

CHARICTERIZATION AND INVESTIGATION OF HEAT TRANSFER ENHANCEMENT IN POOL BOILING WITH WATER-ZNO NANO-FLUID

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ABSTRACT:

The main focus of the present work is to characterize the ZnO nanoparticles further to prepare the ZnO nanofluid with base fluid as deionised water and to investigate enhancement in critical heat flux at different weight concentrations of nanofluids. The size of nanoparticles is found to be 55.25nm. To study Critical Heat Flux (CHF) enhancement using ZnO nanofluid, different weight concentration of nanofluid are prepared. It is observed that maximum enhancement 47.16 percent observed for 1.5 weight percent of ZnO nanofluid. Surface roughness and scanning electron microscopy of heater surface is carried out for all weight concentrations of nanofluid, which shows increase in Ra value up to some extent then it decreases and porosity on the surface of heater observed in SEM, is the source to enhance CHF.

Keywords: Nanoparticles, Nanofluid, Critical Heat Flux (CHF), Enhancement, Deionised Water (DI).

1. Introduction

Heat transfer is an important issue in many industrial applications. The heat transfer in the nucleate boiling regime, the latent heat of vaporization during the change from liquid to gas phase can be exploited and is the most effective way of cooling thermal systems operating at high temperatures [1]. However, the boiling heat transfer is restricted by the critical heat flux (CHF). This is highest heat flux where boiling heat transfer sustains its high cooling performance. When the surface reaches CHF, it becomes coated with a vapour film which isolates the heating surface and the fluid thus the heat transfer decreases drastically [1–3]. In these conditions, the wall temperature rises quickly, and if it exceeds the limits of its constituent materials, system failure occurs. For this reason, every system incorporates a safety margin by running at a heat flux lower than CHF, but this approach reduces system efficiency [1]. This compromise between safety and efficiency is a serious problem in the industry. For this reason, a huge work has been carried out to understand heat transfer mechanisms in nucleate boiling and CHF conditions and to increase the CHF.

Pioro et al. [4, 5] present a very fine review of the parametric effect of boiling surface and prediction methods. They show that it is complex problems involving many inter linked parameters affect heat transfer performances. Their analysis of the literature shows that some results seem contradictory. For example, some researchers conclude that for many practical applications the effect of solid/liquid/vapour interaction on the heat transfer coefficient in nucleate boiling conditions can be ignored (except for the cryogenic fluid), whereas others conclude that these effects are important [4]. Some studies have firm on evaluating the effect of surface characteristics on heat transfer performance. These parameters are typically the contact angle, thermo physical properties, thickness, orientation in space, roughness (surface finish), and microstructure (shape, dimensions, pore density for the vapour bubble generating centre) [4]. All these interlinked parameters simultaneously affect heat transfer performance. At the moment, there is not sufficient information to solve this complex problem and for this reason, only separate effects are considered [4]. Enhancement in CHF also noted in the literature for all nanofluids with different orientation and heater surfaces [6].

In present case, micro-structure and wettability are the most important aspects. These parameters are dealt with in the literature, remarkably; Kim et al. worked with Al₂O₃ and TiO₂ nano-fluid. They concluded that a nano-particle coating on a heating surface is a prime factor in enhancing the CHF of nano-fluids. The main factors that explain this behaviour are wettability and capillary wicking [7].

2.0: Structural and microstructure of ZnO nanoparticles:

The structural and microstructure properties of the ZnO nanoparticles are as follows; Figure 1 Shows XRD patterns of ZnO particles can be used to determine the size of the nanoparticles.

