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Heat transfer coefficient of nucleate boiling in low concentration level of single and hybrid Al_2O_3 - SiO_2 water-based nanofluids

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Abstract. Experiments were conducted to identify the Heat Transfer Coefficient (HTC) in saturated pool boiling of single and hybrid water-based nanofluids. In these experiments, Al_2O_3 and SiO_2 nanoparticles were selected and diluted into two separate single nanofluids, and they were mixed in a different ratio from 0:100, 25:75, 50:50, 75:25 and 100:0 percent to achieve a final total concentration of 0.001 vol. %. Successively, the mixtures were used to obtain the HTC values through experimental works. In the present work, it was found that in the lowest concentration (0.00025 vol.%) of Al_2O_3 nanofluid, the HTC enhanced considerably but deteriorated for SiO_2 nanofluid. Separately, as for the hybrid nanofluids, the HTCs were dramatically enhanced at the initial stage but slowly deteriorated once the time variation increased, especially in a higher ratio of SiO_2 nanofluid. The deposition of the nanoparticles onto the surface heater suggested being the main factor, where in the present case, the significant coexisting effect of the deposited hybrid nanoparticles (Al_2O_3 and SiO_2) on the heated surface to the changes of ΔT_w due to different nanoparticles properties.

1. Introduction

Nucleate boiling involves the high heat transfer rates and coefficients of convection which associated with small values of excess temperature. Therefore, it is desirable to operate many engineering devices such as chiller, heat exchange and many more which in the nucleate boiling regime. Currently, there are two significant heat transfer characteristics in the nucleate pool boiling that attracts most of the researchers which are the Heat Transfer Coefficient (HTC) and Critical Heat Flux (CHF). Most of the studies are focusing on the enhancement of the HTC and CHF which gives the significant benefits to the industries. One of the methods is using nanoparticles.

Various of studies about the effect of nanoparticles on the HTC and CHF were conducted after the discovery by You et al. [1] on the heat transfer enhancement that up until more than 100% for saturated pool boiling using Al_2O_3 . Several significant factors were identified to affect the heat transfer in the pool boiling especially related to the surface characteristics such as wettability [2–5], capillarity [6], surface roughness [7–9] and porosity [10–13]. All of these factors significantly altered by the nanofluids deposition proces on the heated surface during nucleate boiling. While the reported results



showed in nucleate pool boiling, the heat transfer demonstrated not only an enhancement but also can be deterioration, and their performance depended on the experimental condition [14–16].

Quan et al. [17] indicated that the pool boiling experiment of SiO₂ with surfactant resulted in the deterioration in HTC but drastically increase for CHF. It is differently finding by Kim et al. [15] where both for CHF and HTC for Al₂O₃ with distilled water was increased for a particular condition, and both also decrease for a particular condition. Separately, Watanabe et al. [18] proved that the HTC was enhanced, but CHF was deteriorated for nanofluids water based due to the nanoparticles deposited on the heater surface.

Based on the reported study by Sulaiman M. Z. et al. [19], the nanofluid was enhanced for Al₂O₃ nanofluids but deteriorated in a SiO₂ nanofluid. However, there is no experimental work conducted on the hybrid nanofluids to investigate the effect of nanofluids to the increment or reduction in HTC. An extension to that study, the present work will be emphasised on the performance of hybrid nanofluid (Al₂O₃ and SiO₂ nanofluids) at low concentrations respective to the heat transfer performance.

In this study, an experimental investigation was performed in nucleate pool boiling of water-based nanofluids. Thus, the paper aims to clarify the influence of concentration for single and hybrid nanofluids on the boiling heat transfer and evaluate the possible mechanism that responsible for the HTC enhancement or deterioration.

