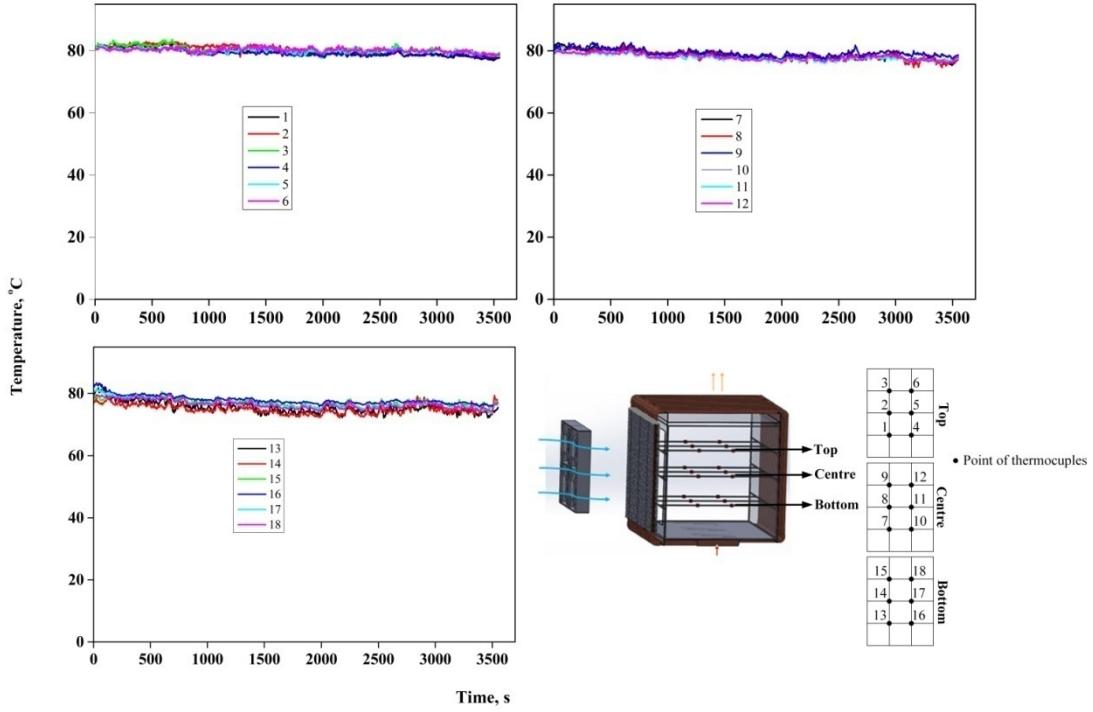


Figure 46 The distribution temperature inside the OALT/CV - NP containing 1.0 wt% of OAK+

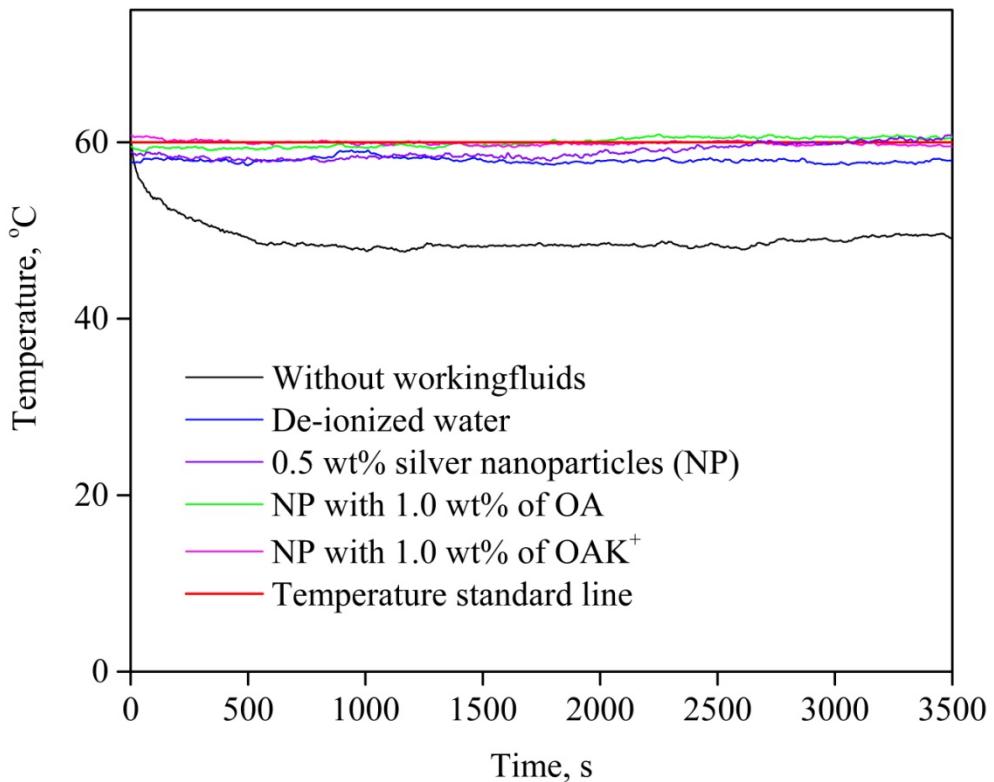


Figure 47 The distribution temperature inside the OALT/CV - Compared temerature contour

Table 10 Standard deviation of temperature inside of OALT/CV

Type of OALT/CV by workingfluids	Standard deviation		
	Top	Centre	Bottom
Without working fluids	±0.81	±0.67	±0.82
Deionized water	±0.50	±0.61	±0.72
0.5 wt% silver nanoparticles (NP)	±0.30	±0.30	±0.34
NP containing 1.0 wt% of OA	±0.20	±0.47	±0.33
NP containing 1.0 wt% of OAK+	±0.25	±0.33	±0.42

Figure 42, Figure 43, Figure 44, Figure 45 and Figure 46 of the OALT/CV shown temperature distribution at inside temperature controlled at 80°C with air velosity of 1 m/s. On the other hand, Figure 42 as a without working fluids was generated close inside average temperature at

60°C with air velocity of 1 m/s which was maximum and properly for OALT/CV- without working fluids. However, Figure 47 shown compared temperature distribution with operating temperatures at 60°C with line standard from OALT/CV- without working fluids. It was found that the temperature distribution around inside of the OALT/CV filled NP containing OAK⁺ was more regular than the other working fluids. The maximum temperature difference of OALT/CV was according to Table 10 in each type of OALT/CV by working fluids. Consequently, the OALT/CV filled NP containing OAK⁺ helped temperature distribution and gave higher quality performance than the OALT/CV without a working fluids.

3.2. The Quality of chilli

A chilli quantities study can be explained as follows into 3 groups including manufacturing standards: Colour quantities, Shear force and Sensory test. All groups was controlled by moisture lower than 13% wet basic. The quality of chilli measurement diagram shown in Figure 48.

The colour measurement of chilli is shown in Figure 49. The total colour difference (ΔE^*) will be according to the manufacturing standard. Thus, the statistical analysis points out that ΔE^* of chilli oven were significantly different ($p \leq 0.05$). Furthermore, the colour measurement of fresh chili, dried chilli from manufacturing and OALT/CV is shown in Table 11. Thus, each oven type have produced ΔE^* chilli similar to the manufacturing. Comparing both through statistical analysis points out that the colour measurement of chilli drying prepared by all conditions are significantly different ($p \leq 0.05$) which is according with Kiniman et al. [117]. The quality of colour of the chilli form OALT/CV exceeded the manufacturing standard. Definitely, the OALT/CV is confirmed as having good performance with short time.

Table 11 Total colour difference of chili

OALT/CV type	Colour quantities			
	L*	a*	b*	ΔE
French chili (Oven total time = N/A)	34.02 ^{ab} ±3.824	38.53 ^{cd} ±5.61	24.33 ^g ±5.23	-
Manufacturing (Oven total time = N/A)	26.86 ^b ±0.131	23.59 ^d ±0.541	13.03 ^{ab} ±0.39	28.88 ^{ag} ±0.68
Without working fluid (Oven total time = 21 hr)	34.96 ^{ca} ±2.47	25.87 ^{de} ±1.68	15.91 ^{cb} ±1.40	26.47 ^{fg} ±1.68
Deionized water (Oven total time = 17 hr)	30.58 ^a ±2.59	26.92 ^{eg} ±3.67	19.54 ^{cg} ±4.58	29.78 ^{fb} ±5.02
0.5 wt% silver nanoparticles (NP) (Oven total time = 15.5 hr)	31.86 ^{abc} ±1.44	22.54 ^{cde} ±1.13	15.42 ^c ±1.05	25.93 ^{be} ±2.32
NP containing 1.0 wt% of OA (Oven total time = 14 hr)	31.53 ^a ±1.06	24.23 ^{fg} ±4.49	17.06 ^{ag} ±0.88	24.35 ^{ce} ±1.56
NP containing 1.0 wt% of OAK ⁺ (Oven total time = 11)	32.93 ^{ab} ±2.42	22.65 ^f ±4.49	14.4 ^c ±3.59	27.92 ^{fg} ±4.70

Difference superscript (a,b,c,d,e,f and g) in the same column means that the values signification different ($p \leq 0.05$)