2.1 Structure analysis ZnO of Nanoparticles:

Table 1: Properties of ZnO nanoparticles

Items	ZnO
Content of ZnO	99.9 %
Average particle size	55.25 nm
Specific surface area	80 m ² /gm

Figure 1 shows the X-ray diffraction pattern (XRD) pattern of ZnO nanoparticles. All the peaks in diffraction pattern shows monoclinic structure of ZnO and the peaks, average grain size calculated by using Debye-Scherrer formula is approximately 55.25 nm. Debye-Scherrer formula,

$$D = 0.9\lambda / \beta \cos \theta \quad (1)$$

Where β is full width at half maxima of the peak in XRD pattern, θ is angle of the peak, wavelength of X-rays. Elastic strain is also calculated from XRD results. The strain results suggested that if the particle size is less than 20 nm than they have more strain and greater than 50 nm particles have less strain.

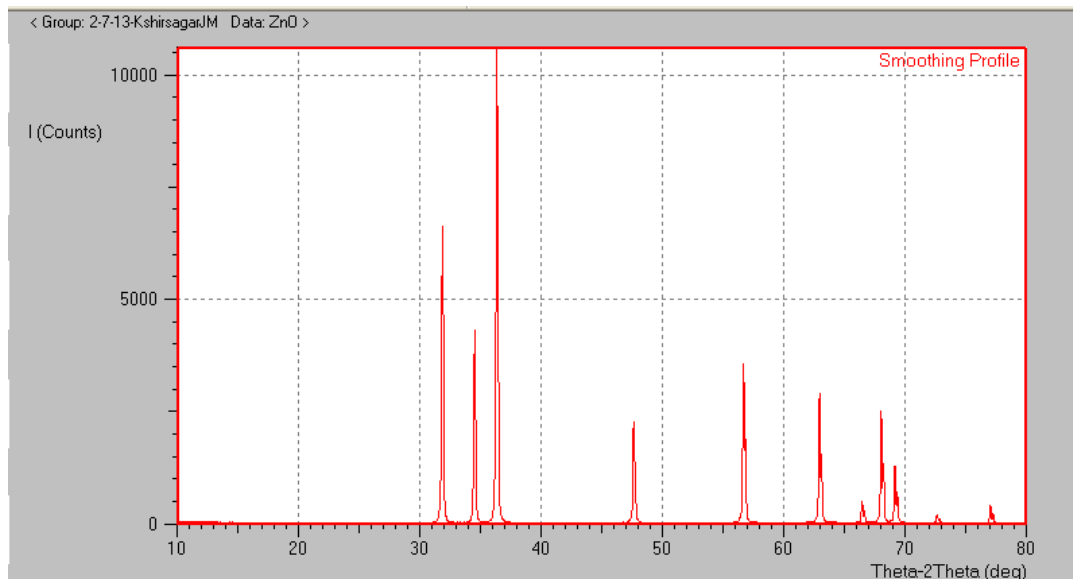


Figure 1: XRD pattern of ZnO nanoparticle

2.2 Preparation of Nano Fluid and Characterization.

The Zinc Oxide nanoparticles were dispersed in deionized water for 12 hours under high speed mechanical stirrer (Toshiba, India). No surfactant or stabilizer is used during the preparation of nanofluid as they have some influence on forced convective heat transfer coefficient as well as on overall heat transfer coefficient. After 24 hour no sedimentation of nanoparticles are found. Generally, the properties of the nano-fluid depend on the properties of the nano-particles and the surface molecules taking part in the heat transfer procedure depend on the size and shape of the particles themselves, which are also affected by the agglomeration of the particles. As shown in the figure3 taken by Transmission Electron Microscopy (TEM), the size has a normal distribution in a range from 50 nm to 55nm.