2. Experimental method

2.1 Preparations of nanofluids

In this study, Aeroxide Alu C (Al₂O₃ (Alumina) particle with a size of 13 nm) and Aerosil 90 (SiO₂ (Silica) particles with a size of 20 nm) were chosen as the enhancement ability shown by previous study [19–21]. These two nanoparticles were in white colours and manufactured by Aerosil Corporation. First, the nanoparticles were diluted in a series of low concentrations dilutions as depicted in Table 1. For hybrid nanofluids, the mixture was by mixing Al₂O₃ and SiO₂ using the ratio shown in Table 2. Then, the nanoparticles were weighted on a weight scale (Sartorius Practum213-1S) and then mixed with 75 ml of distilled water in a cup. The solution was stirred by hand for several seconds to make sure the nanoparticles adequately dispersed in the distilled water. Then the nanofluid was poured into the 100 ml test tube and then placed in the ultrasonic bath (CPX2800H, Branson). Each dilution was experiencing 1 hour of ultrasonic excitation to get the high stability dispersion of nanoparticles. Later, 75 ml nanofluids were heated up until it achieved its saturation temperature then were injected into the 1425 ml boiled distilled water which located in the test vessel [19].

Table 1: Volume % for single nanofluids

Nanoparticle	Volume %	Type
Al ₂ O ₃	0.00025 vol.%	A25
Al ₂ O ₃	0.0005 vol.%	A50
Al ₂ O ₃	0.001 vol.%	A100
SiO ₂	0.00025 vol.%	S25
SiO ₂	0.0005 vol.%	S50
SiO ₂	0.001 vol.%	S100

Table 2: Ratio of volume % for hybrid nanofluids

Ratio					Types
Nanoparticle	Volume %	Nanoparticle	Volume %	Total volume %	
Al ₂ O ₃	0.001	SiO ₂	0	0.001 vol.% : 0 vol.%	A100S0
Al ₂ O ₃	0.00075	SiO ₂	0.00025	0.00075 vol.% : 0.00025 vol.%	A75S25
Al ₂ O ₃	0.0005	SiO ₂	0.0005	0.0005 vol.% : 0.0005 vol.%	A50S50
Al ₂ O ₃	0.00025	SiO ₂	0.00075	0.00025 vol.% : 0.00075 vol.%	A25S75

Al_2O_3	0	SiO_2	0.001	0 vol.% : 0.001 vol.%	A0S100
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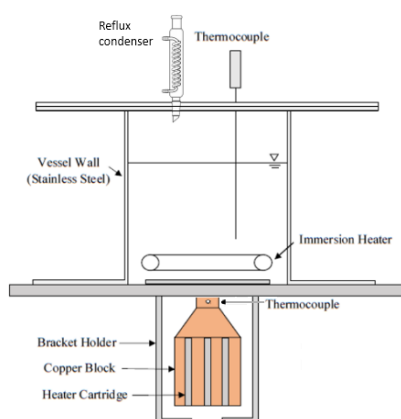
2.2 Experimental apparatus

In the present experiment, an experimental setup was fabricated nucleate boiling to investigate the boiling performance in nanofluids. The test sections consisted of a cylindrical vessel, a Perspex cover, a copper block attached with cartridge heaters, several copper block holders, several thermocouples and ceramic fiber insulation blanket. The detail schematic diagram and a photo of the experimental apparatus were shown in Figure 1 (a) and (b).

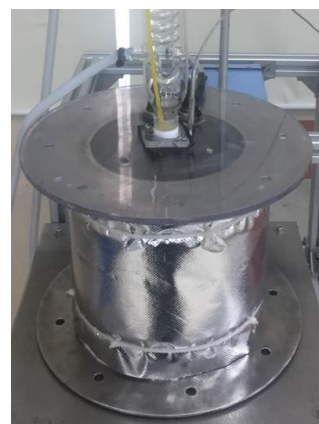
The main body, the vessel, was made from stainless steel with a thickness of 3 mm and the outer diameter of 145 mm of inner cylindrical diameter and the height of the vessel was 185 mm. Outside the cylindrical vessel, the outer surface was covered with thick ceramic fiber insulation blankets. At the bottom of the cylindrical vessel, a heating component was mounted concentrically. A modified one end of the copper block from 40 mm original diameter and shaped it to the cylindrical shape of 20 mm diameter. The copper block surface was used as the heater surface and was in direct contact with the fluids in the main vessel. This copper block attached with a circular stainless-steel plate as a flange with an outer diameter of 100 mm and holes through the middle 20 mm diameter hole. Both components assembled and cured with a reliable and high temperature resistant resin to keep the heater block and stainless-steel flange attached as well as eliminating the clearance gap.