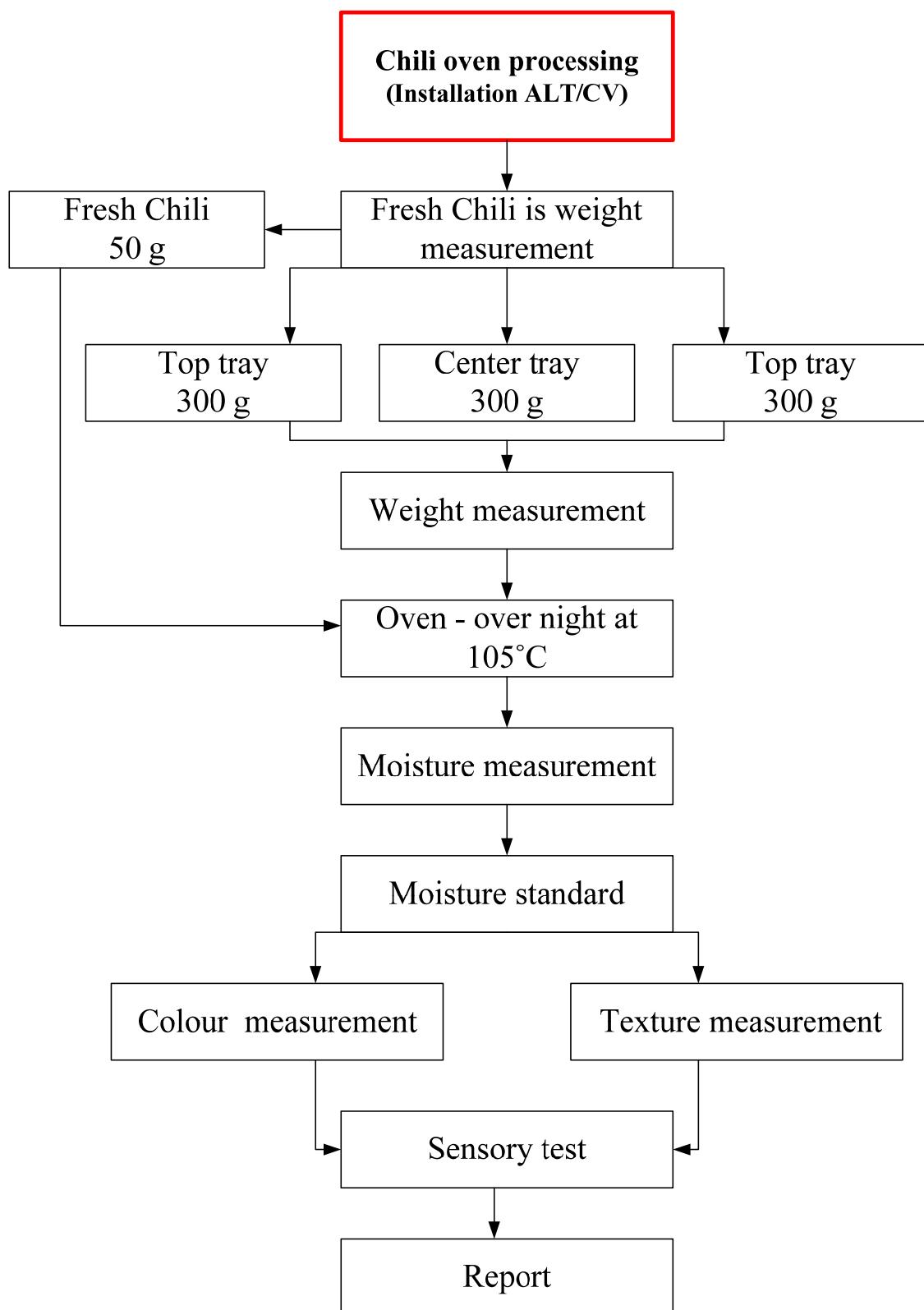


Figure 48 The quality of chilli measurement

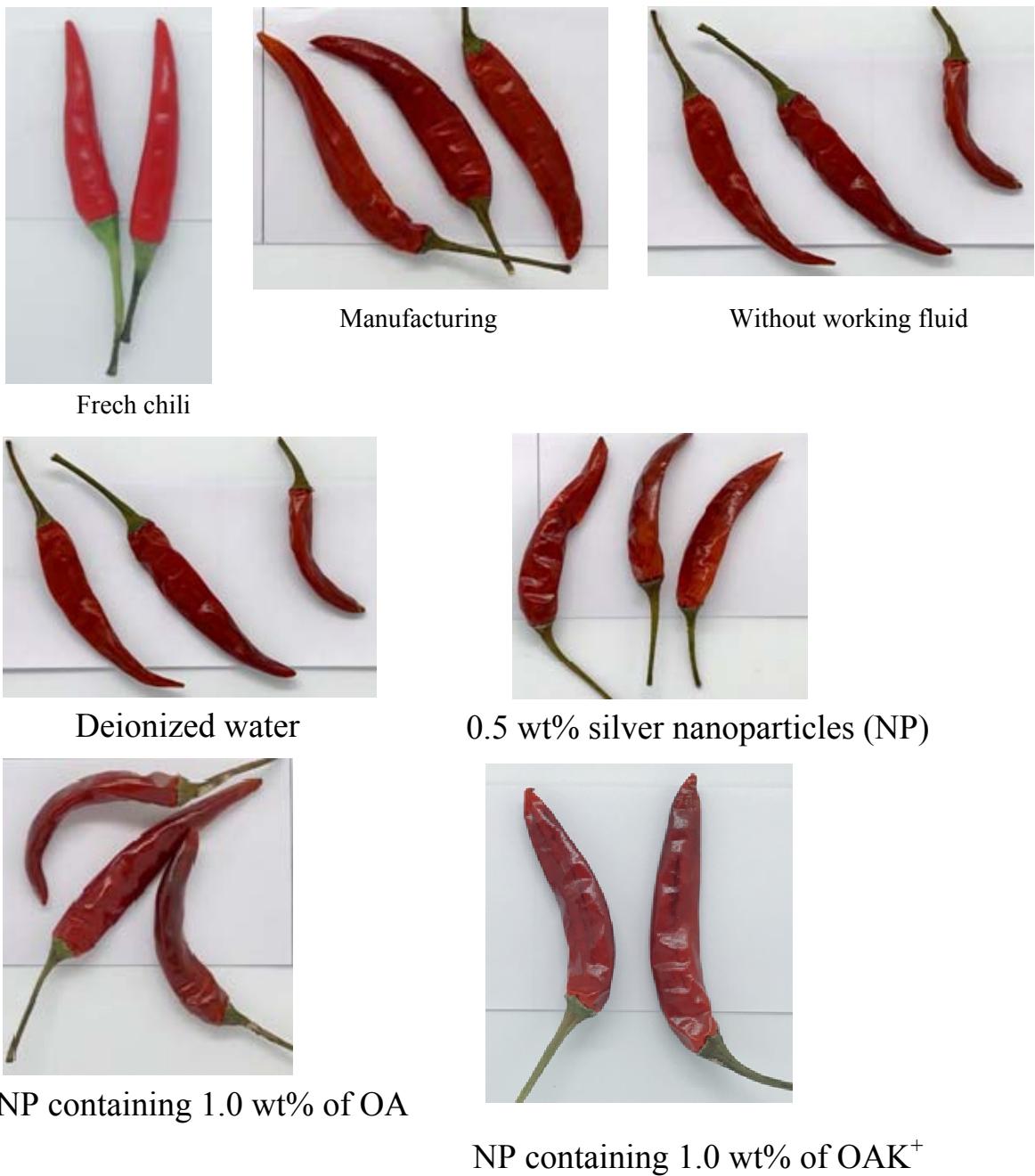


Figure 49 The colour chilli of each type oven

The result of shear force is shown in Table 12 and Figure 50. The values showed that shear force of OALT/CV is close the manufacturing. Moreover, shear force of OALT/CV did not significant different when comparing with that of manufacturing standard. This result revealed that blending of O/LTCV was easy and consumed less energy. The quality of chilli powder of the chilli form O/LTCV exceeded the manufacturing standard.

Table 12 Shear force (N) of dried chili

Type of OALT/CV by workingfluids	Shear force (N)
Manufacturing	25.62 ^a ± 1.11
Without working fluids	27.28 ^a ± 2.36
Deionized water	26.48 ^a ± 2.31
0.5 wt% silver nanoparticles (NP)	25.32 ^a ± 3.25
NP containing 1.0 wt% of OA	26.72 ^a ± 2.89
NP containing 1.0 wt% of OAK ⁺	25.78 ^a ± 3.15

Table 13 The result of Sensory score

Type of OALT/CV by workingfluids	Sensory score			
	Colour	Texture	Spicy and Smell	Odor
Manufacturing	4.12 ^a ± 0.65	4.20 ^a ± 0.72	N/A	N/A
Without working fluids	4.45 ^a ± 0.19	4.78 ^a ± 0.48	N/A	N/A
Deionized water	4.33 ^a ± 0.71	4.72 ^a ± 0.43	N/A	N/A
0.5 wt% silver nanoparticles (NP)	4.39 ^a ± 0.22	4.23 ^a ± 0.59	N/A	N/A
NP containing 1.0 wt% of OA	4.00 ^a ± 0.63	4.30 ^a ± 0.33	N/A	N/A
NP containing 1.0 wt% of OAK ⁺	4.68 ^a ± 0.12	4.16 ^a ± 0.43	N/A	N/A

Difference superscript (a and b) in the same column means that the values signification different ($p \leq 0.05$)



Manufacturing



Without working fluid



Deionized water



0.5 wt% silver nanoparticles (NP)



NP containing 1.0 wt% of OA



NP containing 1.0 wt% of OAK⁺

Figure 50 Shear force of dried chili

Sensory test in Table 13 is normally used to indicate consumer acceptability in food product. To evaluate and compare consumer acceptability of ovened chili which were produce from different oven. Sensory test was then performed. The result presented that the sensory which were produce from different ovened chili were in the range of 4.00 - 4.46. The values of colour, texture, spicy and smell which was product from OALT/CV ovened chili and manufacturing were significant different. The consumer agreed with the results of shear force, L^* , a^* , b^* , ΔE^* , smell and tester to check the level of spicy. This result indicated that OALT/CV oven can produce high quality of ovened chili. Consumer accepted OALT/CV oven chili similar with manufacturing.

3.3. The OALT/CV effectiveness

Table 14 The OALT/CV effectiveness

Type of OALT/CV by workingfluids	OALTCV effectiveness (%)		
	60	70	80
Without working fluids	-	-	-
Deionized water	18.92	25.23	35.33
0.5 wt% silver nanoparticles (NP)	28.36	30.45	444.58
NP containing 1.0 wt% of OA	37.45	46.26	52.36
NP containing 1.0 wt% of OAK ⁺	48.65	55.23	65.46

Table 14 shows the effectiveness of the OALT/CV with calculated form air velocity and LPG. It was found that the effectiveness dependent on air velocity and LPG. The maximum effectiveness of the OALT/CV was 65.46% with NP containing 1.0 wt% of OAK⁺. Consumer accepted OALT/CV dependent on condition in oven application which was selected.