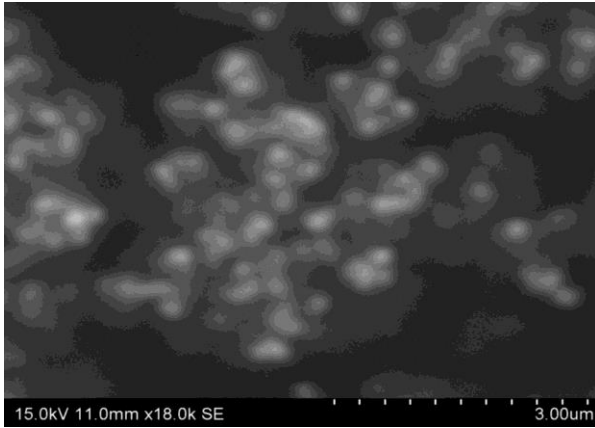


Figure.2. SEM image of ZnO particles

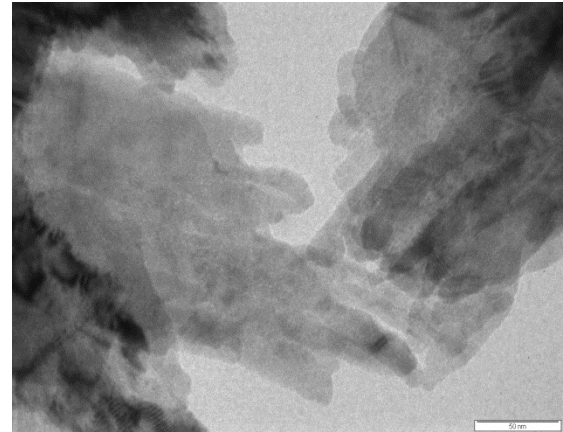


Figure.3. TEM image of ZnO particles.

The graph shown in Figure.1 depicts the X-ray diffraction spectra has highest intensity of 9177 counts at 36.39°. It is also observed that all zinc nanoparticles remained in pure zinc state. The SEM photograph of nanopowder is shown in Figure.2.

3.0 Determination of enhancement in CHF

The Ni-Cr wire having 0.321 mm diameter is used as heater surface. The length of the heater is 110 mm. For benchmarking the experiment the Zuber's correlation is used initially for deionised water, average CHF for ten experiments is found to 1.2 MW/m².

3.1 Theoretical determination of CHF

A number of experiments on bare Ni-Cr wires of 0.321 mm in diameter are carried out to examine the reproducibility of the experimental apparatus and get insights about the fundamental mechanism of the CHF phenomenon on the thin wire used in this study. The CHF values of pure water on bare wires showed good repeatability.

Methodology and correlation used are as below:

The well known Zuber's correlation is used for validation of the test set up. Experimental values of q''_{CHF} is compared with that as predicted by Zuber's correlation:

$$q''_{CHF} = \frac{\pi}{24} \rho_g^{0.5} h_{fg} [\sigma g (\rho_l - \rho_v)]^{0.25} \quad (2)$$

But, it is known that the effect of cylinder radius on the CHF for wires is significant [8-12]. You et al. [12] reported in their photographic studies that the CHF on small wires were two different mechanisms which proceed to film boiling: Hydrodynamic CHF and local dry out in which Power is transferred from a heated surface to deionized water it is desired to obtain high heat fluxes with low temperature difference, there is linear relationship between heat fluxes and temperature difference. If heat fluxes are increased bubbles nucleate at hot surface of heater wire and depart to the sub cooled fluid and collapse. If the heat flux more increased at some point a vapour film is formed on the surface of heater. the heat transfer rate suddenly decreased and wall temperature increased the value at which it occurs is called critical heat flux (CHF).

During the experimentation, condition at breaking of wire due to critical heat flux is noted and corresponding voltage and current are recorded. The CHF is calculated by following formula,

$$q'' = \frac{V \times I}{\pi DL} \quad (3)$$

Validation of experimental set is done for ten trials, result of that is shown in the figure 4.