Three type-K thermocouples were positioned along the central axis at this copper end to measure the temperature in the copper block which determines the heat flux, q_w and wall temperature, ΔT_w . The thermocouples were inserted into the drilled holes from top to bottom. While for the other end of the copper block, drilled four holes and inserted cartridge heater (900 W) into all the holes. The copper block was attached to 3 holders to prevent from a slip throughout the experiments. The copper heater and the holder were wrapped a with several layers of ceramic fiber insulation blankets with an anti-burning layer sheet covering outside. The cover was made from Polycarbonate sheet with 6 mm thickness and 250 mm diameter. Using the uncertainty analysis method described by Cooke et al. [22], the measurement uncertainties of q_w and ΔT_w were estimated within less than 84 kW/m² and 2.4 K, respectively.

Additional immersion heater with 1 kW was attached to the top cover and located concentrically at the centre of the lower part of the vessel. The immersion heater used to maintain the bulk liquid temperature in the vessel at the saturation condition. A Reflux condenser was equipped on the top cover to prevent the vapour from the test vessel and its cooled with tap water circulation. The pressure inside the vessel was assumed to equal to the atmospheric pressure as the top condenser was open to the atmosphere. Another type-K thermocouple was attached to the cover with its end enter into the vessel. The thermocouple's end was positioned close to the outer diameter of the modified copper surface. A 5 mm diameter hole was drilled onto the top cover, functioned to allow the injection of nanofluid into boiling water in the vessel by using a syringe, then the hole was closed with a rubber plug except during the nanofluid injection.



(a) Schematic diagram



(b) Photo

Figure 1: Experimental apparatus

2.3 Experimental procedure

The experiment began with preparation of the heated surface. The heated surface was polished by using metal polishing paste and Kim Wipes, and successively with the surface cleaning by using acetone. Then the vessel cylinder was mounted. Later, the first variable transformer that was attached to the cartridge heater was regulated to supply a 600 kW/m^2 of heat flux to the copper block. After that, 1425 ml of distilled water were poured into the vessel, and the vessel was closed using Polycarbonate sheet cover attached on the top. Next, the second variable transformer that attached to the immersion heater was regulated to the maximum capacity of 1 kW to boil and degas the distilled water. In order to keep the water level inside the vessel and eliminate the evaporated liquid to escape during the experiment, a Reflux condenser used to condense the water, and it was attached in the middle of the Polycarbonate cover. The degassing process was set to be 20 minutes for each experimental work, and the voltage output from the variable transformer for immersion heater was reduced to the appropriate level enough to keep the water at the saturated condition.

In this experimental stage, the copper block temperature and fluids temperature were measured and monitored by using a temperature module datalogger from National Instrument and DASY LAB software, respectively. The temperatures data were used to calculate the value of wall superheat, ΔT_w and the steady state condition was determined by observing the scattering of the ΔT_w value in the experimental run was about less than 0.3 degrees C for the approximate duration of 10 minutes. Finally, after a steady state of the copper block was confirmed, the experiment was ready for the next step.