5. Conclusions

The OALT/CV gave a total colour difference similar with the chilli from the manufacturing. All chilli at each tray level in the oven from OALT/CV had a total colour, texture, spicy and smell difference with the statistical analysis pointing out significant difference ($p \leq 0.05$). However, the quality of chilli ovened will exceed the manufacturing standard. Moreover, the ALT/CV improvement in the oven was more economical in the energy consumption cost which is increasing effectiveness. In conclusion, the design and improvement of the OALT/CV led to performance in oven process. The OALT/CV has maintained a quality exceeding the manufacturing while saving in energy consumption cost by a short time.

CHAPTER VI CONCLUSION

The objective of this study is based on the heat transfer rate of ALT/CV applying silver particles which containing oleic surfactant (OA) and potassium oleate surfactant (OAK^+) as working fluid. By solving the problem of traditional TPCT with advanced loop and check value will be led the heat transfer rate enhancement.

Finally, in conclusion, the thermal properties of NF containing OAK^+ was superior in thermal behaviour, over water studied in all experimental conditions. The presence of OAK^+ had clearly contributed to the rise in the heat transfer rate. By improving the properties of the working fluid with the OAK^+ this led to thermal properties enhancement, thus giving better properties than the OA. The OAK^+ showed compatibility with silver nanoparticle. The OAK^+ was Newtonian fluid. The optimum concentration for the addition of OAK^+ in the working fluid was 1 wt%. Expectedly, the heat transfer rate of ALT/CV was superior in heat transfer rate over of all experimental conditions studied. The design and improvement of the OALT/CV led to performance in oven process. The OALT/CV has maintained a quality exceeding the manufacturing while saving in energy consumption cost by a short time.

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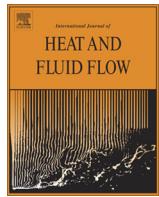
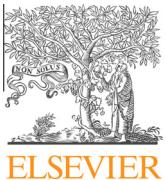
NOMENCLATURE

A_{eo}	outside wall area of evaporator section	m^2
A_{co}	outside wall area of condenser section	m^2
A	Surface area	m^2
C_p	Specific heat capacity constant	$\text{kJ/kg}^{\circ}\text{C}$
C_{pl}	Specific heat capacity of the working fluid at liquid phase	$\text{kJ/kg}^{\circ}\text{C}$
D_o	outside diameter of the pipe	m
D_i	inside diameter of the pipe	m
g	Gravity	m/s^2
h	Heat transfer coefficient	W/m^2
h_{eo}	Heat transfer coefficient at the evaporator section	$\text{W/}^{\circ}\text{C}\cdot\text{m}^2$
h_{co}	Heat transfer coefficient at the condenser section	$\text{W/}^{\circ}\text{C}\cdot\text{m}^2$
k_x	thermal conductivity of the material	$\text{W/}^{\circ}\text{C}\cdot\text{m}$
k_l	thermal conductivity of the working fluid at liquid phase	$\text{W/}^{\circ}\text{C}\cdot\text{m}$
L	Latent heat of working fluid	kJ/kg
L_e	Evaporator length	m
L_c	Condensor length	m
\dot{m}	Mass flow rate	kg/s
N	Number of fin	
NP	Silver nanoparticles 0.5 wt% contained in ethanol	
OA	Oleic acid surfactant	
P_v	Vapour pressure of the working fluid	Pa
P_a	Atmospheric pressure 101.3 kPa	
Q	Heat transfer rate	W
R_h	Radius of hydraulic	m
RT	relative thermal efficiency	
T_{out}	Outlet temperature of the condenser section	$^{\circ}\text{C}$
T_{inlet}	Inlet temperature of the condenser section	$^{\circ}\text{C}$
V_l	Volume of the working fluid	m^3
$W_{Perimeter}$	Wetted perimeter	m
X	Section width of RTPCT	m
Y	Section length of RTPCT	m
ρ_l	Density of the working fluid at liquid phase	Kg/m^3
ρ_v	Density of the working fluid at vapour phase	Kg/m^3
μ_l	Viscosity of the working fluid at liquid phase	N/m
η	Fin efficiency	
θ	Excess temperature	
ΔE^*	total colour difference	
l^*	Sample are lighter and darker standard	
a^*	Sample are redder and greener standard	
b^*	Sample are yellower and bluer standard	

APPENDIX

Journal publication

Parametthanuwat, T.; Bhuwakietkumjohn, N.; Rittidech, S.; Ding, Y. Experimental investigation on thermal properties of silver nanofluids. *International Journal of Heat and Fluid Flow*, 2015. 56: p. 80-90.



Experimental investigation on thermal properties of silver nanofluids



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ABSTRACT

This paper reports on an experimental investigation of the thermal properties behavior of 0.5 wt% silver nanoparticle-based nanofluids (NF) containing oleic acid (OA) and potassium oleate surfactant (OAK⁺) with concentrations of 0.5, 1, and 1.5 wt% respectively. The experiments were conducted from 20 °C to 80 °C. It was shown that the NF with 1 wt% OAK⁺ yielded the highest thermal behavior enhancement of about 28% at 80 °C compared to deionized water. The thermal performance had higher than the base fluid/nanofluids at approximately 80%. Moreover, the NF containing OAK⁺ showed higher thermal conductivity and dynamics of specific heat capacity than deionized water in all of the experimental conditions in this study. The rheological experiment showed that viscosity of NF was significantly dependant on temperature. As shear rate increased, the shear stress of the NF increased; however, the viscosity of the nanofluids decreased first and then stabilized. It was further found that NF containing OAK⁺ at a range of operating temperatures produced Newtonian behavior.

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1. Introduction

Cooling is one of the most important challenges facing numerous industrial sectors. Despite the considerable amount of research and development focusing on industrial heat transfer requirements, major improvements in cooling capability are still insufficient because conventional heat transfer fluids possess poor heat transfer properties. Nanofluids, which are engineered by suspending ultrafine metallic or non-metallic particles of nanometer dimensions in traditional cooling fluids, have shown great enhancement in thermal conductivity and convective heat transfer coefficient (Choi and Eastman, 1995; Khandekar et al., 2008; Parametthanuwat, 2012; Seeremya et al., 2014).

This section also contains the literature review for thermal properties. Many researchers have discussed the thermal properties points on which this study is based, together with the background of the research and explanations of the problems faced. This study highlights the theories and experiment for investigating the characteristics of thermal properties. Points of importance will be emphasized, with significance given to the properties of nanofluids and surfactants and their use in this experiment. Also included is the explanation of the characteristics of nanofluid behavior in silver nanofluids containing surfactant. Thus, the

researchers used different methods depending on the base fluids, nanofluid/nanoparticle type, etc. Recently, Thermal conductivity of 0.1–0.4% volume concentration silver (Ag) nanoparticles in water were investigated. The nanofluids were formulated using the ultrasonic vibration method for 3 h and thermal conductivity enhancement showed 10% at 0.4% of concentration (Kang et al., 2006). Using a different method, the synthesis of silver nanofluids was performed using high-pressure homogenization with a volume fraction 0.1–0.3% in water. The highest thermal conductivity of the nanofluids showed an 18% increase at the concentration of 0.3% (Oliveira et al., 2014). Moreover, regarding the difference in base fluid, Ag nanofluids in toluene have shown 9% thermal conductivity enhancement with a very low loading of 1.10–3 vol% (Daungthongsuk and Wongwises, 2007; Patel et al., 2003; Trisaksri and Wongwises, 2007; Wang and Mujumdar, 2007). Consequently, nanofluids show better cooling capacity with respect to water in conventional heat pipes since nanoparticles can flatten the temperature gradient of the fluids and reduce the boiling limit (Daungthongsuk and Wongwises, 2007; Khandekar et al., 2008; Trisaksri and Wongwises, 2007). In addition, the concentration of nanofluids may affect the enhancement of thermal conductivity. The studied silver nanofluids in ethylene glycol (EG) with 10,000 ppm concentrations showed 18% thermal conductivity enhancement (Sharma et al., 2011). Then, the investigated carbon black (CB) in deionized water with sodium dodecylsulfate (SDS) as well as Ag nanoparticles in silicon oil with

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Nomenclature

NF	silver nanofluids
NP	silver nanoparticles
OAK ⁺	potassium oleate surfactant
OA	oleic acid
K and n	consistency index and low power index
k_{eff}	the effective thermal conductivity
k_p	thermal conductivity of particle, $W(m\ K)^{-1}$
k_l	Thermal conductivity of liquid, $W(m\ K)^{-1}$
V_p	volume of nanoparticles in fluid, m^3
V_l	volume of base fluid, m^3
$C_{p,nf}$	specific heat capacity of nanofluids, $J(kg\ ^\circ C)^{-1}$
$C_{p,n}$	specific heat capacity of nanoparticles, $J(kg\ ^\circ C)^{-1}$
$C_{p,bf}$	specific heat capacity of base fluid, $J(kg\ ^\circ C)^{-1}$
$C_{p,eff}$	the effective specific heat capacity, $J(kg\ ^\circ C)^{-1}$

x_i	the independent variable to be estimated
μ_i	the manufacturer reported precision of the measurement
Q	heat transfer rate, W
T_{out}	outlet temperature at condenser section, $^\circ C$
T_{in}	inlet temperature at condenser section, $^\circ C$
\dot{m}	mass flow rate, $kg\ s^{-1}$

Greek symbols

ρ_n	density of nanoparticles, $kg\ m^{-3}$
ρ_{bf}	density of base fluid, $kg\ m^{-3}$
η	apparent viscosity
$\dot{\gamma}$	shear rate
ϕ	volume fraction of suspension

oleic acid (OA), and with the maximum enhancement of thermal conductivity compared to the base liquid, was 9% for the wt% of the carbon black (CB) nanofluids and the wt% of the Ag nanofluids respectively (Hwang et al., 2008). The .1 wt% copper (Cu) aqueous nanofluids with 0.14 wt% of sodium dodecylbenzene sulfonate (SDBS) as surfactant can generate maximum thermal conductivity enhancement up to 10.7% (Li et al., 2008).