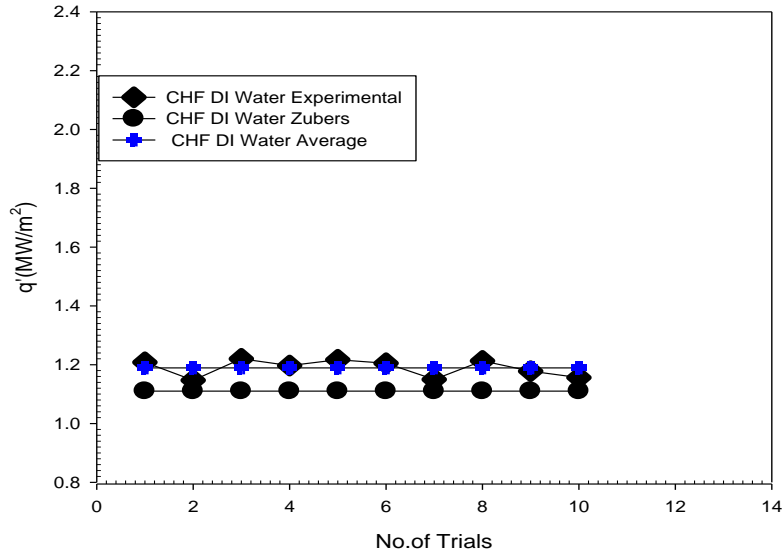


Figure 4: Critical heat flux curve experimental and theoretical Zuber’s correlation of deionised water

3.2 Uncertainty in CHF:

The main source of uncertainty of the applied voltage and current only due to contact resistance between the wire heater and electrodes connected with the clamps in addition to this uncertainty also associated with length and diameter of the Ni-Cr wire heater [13].

In this study, the uncertainties of the measured parameters were analyzed by the error propagation method. For example, Uncertainty of the heat flux was calculated as follows: Heat flux is calculated using Eq.(3). Thus the main source of heat flux uncertainty is found as voltage (V), current (I), diameter of heater (D) and length (L). Heat flux uncertainty can be calculated using the following equation:

$$\frac{\Delta(q)}{(q)} = \sqrt{\left(\frac{\Delta V}{V}\right)^2 + \left(\frac{\Delta I}{I}\right)^2 + \left(\frac{\Delta d}{d}\right)^2 + \left(\frac{\Delta L}{L}\right)^2} \dots\dots\dots(4)$$

The elemental percentage uncertainty of V, I, D and L is less than 1.63%, 3.49%, 0.31%, and 1.81% respectively. Thus:

$$\frac{\Delta q}{q} = \sqrt{(1.63)^2 + (3.49)^2 + (0.31)^2 + (1.81)^2}$$

$$\frac{\Delta q}{q} = 4.26\%$$

Therefore, the measurement uncertainty on the calculated heat flux is $\leq \pm 4.26\%$. For other parameters, a similar approach was used for calculating the measurement uncertainty.

4.0 Experimental setup

Figure 5 shows a schematic illustration of the experimental setup. The test facility consists of Borosil glass vessel of capacity 3 litres. The glass vessel is transparent to observe the actual phenomenon of bubble formation. On top of the vessel 16 mm thickness Bakelite sheet is used to cover test vessel. Rubber gasket is used for the perfect alignment on test vessel. On sheet two holes of 5 mm diameter each, are drilled at a distance of 55 mm from centre, i.e. the distance between two holes are 110 mm. These holes are used for inserting and holding copper electrodes. One hole of 20

mm diameter is drilled at the centre of the sheet to hold the condenser inlet, which collects the vapours generated in the test cell. The condenser is placed at the top of test cell. One 5 mm hole is drilled to accommodate K- type thermocouple is inserted in the test cell to measure the bulk temperature of the fluid during experimentation. An arrangement is done to hold the pre heater in the test cell on Bakelite sheet. This pre heater is used to heat bulk fluid in the test cell to saturation temperature. Vent is provided on the Bakelite sheet, so that atmospheric pressure always acts on the test cell. Two copper electrodes of size 5 mm (diameter) × 150 mm (length) are used for holding the test heater. The Ni-Cr wire of 28 gauge i.e. 0.321 mm diameter is used as heater surface. The length of the heater is 110 mm. The DC power supply with a capacity (230 V, 30 A) is used. The power supply has a provision to increase the power with a step of 0.1 V. The copper electrodes are connected to power supply with 10 gauge copper wire and copper lugs to have minimum resistance. Two pipes of ¼ inch diameter are used for circulation of cold water and hot water from the condenser, so that, the vapours form are converted to water and the volume of test cell remains unchanged. K- Type thermocouple used for data logging. The Data recorded by using RISHABH multi SI 232 data card. The shunt register (200A by 75 mV rating), connected in series with the test wire, that is used to measure the current through the wire.