Since the experimental procedures were repeated for each experimental run, the scattering of ΔT_w was non-negligible in a separate experimental run since the condition of the heated surface could not be the same. In the experimental setup, the mean ΔT_w was found to be in the range of $14 \pm 2 \text{ }^\circ\text{K}$ and at this state, the nanofluids addition was conducted. This step was necessary because it reduced the influence of the scattering of the initial wall superheat. Separately, the prepared nanofluids in a test tube were heated in another hot boiling water to ensure that the temperature of the nanofluid did not affect the saturated temperature of the distilled water in the vessel. Once the steady state condition was of the copper block was confirmed, the nanofluid was poured into the vessel by using the modified funnel syringe through a small hole located at the top cover of the vessel. After that, the hole was closed with a rubber plug, and the experiment was run for 1 hour, and temperatures were recorded for every second and the time variation of wall superheat, ΔT_w was analysed.

3. Result and discussion

3.1 Effect of concentration variations at relatively low concentration level to the HTC in single nanofluids

Figure 3.1 depicted the time variation of wall superheat ΔT_w for two types of single nanofluids ($\text{Al}_2\text{O}_3/\text{water}$ and $\text{SiO}_2/\text{water}$) in several levels of low concentrations between 0.00025 and 0.001 volume %. They were coded to be A25, A50, A100, S25, S50 and S100 as described in the previous section (see Table 1). Boiling time t_b was set to be the time duration after the injection of nanofluids into the vessel with a duration of 1 hour present experimental setup. As can be seen, the initial ΔT_w of distilled water achieved steady state from the beginning and did not change upon the addition of 75 ml distilled water from the test tube, and it is considered as a reference line. While one could noticeable seen that ΔT_w drastically change once the nanofluid was added as shown in Figure 3.1 (a) and (b).

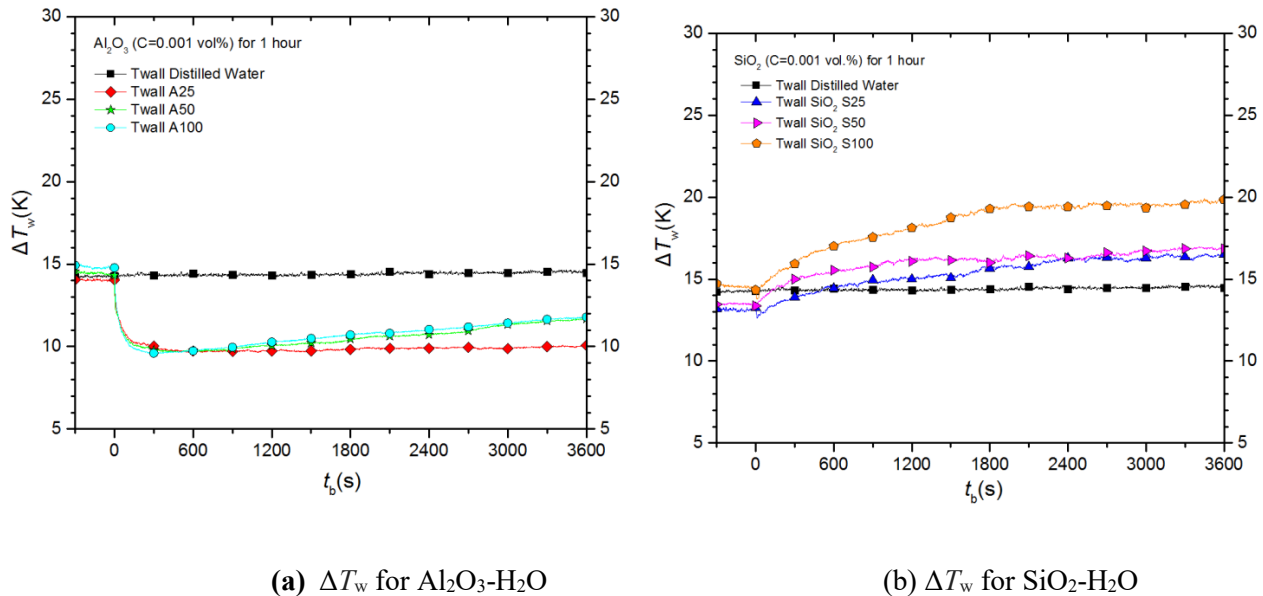
(a) ΔT_w for Al₂O₃-H₂O(b) ΔT_w for SiO₂-H₂OFigure 3.1: Time-variations of wall superheat after adding single nanofluids ($q_w = 600 \text{ kW/m}^2$)