The rheological behavior of nanofluids is essential in establishing adequate application and design of processing. The 8 wt% titania nanoparticles in the EG showed Newtonian behavior at a low shear rate, and the shear viscosity was strongly dependent on the temperature and concentration of the nanoparticles (Chen et al., 2007a). Then, the studied 1 vol% silver NP in ethanol with polyvinylpyrrolidone (PVP) was stabilized (Singh and Raykar, 2008). The rheological results suggest that the PVP helped to decrease the nanoparticle's size, resulting in low fluid viscosity and Newtonian fluid behavior but remarkably high thermal conductivity. Meanwhile, the 4.38 vol% silver nanofluids in the diethylene glycol (DEG) showed Newtonian behavior at high viscosity (Tajjid and Guenther, 2010). However, different literature data have shown that nanofluids have non-Newtonian behavior, particularly at a low shear rates. The most important influence could be the effective particle concentration, the range of shear rate, and the viscosity of the base liquid (Singh and Raykar, 2008). Then, it was found that the TiO₂ nanoparticle in the EG exhibited shear thinning behavior when the particle concentration was higher than ~2% (Chen et al., 2007a). The investigated shear thinning behavior was 3% γ -Al₂O₃ and 10% TiO₂ in water (Pak and Cho, 1998). Another main reason for the non-Newtonian behavior could be the aggregation of nanoparticles in the nanofluids. Lu reported that physical properties may change when the surfactant affects surface tension and viscosity. For instance, Al₂O₃ in water, at a 1:10 weight ratio with ammonium poly (PMAA-NH4), has demonstrated shear thinning behavior (a decrease in viscosity with an increased shear stress rate), which yields a good dispersion rate when using PMAA suspension up to 47.5 vol% (Lu and Kessler, 2006). Then it was found that the 4 vol% of γ -Al₂O₃, TiO₂, and CuO nanofluids with 0.5 wt% of carboxymethyl cellulose (CMC) in deionized water containing up to 4 vol% of particle concentration showed non-Newtonian behavior with shear thinning (Hojjat et al., 2011).

In this paper, 0.5 wt% silver nanoparticle-based aqueous nanofluids with oleic acid (OA) and potassium oleate surfactant (OAK⁺) as surfactant were prepared by sonicating in water bath with a cooling technique for a period time of 12 h. The effect of

the additive concentration on the thermal properties was studied experimentally (thermal conductivity, specific heat, density, viscosity, contact angle, and application of thermal enhancement), and the rheological behavior (the correlation between shear stress and shear rate) was investigated experimentally and theoretically. Moreover, the heat enhancement cooling of the fluid (HEC) was investigated experimentally and it was confirmed that nanofluids/nanofluids containing surfactant could be used in the application of heat transfer. The methods of the experiment are briefly explained in Section 2. Section 3 shows the experimental results and offers a discussion. The conclusions to the study are in Section 4.

2. Materials and methods

2.1. Nanofluids and thermal property study

Fig. 1 shows a schematic diagram of the preparation the nanofluids. Water-based silver nanofluids were formulated with dry silver nanoparticles (Sigma-Aldrich, USA), OA, and OAK⁺ (Sigma-Aldrich, USA) by using a two-step method (Hwang et al., 2008; Moghaddam et al., 2013). The 0.5, 1, and 1.5 wt% of OA and OAK⁺ were added to the 0.5 wt% silver nanofluids, which showed controlled and variable parameters as seen in Table 1. After sonicating for 12 h with a cooling technique, the particle size was measured using a nano-size particle analyzer (ZEN 3600 MALVERN, USA) in the range between 0.6 nm and 6.0 μm . The thermal properties of the nanofluids were measured using the hot-wire method (PSL Systemtechnik GmbH) from 20 $^\circ C$ to 80 $^\circ C$. The rheological characteristics of the NF were analyzed using a Rheo-microscope Physica MCR301 (Anton Paar GmbH). The measurements were based on the controlled shear stress model with the stress ranging from 0.05 to 5 Pa. The maximum uncertainty was found to be 1.7% (Chen et al., 2007b; Moghaddam et al., 2013; Parametthanuwat, 2012).

The rheological behavior of the NF containing OAK⁺ can be expressed with the power model in Eq. (1) with the viscosity as following the power law model indices less than $n \leq 1$.

$$\eta = K\dot{\gamma}^{n-1} \quad (1)$$

In Eq. (1), η is the apparent viscosity, $\dot{\gamma}$ is the shear rate, K is the consistency index, and n is the power law index. The power law index of the nanofluids decreases with increasing nanoparticle concentration, and increases with increasing temperature (Hojjat et al., 2011). Apparently, the viscosity of the NF decreases as the shear rate increases.

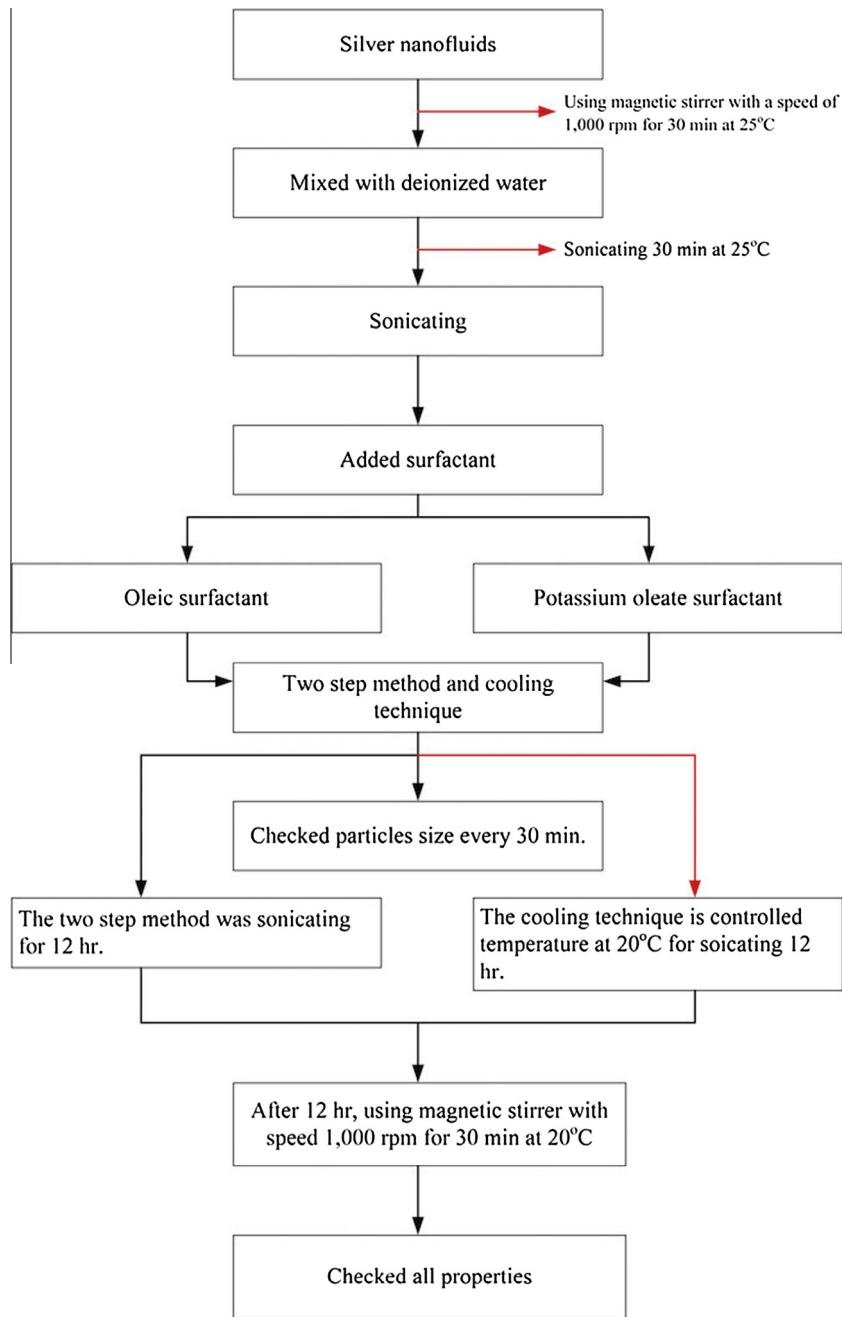


Fig. 1. The schematic diagram of preparation the nanofluids.

2.2. Thermal conductivity

The idea of thermal conductivity is non-existent in nanofluids theory scientists have used an existing model for estimated. The Maxwell model was developed to explain the heat transfer characteristics of larger particles in nanofluids research. This model has served as a foundation in the development and explanation of the much higher conductivity increase observed in nanofluids. The effective thermal conductivity (k_{eff}) can be defined by the following (Kleinsteuer and Feng, 2011; Kwak and Kim, 2005):

$$k_{eff} = \frac{k_p + 2k_l + 2(k_p - k_l)\phi}{k_p + 2k_l - (k_p - k_l)\phi} k_l \quad (2)$$

Thus, the ϕ can be defined by:

$$\phi = \frac{V_p}{V_p + V_l} \quad (3)$$

2.3. Contact angle instrument

In order to measure the contact angle of the sample fluids, the values are required to be at room temperature. The temperature was controlled with a precision of ± 1 °C. In this study, the drops of fluids were measured using a Contact Angle Meter Model: DM-CE1; Kyowa Interface Science. The accuracy was ± 0.5 ° (repeatability described in standard deviation). The following liquids were used in the experiment: a copper plate with a diameter of 60 mm and a thickness of 0.3 mm were used as a test surface. A droplet of nanofluid was generated at a very low rate (1 μ l/s) and detached from the syringe needle tip as soon as it touched the copper plate. Consecutive photographs were used to measure the contact angles. The spatial resolution was estimated to be about 50 μ m on the basis of the focused area and camera pixel size. A video was taken while the droplet was spreading over

Table 1
Controlled and variable parameters.

The controlled parameters	Silver nanofluid concentration of 0.5 wt% (NF) Operating temperature of 20–80 °C Shear rate ranges were 10 ⁰ s ⁻¹ to 10 ³ s ⁻¹
The variable parameters	Working fluid was: <ul style="list-style-type: none">• Deionized water• Deionized water containing surfactant• NF• NF containing surfactant Surfactant were: <ul style="list-style-type: none">• Concentration of Oleic acid was 0.5, 1, and 1.5 wt%• Concentration of Potassium oleate was 0.5, 1, and 1.5 wt%
Dependent variable	The dependent variable was: <ul style="list-style-type: none">• Thermal conductivity, specific heat, density, viscosity, contact angle and application of thermal enhancement• Rheological behavior

the copper plate from initial contact to equilibrium position. The temporal resolution was estimated based on the frame speed of the CCD camera at 30 fps. For each concentration, three experiments were performed and the average was ascertained. The measurement settings were then adjusted and the software was initialized (Lock, 1992; Ramesh and Prabhu, 2011).