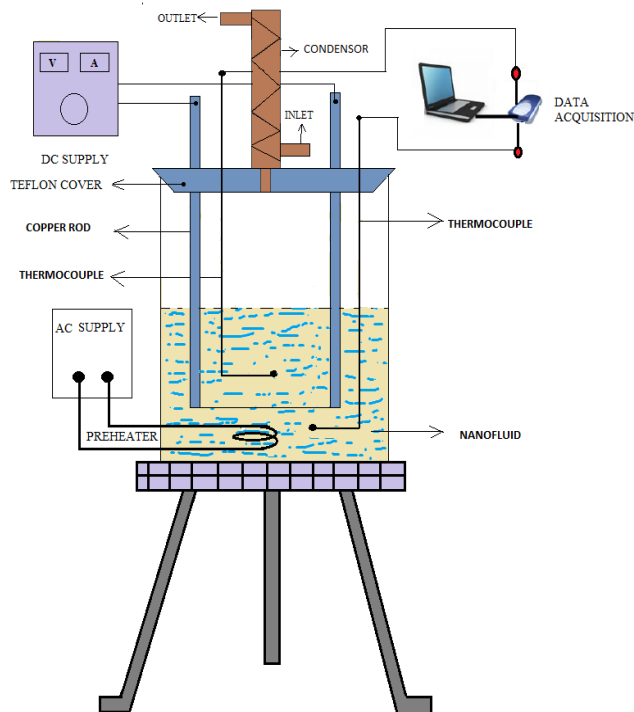


Figure 5: Schematic diagram of test rig

According to Asakura et al. [17,14], nanoparticles deposition on a surface during nucleate boiling is mainly influenced by heat flux of the heated surface, boiling time, concentration of nanoparticles and pH of the suspension.

5.0 Results and discussions:

Pool boiling experiment is carried out using ZnO nanofluid and critical heat flux is compared with deionised water. Significantly improvement in critical heat flux is observed. Figure 6 shows critical heat flux values at different weight concentration of ZnO nanofluid.

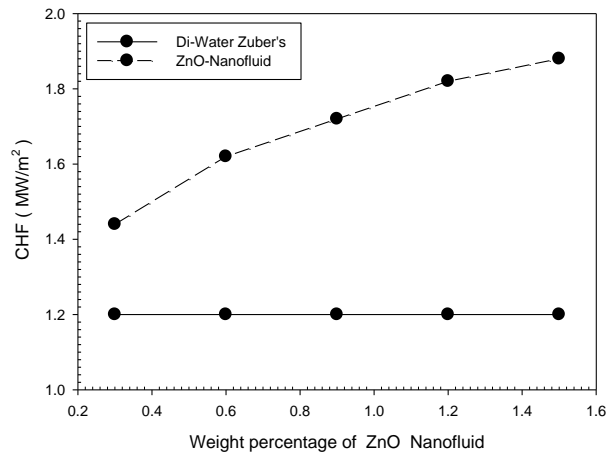


Figure 6 : Variation in CHF with different weight concentration of ZnO nanofluid

The critical heat flux (CHF) of nanofluid has been compared with pure water by various researchers. The critical heat flux enhancement is nearly 73 percent for stain steel wire with Al_2O_3 , ZrO_2 , and SiO_2 nanofluid, Kim et al.[9] the enhancement in CHF is only due to increase of contact area by deposition of nanoparticles over the heating surface. The nanoparticles generate porous layer on the test section tube surface thus reducing the contact angle between the fluid and heater surface [15]. The existence of sorption layer enhance the trapping of liquid in nano porous sorption layer and prevents the vapour blankets formation, so that the CHF increases with increasing the sorption layer thickness at lower particle concentration range.