Figure 3.1(a) demonstrated the time variation of wall superheat after the injection of Al₂O₃/water nanofluid. As can be seen, the ΔT_w was instant and sharply reduced after the nanofluid addition for all low level concentration of Al₂O₃ nanofluid. The results indicate for all those three levels of the low concentrations; each respective ΔT_w were kept lower than the distilled water after 1 hour. One could notice that ΔT_w for A50 and A100 show similar patterns where after sharp reduction of ΔT_w value at the earlier stage a gradually increased trend could be seen after 10 minutes of experiments and end up at the same ΔT_w after 1 hour. It could be seen that and they need longer than 1 hour to reach the steady state conditions. While for A25, the ΔT_w was kept low throughout 1 hr experiment duration and maintaining a steady state condition. The figure indicates that the smallest level of concentration (A25) could improve heat transfer dramatically. Although both A100 and A50 also showed similar enhancement at the initial stage, the ΔT_w was observed keep on increasing in the 1 hour experimental duration.

Figure 3.1 (b) showed the time variation of wall superheat ΔT_w for SiO₂/water nanofluid with three different low concentrations, coded as S25, S50 and S100. In contrast with ΔT_w of Al₂O₃/water nanofluid, the addition of SiO₂ seems to increase the value of ΔT_w . For a maximum concentration of SiO₂/water in the present experiment (S100), the value for ΔT_w was dropped drastically for a few seconds and increased monotonically after that respect to time. After 30 minutes of experiment duration, it achieved the steady state condition where the value of ΔT_w was kept maintained in nearly similar range as shown in Figure 3.1 (b). Similarly to the result for the lowest concentration of SiO₂/water (S25) in present experiment where in the first few seconds, the ΔT_w was decreased drastically but increased monotonically to achieve steady state condition. However, compared to the other two higher concentrations, the value of ΔT_w was the lowest along the 1 hour of the experiment. For S50 type, it is unnoticeable in Figure 3.1 (b) that the value of ΔT_w dropped for the first few second.

After the nanofluids injection, the graph line seems increased monotonically. All graph lines of three different concentration from SiO₂/water nanofluids shows the same parallel movement of the ΔT_w which increased monotonically. However, for S25 and S50 concentrations still did not achieve the steady state condition and keep on increasing slowly. Therefore, further study must be performed to identify the time needed for these two concentrations of SiO₂/water nanofluid to achieve steady state.

In the present experiments, it was noticeable that after the presence of Al₂O₃ and SiO₂ in the based liquids, the wall temperature, ΔT_w changed depends on the types of nanofluids and its concentrations, supported by the previous studies [23–27]. It can be seen that the value for ΔT_w of Al₂O₃/water nanofluid decreased as shown in Figure 3.1 (a), same as reported by Hassanpour et al. [28].

Therefore, the HTC was enhanced in that situation. In contrast with $\text{SiO}_2\text{-H}_2\text{O}$ nanofluids, the HTC was deteriorated because of the pattern of the graph which keeps on increased more than the reference line respect to time. Although the present experiments used a relatively low concentration of nanofluids; it did affect the ability of HTC in the nucleate pool boiling as proven in Figure 3.1.

3.2 Composition ratio effects to the HTC of hybrid ($\text{Al}_2\text{O}_3\text{-SiO}_2$)/water nanofluid at low concentration (0.001 vol. %)

Base on the unique result of contradicting HTCs in two types of single nanofluids as discussed in section 3.1, experimental work was extended by mixing two types of particles material called hybrid nanofluids. In the present work, Al_2O_3 and SiO_2 were diluted into a series of combination to obtain a single and hybrid water-based nanofluids in order to investigate their HTCs performance in depth. It should be noted that in some cases especially for both single nanofluids, a similar trend also obtained in the previous investigation by Sulaiman M. Z. et al. [19]. The present water-based nanofluids mixing was labelled as A100S0, A75S25, A50S50, A25S75 and A0S100 (see Table 2).