2.4. Specific heat

The dynamic of specific heat was applied in the experiment. This was according to Rajabpour et al. (2013) regarding the application of the theory model to nanofluids.

$$C_{p,nf} = \phi C_{p,n} + (1 - \phi) C_{p,bf} \quad (4)$$

This second model has served as a foundation in the development and explanation of the much higher specific heat observed in nanofluids from nanoparticles. The effective specific heat ($C_{p,eff}$) can be defined as follows (O'Hanley et al., 2012):

$$C_{p,eff} = \frac{\phi(\rho C_p)_n + (1 - \phi)(\rho C_p)_{bf}}{\phi\rho_n + (1 - \phi)_{bf}} \quad (5)$$

The measurement uncertainty for the specific heat was calculated by propagating the precision uncertainties of all the individual measurements required to determine the specific heat in Eq. (5) and can be defined as follows (O'Hanley et al., 2012):

$$\mu_{total} = \sqrt{\sum_i^n \left(\frac{\partial C_{p,sample}}{\partial x_i} \mu_i \right)^2} \quad (6)$$

Eqs. (5) and (6) should be noted that for Newtonian nanofluids. For the non-Newtonian fluid, the variation of the rheology does not depend on direct models but on the volume fractions of the nanoparticles (Cabaleiro et al., 2013; Lu and Huang, 2013).

2.5. The heat transfer enhancement

Fig. 2 shows a schematic diagram of the experimental apparatus, which consists of the heat enhancement cooling of the fluid (HEC) and peripheral devices. The heat enhancement cylinder was made from Stan less steel (AISI 304) with a diameter and height of 1500 mm and 3000 mm. The HEC was the heat source from the Stan less heater (2000 W) with a diameter and height of 500 mm and 2000 mm. The heat was supplied by circulating the Stan less through to 20%, 40%, 60%, 80%, and 100% respectively of the heat source. The cooling and pre-cooling section was heat sink from a cold bath. The cooling fluids are shown in Table 2. Eighteen thermocouples were connected through a data logger (Yokogawa DX200 with ±0.1 °C accuracy, 20 channel input, and –200 °C to 1100 °C measurement temperature range). Type K thermocouples (OMEGA with ±0.1 °C accuracy) were attached to the inlet, the outlet, and the surface of the heating and cooling as the HEC. The inlet temperature of the cooling fluids was maintained at 20 °C and a floating Rota meter (PLATON PTF2 ASS-C with a volumetric flow rate of 0.2–1.5 l/min) was used to control the flow rate of the cooling fluid during the experiments. During the experiment, the volumetric flow rate was set at 0.25 l/min in order to calculate the heat transfer enhancement of the cooling fluid using the calorific method. The following Eqs. (7) and (8) were used for calculating one of the heat-transfer rates and for error analysis (Parametthanuwat et al., 2011).

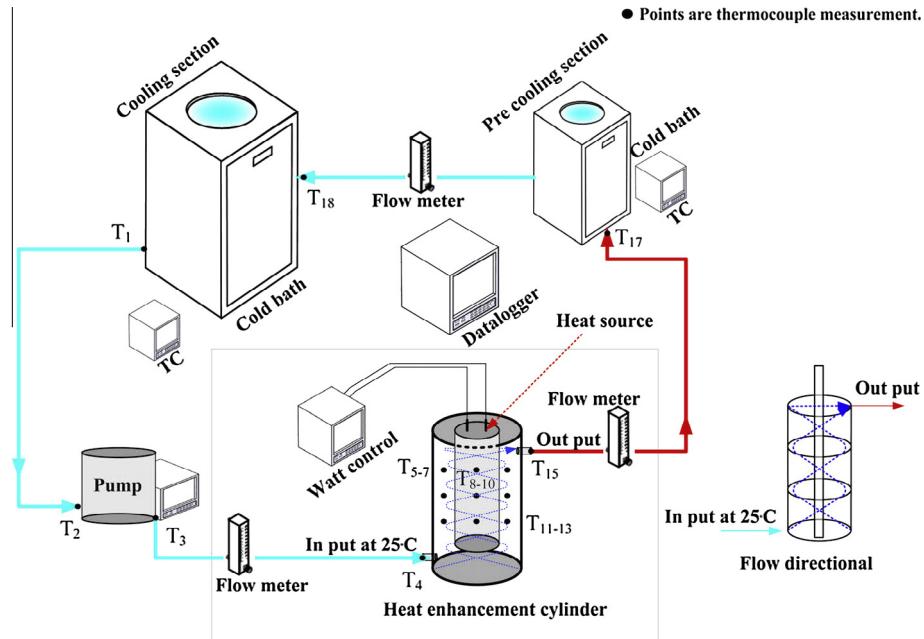


Fig. 2. Schematic diagrams of the HEC experimental apparatus.

Table 2
Controlled and variable parameters.

The controlled parameters	<ul style="list-style-type: none"> • Cylinder diameter and height of 1500 mm and 3000 mm. • Heater (2000 W) with diameter and height of 500 mm and 2000 mm. • The heat was supplied of 20%, 40%, 60%, 80% and 100% of heater.
The variable parameters	<p>The cooling fluids were:</p> <ul style="list-style-type: none"> • Deionized water • Deionized water containing surfactant • NF • NF containing surfactant <p>Surfactants were:</p> <ul style="list-style-type: none"> • Concentration of Oleic acid was 0.5, 1, and 1.5 wt% • Concentration of Potassium oleate was 0.5, 1, and 1.5 wt%
Dependent variable	Heat transfer enhancement

$$Q = \dot{m}C_p(T_{out} - T_{in}) \quad (7)$$

Thus:

$$Q = f(\dot{m}, T_{out}, T_{in}) \quad (8)$$

The error analysis of the heat transfer can be obtained from (Parametthanuwat et al., 2010):

$$Q_{Error} = \left[\left(\frac{\partial Q}{\partial \dot{m}} \times \dot{m} \right)^2 + \left(\frac{\partial Q}{\partial T_{out}} \times T_{out} \right)^2 + \left(\frac{\partial Q}{\partial T_{in}} \times T_{in} \right)^2 \right]^{0.5} \quad (9)$$

In order to experiment with a wide range of aspect ratios, the following parameters were corresponding set, as shown in Table 2, to formulate the heat transfer characteristics of the HEC (Parametthanuwat and Rittidech, 2013).

3. Results and discussion

3.1. The nanoparticle size

Fig. 3 shows the average particle size as a function of sonicating time. Zero point 5 wt% silver nanoparticles based nanofluids (NF) with surfactants as a stabilizer have an average particle size of ~100 and ~95 nm respectively. It can be seen that the average particle size decreases as sonicating time increases. Moreover, the red ellipse in Fig. 3a, NF+0.5OAK⁺ and NF+1OAK⁺ was seen to cause smaller particle size for a continuous period of 12 h when compared with Fig. 3b. This indicated that the two-step method and cooling technique did not break the agglomerate into primary particles (Bönnemann et al., 2005; Chen et al., 2007a). After sonicating for 12 h, the sample was put into the TEM (Oxford Instruments) to check for average the particle size, as shown in Fig. 4a, suggesting that the size distribution of particles NF+1OAK⁺ was between 5 and 25 nm. The TEM image also shows that the long chains of potassium oleate combined with outside nanoparticles and prevented them from aggregating together (Salehi et al., 2011; Vekas et al., 2006), which means that the viscosity and surface tension of the surfactant provided enough support to stabilize the dispersion of NP in deionized water (Parametthanuwat, 2012).

3.2. Rheological properties of nanofluids

The shear rate and shear stress had an effect on the rheological properties, such as viscosity. The rheological properties of the nanofluids containing surfactant are important to its thermo physical property. In this study, a surfactant was used to employ the NF's heat transfer rate. The OA and OAK⁺ are known for their ability to decrease viscosity and surface tension due to the organic and hydrocarbon interaction with oxygen which exists in deionized water (Huminic and Huminic, 2011; Salehi et al., 2011). However, the case was compared with the same group of surfactant (OA and OAK⁺) but with a difference in potassium salt (K⁺). The potassium salt was helpful in balancing the pH value of the

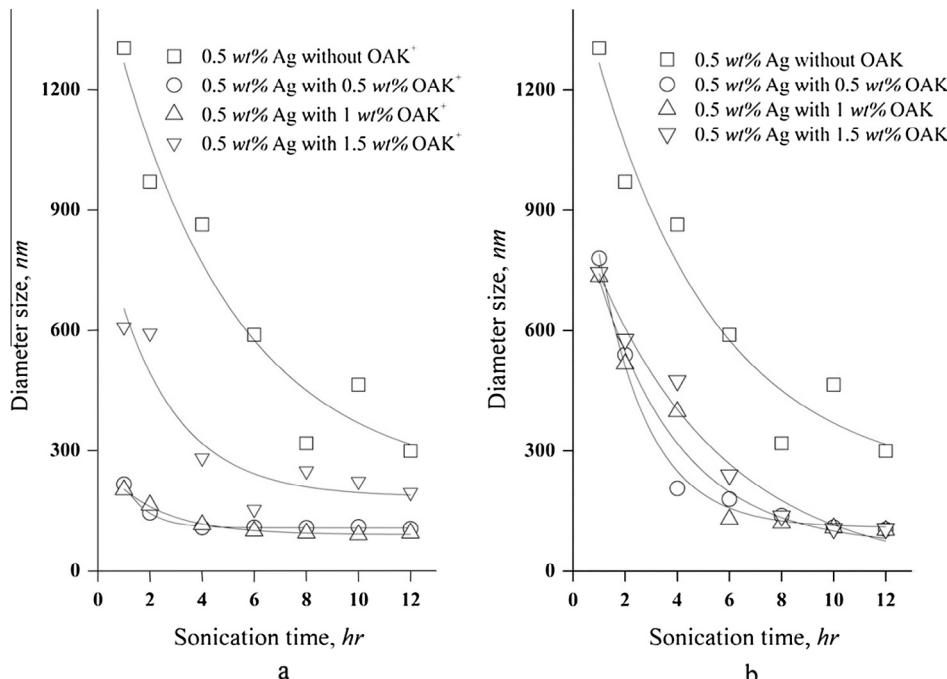


Fig. 3. Relationship between sonication times and particle size.