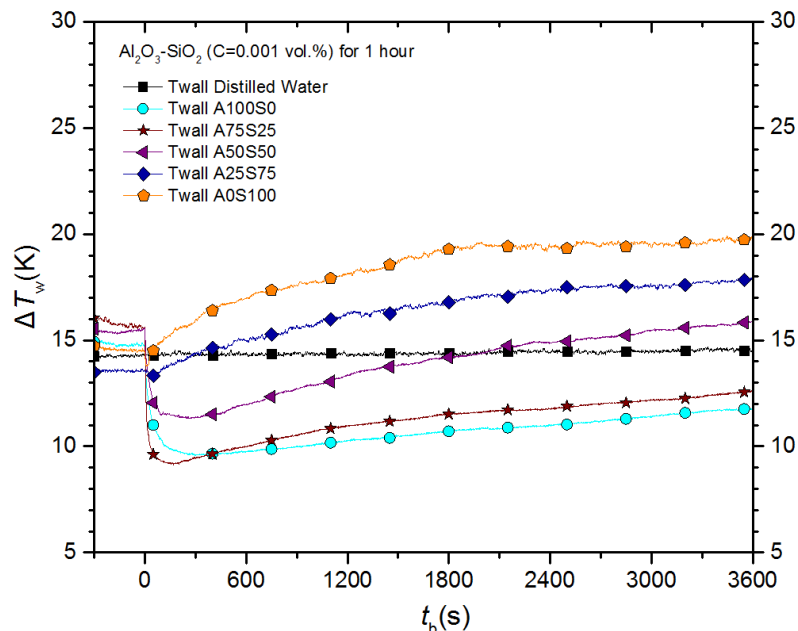


Figure 3.2: Time-variations of wall superheat after adding hybrid nanofluids ($q_w = 600 \text{ kW/m}^2$)

Figure 3.2 depicted the result of the wall super heat ΔT_w of two single nanofluids and their hybrid at a low-level concentration of 0.001 volume %. For the hybrid nanofluids with type A25S75, the value for ΔT_w was dropped first and then slowly increased monotonically respective to time. After 1 hour of experiment duration, the steady state condition was not achieved where the value of ΔT_w keep on increasing and the line (marked with a diamond symbol) located on top of the reference line (ΔT_w for distilled water) therefore it has a higher value of ΔT_w as shown in Figure 3.2. Same goes for the A75S25 type. The figure also demonstrated the ΔT_w was slight reduced for all nanoparticles addition in the first few seconds, however at later stage increased asymptotically respective to the time. In the present case, it could be seen that the effect of Al_2O_3 showed a considerable reduction in ΔT_w especially in the initial stage of nanofluids addition. However, the present of SiO_2 later diminished the positive effect. For example, one could be seen that in the similar hybrid volume ratio of ($\text{Al}_2\text{O}_3\text{-SiO}_2$)/water (A50S50 type), the ΔT_w was decreased and gradually shows the increasing value of ΔT_w . After 30 minutes, it passed above the ΔT_w value of reference line. The trend line for this particular ratio (A50S50 type) falls in the middle of those two single Al_2O_3 /water and SiO_2 /water nanofluids. It is interesting to note that, the ΔT_w

after 1 hour boiling time were at the consistent magnitude and depended on the ratio of nanofluids mixture. Therefore, it can be concluded that after the injection of hybrid nanofluids, the HTC was enhanced at the beginning, however, the HTC starts to deteriorate slowly respect to time.