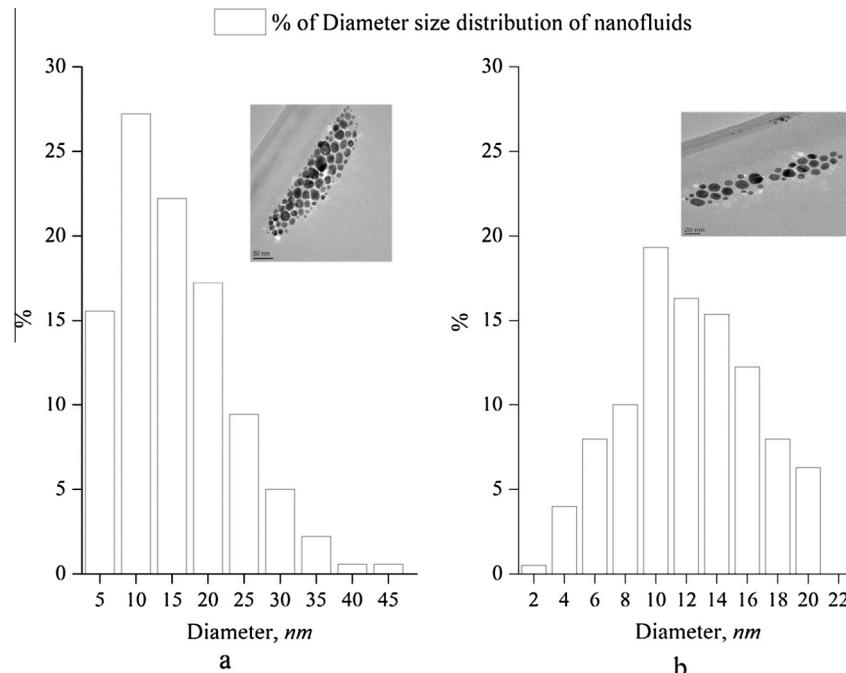


Fig. 4. Relationship between Diameters with percentage of dispersed size and TEM micrograph at silver nanofluids containing surfactant at concentration 1 wt%.

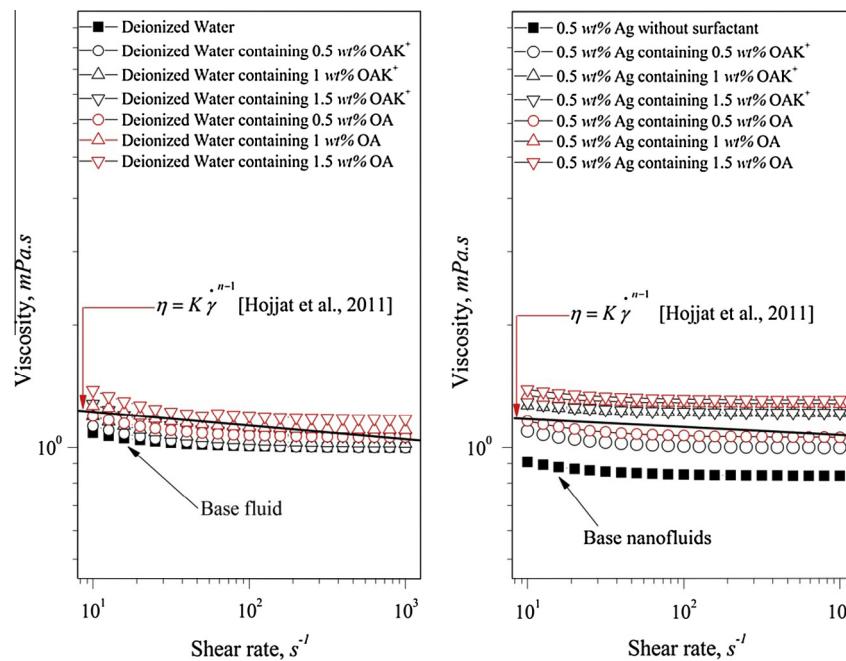


Fig. 5. Relationship between shear rates with viscosity of silver nanofluids at operating temperate 30 °C.

liquid and dissolving the solutions of fat oil catalysis interaction with hydrogen. Thus, the OAK^+ has the ability to span the NP's random motion throughout the deionized water (Chiu et al., 2008; Moghaddam et al., 2013; Molchanov et al., 2005; Mondragon et al., 2012; Paramethanuwat, 2012).

The rheology measurement results are shown in Fig. 6 and the shear rates with the viscosity of silver nanofluids were measured at 30 °C according to Paramethanuwat (2012). As shown in Fig. 5, the viscosity of all samples decreased in the first 10^1 s^{-1} to 10^3 s^{-1} intervals and the NF showed Newtonian behavior. This behavior could have been caused by the change of concentration

in the OA and OAK^+ , which was 0.5, 1 and 1.5 wt%. The most preferable OAK^+ concentration was 1 wt%, which was sufficient to distribute the NP with the lowest and most stable viscosity. The long chain nature of the OAK^+ molecular structure helped to decrease the NF's surface tension. It was concluded that there was an apparent change in the viscosity of the NF and deionized water; however, the NF's viscosity was still larger than that of the deionized water when the shear rate rose (Singh et al., 2013) according to Eq. (1) (Hojjat et al., 2011). Thus, the NP existing in the deionized water containing OAK^+ , which affected the flowing behavior of the nanofluids, was the main cause of the decrease in

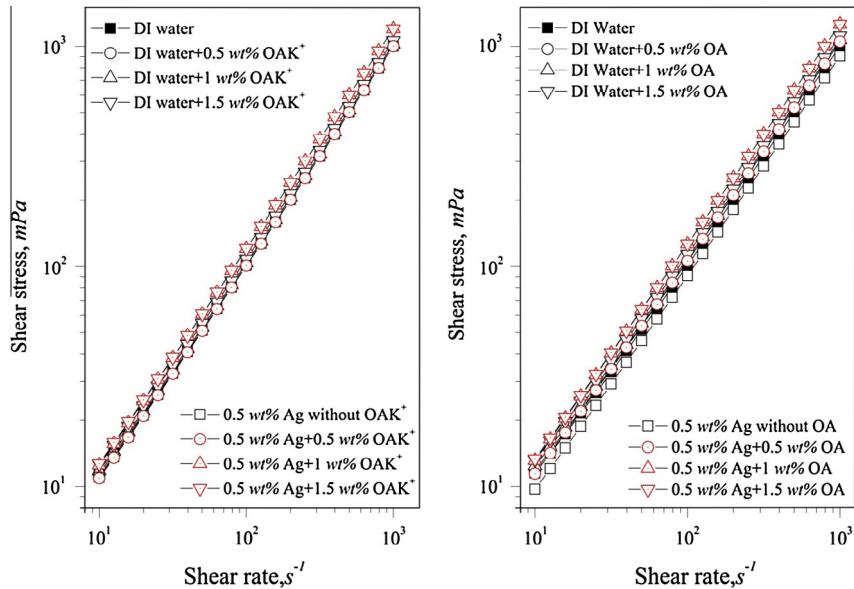


Fig. 6. Relationship between shear rates with shear stress of silver nanofluids at operating temperate 30 °C.

viscosity (Anoop et al., 2009; Lo et al., 2007). The potassium salt which produces an effect properly surfactant was helpful in balancing the physical properties of the base liquid. As can be seen in Fig. 6, along with the increasing shear rate, the nanofluids with OAK⁺ concentration smaller or larger than 1 wt% possessed larger shear stress. From the cross point onward, all of the results of the OAK⁺ were of almost the same value and the R^2 was close to 1. This was in line with the study of Hojjat et al. (2011), who stated that the transition metal in the same group with silver nanofluids containing OAK⁺ produced the Newtonian fluid (Parametthanuwat, 2012).

Fig. 7 shows the viscosity of NF+1OA compared with 0.5 wt% NF+1OAK⁺ as a function of shear rates. It was observed that for all operating temperatures and shear rate larger than 10^1 s^{-1} , the viscosity became stable and NF showed Newtonian behavior.

Thus, it was well established that for all operating temperatures, the system's rheological behavior exhibited a similar trend. Fig. 8 shows the relationship of the shear rates and shear stress in accordance with Fig. 7. It could be explained that the higher temperature increased the intermolecular distances, which decreased the interaction between the molecular structures of deionized water and OAK⁺, resulting in decreased viscosity and surface tension (Chen et al., 2007a). Obviously, the OAK⁺ could help decrease the physical properties more than the OA. The surfactant behaved like an interfacial shell between the nanoparticles and base fluids and modified the surface tension of the nanofluids. The surface tension decreased when the concentration of surfactant increased (Parametthanuwat, 2012; Tanvir and Qiao, 2012). The OAK⁺ exhibited good adsorption of the silver particles and the particles uniformly had a direct effect on the shear stress. The optimization of

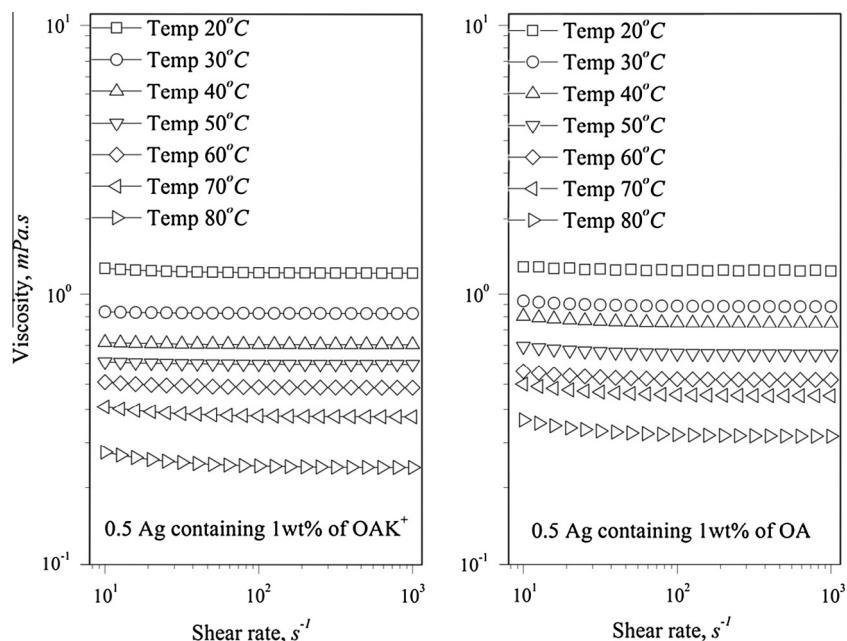


Fig. 7. Relationship between shear rates with viscosity.

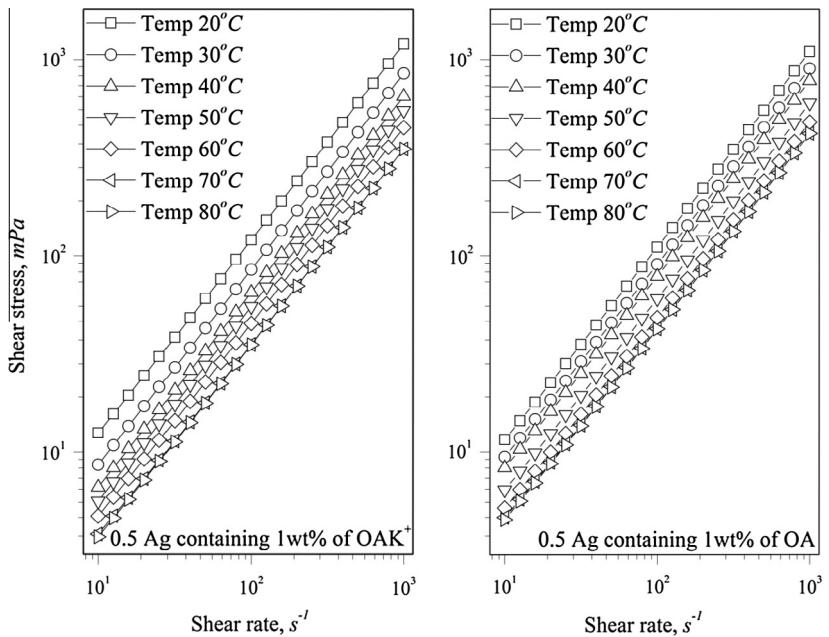


Fig. 8. Relationship between shear rates and shear stress.

the chain length in group OA organic compounds was C18, which was effective for particle dispersing stabilization. The optimized length also improved the colloidal stability and increased the non-precipitation period for the nanoparticles to be uniformly dispersed (Hwang et al., 2008; Li et al., 2010; Vekas et al., 2006). Moreover, OAK^+ achieved more stable suspension than the pure NP did in the deionized water. This might be related to the newtonian property of the nanofluids observed in this study when the rheological properties were observed at operating temperature (Moghaddam et al., 2013; Molchanov and Philippova, 2009; Parametthanuwat, 2012; Fohanno et al., 2011).

3.3. Thermal conductivity of nanofluids

The thermal conductivity of the silver nanofluids containing potassium oleate (NF+OAK^+) as a function of temperature is shown in Fig. 9. It can be seen that the thermal conductivity of the nanofluids depend on the linearity of the temperature, and the

enhancement of the thermal conductivity of NF+OAK^+ was different when the surfactant concentration was 0.5, 1, and 1.5 wt%. In all cases the NF+OAK^+ showed superior performance to that of the water. One wt% OAK^+ showed the lower and highest increase in the thermal conductivity of 15% at 20 °C and 28% at 80 °C throughout all samples, indicating that the thermal conductivity increases independently on surfactant concentration (Chen et al., 2007a). The nanoparticles dispersed in the liquid increased the surface area for the heat absorption. In the case of NF+OAK^+ , the OAK^+ decreased the surface tension of the nanoparticles, stabilized the nanoparticles by uniformly distributing them, and increased the interface area of the nanofluids with the deionized water (Vekas et al., 2006). The surface tension had a significant influence on the thermal process since the property departure and interfacial equilibrium depend on it (Parametthanuwat et al., 2010). The high OAK^+ concentration appeared to hinder the aggregation and entanglement of the NP, which was observed at the bottom of the liquid (Li et al., 2008; Sharma et al., 2011). According to our experimental results, 1 wt% OAK^+ was enough to homogeneously disperse the NP and produce efficient thermal transfer between the particles and deionized water, and consequently resulted in the highest thermal conductivity enhancement (Godson et al., 2010; Kang et al., 2009; Patel et al., 2003; Turkyilmazoglu, 2012; Veilleux and Coulombe, 2011).

However, the current experimental results contrast with those of Kang et al. (2006) and Oliveira et al. (2014), as shown in Fig. 10. In regards to this study, the results were achieved using the same silver nanoparticles however with a difference in concentration and surfactant. The result from Kang et al., demonstrated an increase of relative thermal conductivity. Thus, it is important to note the vast differences among different experimental conditions, especially in regard to the method of preparation and nanoparticle concentration. For example, Kang et al. (2006) showed a maximum k_{nf}/k_{bf} of ~1.11 at 0.4% volume concentration with nanoparticles diameter of 8–15 nm, while Oliveira et al. (2014) showed a maximum k_{nf}/k_{bf} of ~1.17 at 0.3% volume concentration with nanoparticles diameter of 10 and 80 nm. This contrasts significantly with the results achieved in this study of approximately 1.19 with nanoparticles of diameter 5–25 nm at NF+1OAK^+ . Of particular note, the research performed by Kang et al. (2006) and Oliveira

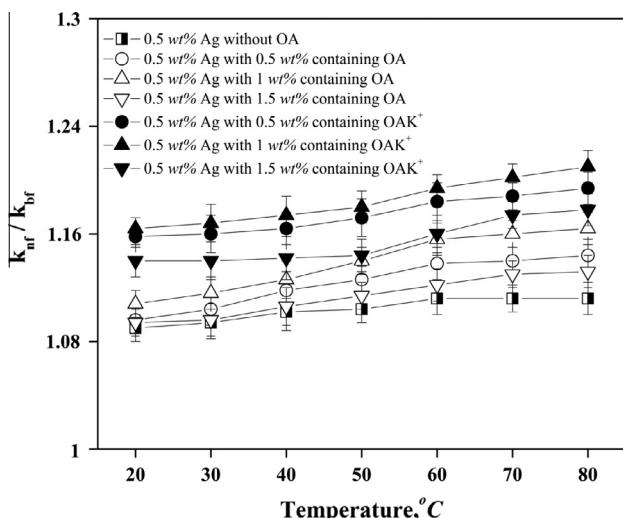


Fig. 9. Relationship between temperature with k_{nf}/k_{bf} .

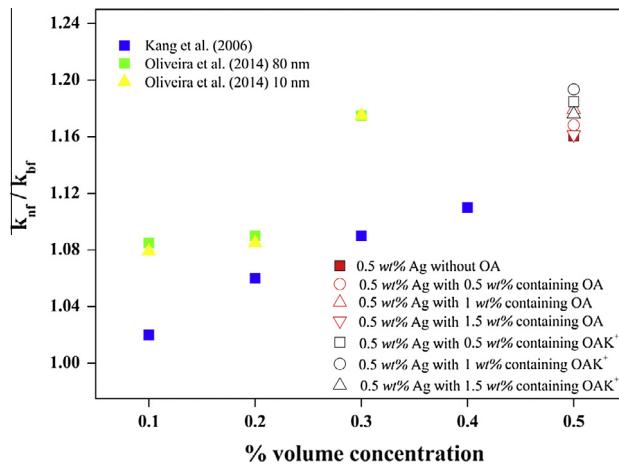


Fig. 10. Comparison different experimental result.

et al. (2014), did not specify the temperature under which the experiment was conducted. Furthermore, relative thermal conductivity was analyzed as a function of nanoparticle concentration and size. Thus, it can be determined that relative thermal conductivity is dependent upon the nanofluid's method of preparation, as well as the nanoparticle concentration and size.

3.4. Dynamic of specific heat capacity (DSC)

The heat energy absorbed ratio is a substance to increase in temperature. Table 3 shows the dynamic of specific heat capacity: nanofluids. The DSC was used to measure the deionized water deionized water containing surfactant (OA and OAK⁺) and the NF containing surfactant (OA and OAK⁺) at 1 wt% respectively, which was then compared with Eq. (6) according to Rajabpour et al. (2013) and O'Hanley et al. (2012). The results indicated a trend similar to that found in Rajabpour et al. (2013) and O'Hanley et al. (2012). It was found that, the good resulted appeared when using OAK⁺ as a surfactant. The silver nanofluids containing OAK⁺ surfactant was subjected to repulsion forces between the positively-charged hydroxyl groups (OA) of the functionalized nanoparticles and the potassium salt hydroxyl groups on the silver. However, the potassium salt of the acidic group functionalizing the nanoparticles became polarized in the water solution. On the other hand, when the positivity-charged potassium cation (K⁺) groups

were attached to the nanoparticles, the free ends (the carboxylic groups) became negatively charged. It is possible that the particle–fluid interactions and long-range electrostatic interactions between the nanoparticles may have affected the capillary properties of the nanofluids. Therefore, the negative forces between the solid and the nanofluids induced a specific heat capacity. Both parameters were dependent on particle size and surfactant concentration.

3.5. The contact angle and surface tension

In this study, the drop contact angle of working fluids was based on the room operating temperature. Furthermore, in order to investigate the effect of adding surfactant to the nanofluids, surface wettability, the static contact angles of the sessile droplets, and the surface tension of the pendant drop (Contact Angle Meter Model: DM-CE1; Kyowa Interface Science) were measured on flat copper plates at room temperature as shown in Fig. 11. This method was based on Khandekar et al. (2008) and Rahimi et al. (2010). When the surfactant was added it caused reduced adhesion between the working fluids dropping and the metal surface and made the total surface free energy. These are the manifestations of the interaction of the different molecular forces. The effects on the bulk thermo physical properties need to be addressed and surfactant dealt with nanofluids thus according to Radiom et al. (2013). It was further found that NF+1OAK⁺ decreased the water drop contact angle and surface tension to 38.23° and 28.69 mN/m, respectively (compared with pure water having 110° and 72.8 mN/m). Thus, the contact angle and surface tension depended on the operating temperature and time. They have an effect on the boiling phenomena in the heat transfer application. The transfer rate improves and decreases the surface tension with wet ability and contact angle (Kondiparty et al., 2011; Parametthanuwat, 2012).