Many possible reasons lied in these peculiar trends of the different ΔT_w between both types nanofluid (Al_2O_3 and SiO_2 nanofluids). Since the intial addition of nanofluids into the main vessel, the ΔT_w decreased drastically due to the nanoparticle deposition process onto the heater surface and possibly promoted activation of new active nucleation sites. Consequently, the heat transfer from heater surface to the liquid was enhanced as discussed by several researchers, i.e. in appendix A in M. Z. Sulaiman et al. [19] and Suriyawong et al. [29]. Similar to the present experiment, the number of active nucleation sites were possibly increased after the addition of the nanofluids even for the lowest concentration of Al_2O_3 /water (A25) in the present experimental work. However, in other cases, Suriyawong et al. [29] revealed that 0.0001 vol.% of concentration, the HTC was not affected after the additional of nanofluids because there was no deposition of nanoparticle occur on the heater surface. However, there is an argument about the increase in the number of active nucleation sites by Raveshi et al. [30] where he suggested that the porous nanoparticle layer reduced the nucleate site density and affected the enhancement of HTC multiplying cavities to smaller sites [31].

As for the highest ratio of Al_2O_3 /water (A100), the HTC was enhanced considerably with slight deterioration could be seen as boiling time increased. It is most probable to infer that the deposition of Al_2O_3 nanoparticle with low thermal conductivity onto the heater surface lowered the HTC value. In addition, the SiO_2 nanofluid with lower thermal conductivity had demonstrated different characteristics compared to Al_2O_3 /water nanofluid where after the injection of nanoparticles, the HTC at the initial stage was slightly increased, but the deposition of SiO_2 nanoparticle onto the heater surface was increased and resulted in the increasing of thermal resistance [32] and caused the HTC to reduced. For hybrid nanofluids, the active nucleation sites were increased at the initial stage due to the effect of improvement of the surface microstructure, however, the deposition of the SiO_2 impeding the positive effect from new active nucleation site activation and forming a thicker layer on top of the surface over the time. Consequently, the transient conduction beneath the bubbling bubble was distracted and the HTC was slowly reduced.

4. Conclusions

Experiments were conducted to investigate the heat transfer characteristics for nucleate pool boiling of two single nanofluids (Al_2O_3 /water and SiO_2 /water) and their hybrid at low level concentration of 0.001 vol.%. The heat transfer characteristics in these experiments represented by the time-variation of the wall superheat during nucleate boiling. Main conclusions of this work were summarised as follows.

- a) In most cases in all of the experiments, the wall superheat first decreased drastically and after a few second, it started to react depends on the type of nanofluids (except for the higher composition of SiO_2). For Al_2O_3 /water nanofluid, the HTC was enhanced and stable for type A25. However, for types A50 and A100, HTC was enhanced at first and then slowly deteriorates respectively with times. Even though the Al_2O_3 nanoparticle deposited on the heater surface will increase the active nucleation site, but due to the high volume of the nanoparticle deposited layer on the heater surface, the HTC observed to slightly decreased over a longer time duration.
- b) For the SiO_2 /water experiment, the HTC deterioration was observed for in all concentration, but the final value of ΔT_w after $t_b = 1$ hr were depended on the concentrations value, which higher for high concentration and vice versa. It could be noticeable seen in Figure 3.1 (b) that even in type S25, the HTC was reduced. This could be the inferred that the deposited SiO_2 /water on the heater surface was thickening the nanoparticles layer on the surface and increased the thermal resistance. Consequently, the heat flows were suppressed from moving from the heater surface to the distilled water.

- c) As for the hybrid nanofluids, the HTC were enhanced at the approximately first 30 minutes. After that, the ΔT_w slowly increase and pass over the reference line for all the experiments. First, the Al_2O_3 nanoparticle will be deposited on the heater surface and will increase the active nucleation sites, and the HTC was enhanced. However, in longer boiling duration, the SiO_2 nanoparticle with low thermal conductivity deposited on the heater surface and formed a thicker layer that possibly prevents the transient heat conduction from underneath the bubbling bubbles.

Conflicts of interest

I declare that there are no known conflicts of interest.

Acknowledgments

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