3.6. Heat transfer enhancement

The experimental results clearly showed the effect of the percentage of heat supplied on the percentage of percentage of thermal heat enhancement, as shown in Fig. 12. When comparing the percentage of heat supplied using the different working fluids, it was seen that the NF+1OAK⁺ showed that the percentage of thermal enhancement was higher than with the other working fluids. Considering the case where the working fluid was NF+1OAK⁺ with a heat input at 100% of heat supplied, the thermal enhancement reached 24.75 ± 0.08%. The increase of maximum percentage of

Table 3

The dynamic of specific heat capacity: nanofluids.

Temperature (°C)	Deionized water containing 1 wt% OA		Deionized water containing 1 wt% OAK ⁺		0.5 Ag containing 1 wt% OA		0.5 Ag containing 1 wt% OAK ⁺	
	Theoretical C_p (kJ/kg K)	Measured C_p (kJ/kg K)	Theoretical C_p (kJ/kg K)	Measured C_p (kJ/kg K)	Theoretical C_p (kJ/kg K)	Measured C_p (kJ/kg K)	Theoretical C_p (kJ/kg K)	Measured C_p (kJ/kg K)
20	–	4.185 ± 0.070 4.182 ± 0.070	–	4.189 ± 0.056 4.182 ± 0.070	4.226	4.227 ± 0.002 4.225 ± 0.055	4.226	4.269 ± 0.02 4.225 ± 0.055
30	–	4.185 ± 0.068 4.183 ± 0.068	–	4.189 ± 0.036 4.183 ± 0.068	4.228	4.227 ± 0.045 4.225 ± 0.103	4.228	4.269 ± 0.120 4.225 ± 0.103
40	–	4.186 ± 0.065 4.182 ± 0.068	–	4.190 ± 0.032 4.182 ± 0.068	4.229	4.228 ± 0.020 4.225 ± 0.003	4.229	4.270 ± 0.025 4.225 ± 0.003
50	–	4.200 ± 0.056 4.182 ± 0.068	–	4.204 ± 0.048 4.182 ± 0.068	4.231	4.242 ± 0.025 4.226 ± 0.028	4.231	4.284 ± 0.008 4.226 ± 0.028
60	–	4.200 ± 0.060 4.183 ± 0.068	–	4.204 ± 0.060 4.183 ± 0.068	4.237	4.242 ± 0.012 4.227 ± 0.040	4.237	4.284 ± 0.023 4.227 ± 0.040
70	–	4.210 ± 0.040 4.187 ± 0.068	–	4.214 ± 0.089 4.187 ± 0.068	4.241	4.245 ± 0.121 4.229 ± 0.104	4.241	4.295 ± 0.016 4.229 ± 0.104
80	–	4.220 ± 0.037 4.189 ± 0.068	–	4.224 ± 0.010 4.189 ± 0.068	4.252	4.262 ± 0.121 4.236 ± 0.112	4.252	4.305 ± 0.039 4.236 ± 0.112

*The bold font second line shows base fluid (Deionized water) and base nanofluids (0.5 Ag without surfactant).

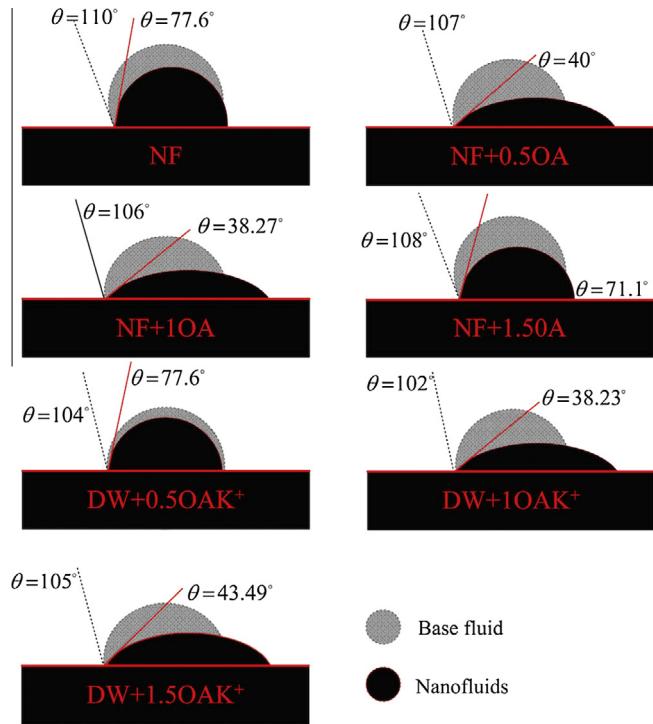


Fig. 11. Wettability at room operating temperature.

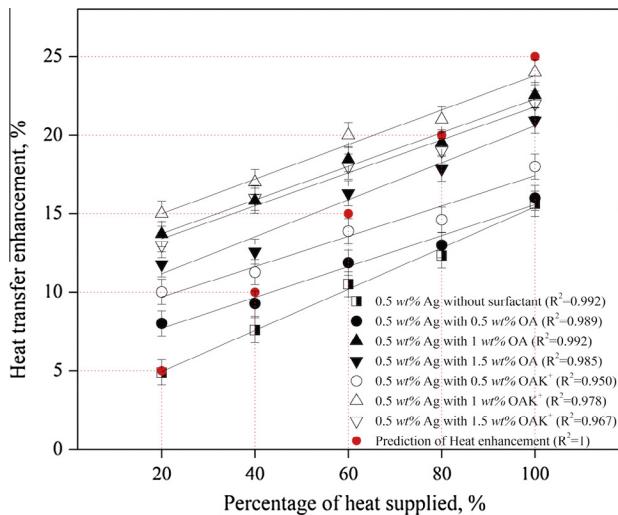


Fig. 12. Thermal enhancement of working fluids.

thermal enhancement with an increase in the percentage of heat supplied can be attributed to the increase in different temperatures (ΔT). In addition, the thermal motion of the nanoparticles enhanced the thermal properties of the nanofluids. The OA group helped with the homogeneous dispersion of the nanoparticles in the nanofluids. Moreover, the potassium cation (K^+) significantly contributed to accruing the thermal property increase, which had effects on the heat transfer rate mechanism more efficiently than the thermal diffusion in the fluid (Guo and Zetterlund, 2011; Hwang et al., 2008; Kwak and Kim, 2005; Wang et al., 2012).

4. Conclusion

Study on NF containing surfactant is leaded to nanoparticles size, the rheological properties of the nanofluids, the thermal conductivity of nanofluids, the dynamic of specific heat capacity (DSC),

wet ability (contact angle and surface tension) and heat enhancement. The important details are discussed below:

- The silver nanofluids containing OA and OAK^+ were conducted on thermal conductivity and rheological properties at various concentrations of OA and OAK^+ and operating temperatures. It was found that the NF containing 1 wt% of OAK^+ yielded better particle size ~ 95 nm.
- At a shear rate range of 10^1 s^{-1} to 10^3 s^{-1} , the samples showed Newtonian behavior, which showed Newtonian behavior, suggesting that the shear stress and viscosity decreased the high solid loading. For a given surfactant concentration, the consistency viscosity and shear stress of the base fluid and all of the nanofluids decreased with an increase in temperature, which confirmed that temperature had a strong effect on the shear stress and viscosity of the nanofluids. The rheological produced Newtonian.
- The NF containing 1 wt% of OAK^+ gave the highest thermal conductivity. It can be seen that the thermal conductivity enhancement was from 11% at 20 °C to 28% at 80 °C when compared with the base fluids. The DSC was increased with respect to operating temperature increase. Explicitly, the thermal conductivity of the NF containing the surfactant was in respect to the operating temperature, showing increments at all concentrations.
- The specific heat of NF containing OA and OAK^+ was superior in specific heat capacity, over water studied in all experimental conditions. The presence of surfactant had clearly contributed to the rise in specific heat capacity.
- It was concluded that the static contact angles of the OA group surfactant used, have better wettability characteristics, dependent on the surfactant concentration. Moreover, the NF containing 1 wt% of OAK^+ could be good at reducing the wettability and the OA group improved the colloidal stability which potassium cation (K^+) increased the non-precipitation period for nanoparticles to be uniformly dispersed in the base fluid. The nanofluids containing 1 wt% of OAK^+ produced a good contact angle of 38.23°.
- It could be concluded that the amount of the thermal enhancement of the nanofluids containing surfactant contributed to the greater rise of the thermal performance over the base fluid/nanofluids by approximately 80%.

Finally, in conclusion, the thermal properties of NF containing OA was superior in thermal behavior, over water studied in all experimental conditions. The presence of OA had clearly contributed to the rise in the heat transfer rate. By improving the properties of the working fluid with the OAK^+ this led to thermal properties enhancement, thus giving better properties than the OA. The OAK^+ showed compatibility with silver nanoparticle. The OAK^+ was Newtonian fluid. The optimum concentration for the addition of OA and OAK^+ in the working fluid was 1 wt%.

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Curriculum vitae



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I am Assistant Professor. Dr. Parametthanuwat. I have been working on several applications of heat pipes, nanofluid and renewable energy for many years. I also got several research funds from Thai government and Thai industrial companies. One of them is the Royal Golden Jubilee PhD Program (RGJ) in which Professor Dr Rittidech is supervisor; Assistant Professor Dr Pattiya and Professor Yulong Ding is an associate supervisor of an outstanding me. According to the RGJ program, I am study abroad to gain experiences from a high quality university in Institute of Particle Science & Engineering, University of Leeds. I am fully funded by the RGJ program and University of Leeds.

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Heat and mass transfer, Thermodynamic, Thermosyphon

Process engineering

Food plant design

Publication

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BOOK CHAPTER

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