Kshirsagar et.al [16] studied pool boiling with water based nanofluid and CuO nanoparticles. The results showed that the water-based nanofluid significantly enhanced CHF compared to that of pure water. The CHF values of the CuO nanofluid were enhanced from approximately 57.26 percent as compared to deionised water. It was found that a sizable layer of nanoparticles deposits were formed on heater surface.

Ramakrishna Hegde et.al[17] investigate CHF enhancement using CuO nanofluid relative to CHF of pure water. CHF values were measured for various volume concentrations, CHF enhancement 130% recorded for 0.2 volume percent of CuO.

The results show that CHF enhancement is definitely possible by using different nanofluid instated of deionised water for cooling fluids. However the surface Coating of heater surface is depending on particles concentrations of the nanofluid, a layer build up which may increase or decrease surface roughness that will depend upon porosity and nanoparticles size. The surface roughness measurement is taken when the test surface exposed to different weight concentration of nanofluid after pool boiling critical heat flux tests made, known that critical heat flux enhancement of nanofluid is closely related with the surface microstructure and enhanced surroundings from the deposition of nanoparticles as shown in figure 7.

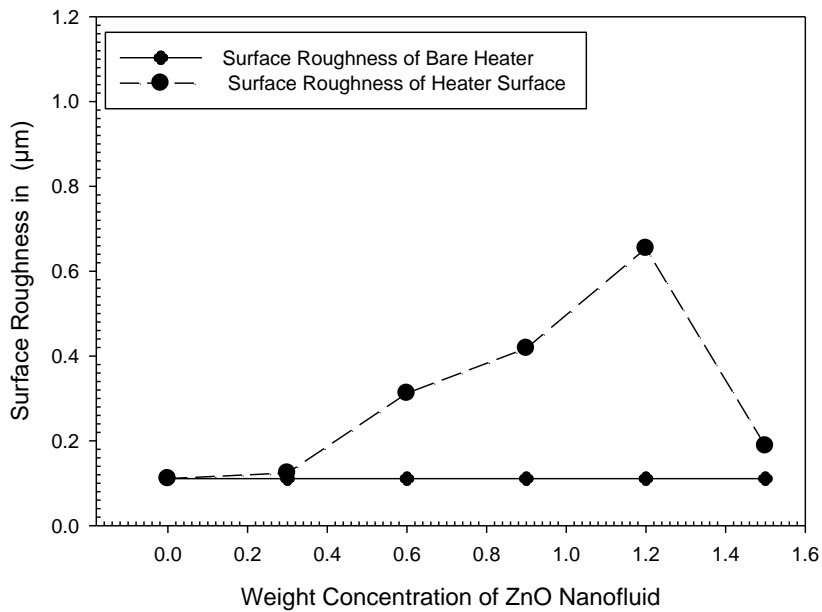


Figure 7: Increase in Surface Roughness with Weight Concentration of Nanofluid

Scanning electron microscopic image of heater surface are taken on which it is clearly observed that increase in surface roughness, SEM of bare heater also taken before starting the experiment. Bare heater roughness is measured is found to be 0.111 µm as shown in figure 8 (a).

For all weight concentration of nanofluid SEM images are taken after conducting the experiment. The increase in roughness from 0.3 weight Percent to 1.2 weight percent observed on the heater surface figure 8(b) to figure 8(e) and for 1.5 weight concentrations surface roughness suddenly decreases up to 0.188µm.

The study of SEM image shows that deposition of nanoparticles on heater surface. So the porosity plays important role in boiling heat transfer even changes roughness and wettability.

Heater Surface	Scanning Electronic Microscopy Image	Surface Roughness
(a) Bare Heater		 Ra=0.111 µm
(b) 0.3 Wt.Percent ZnO		 Ra=0.124 µm

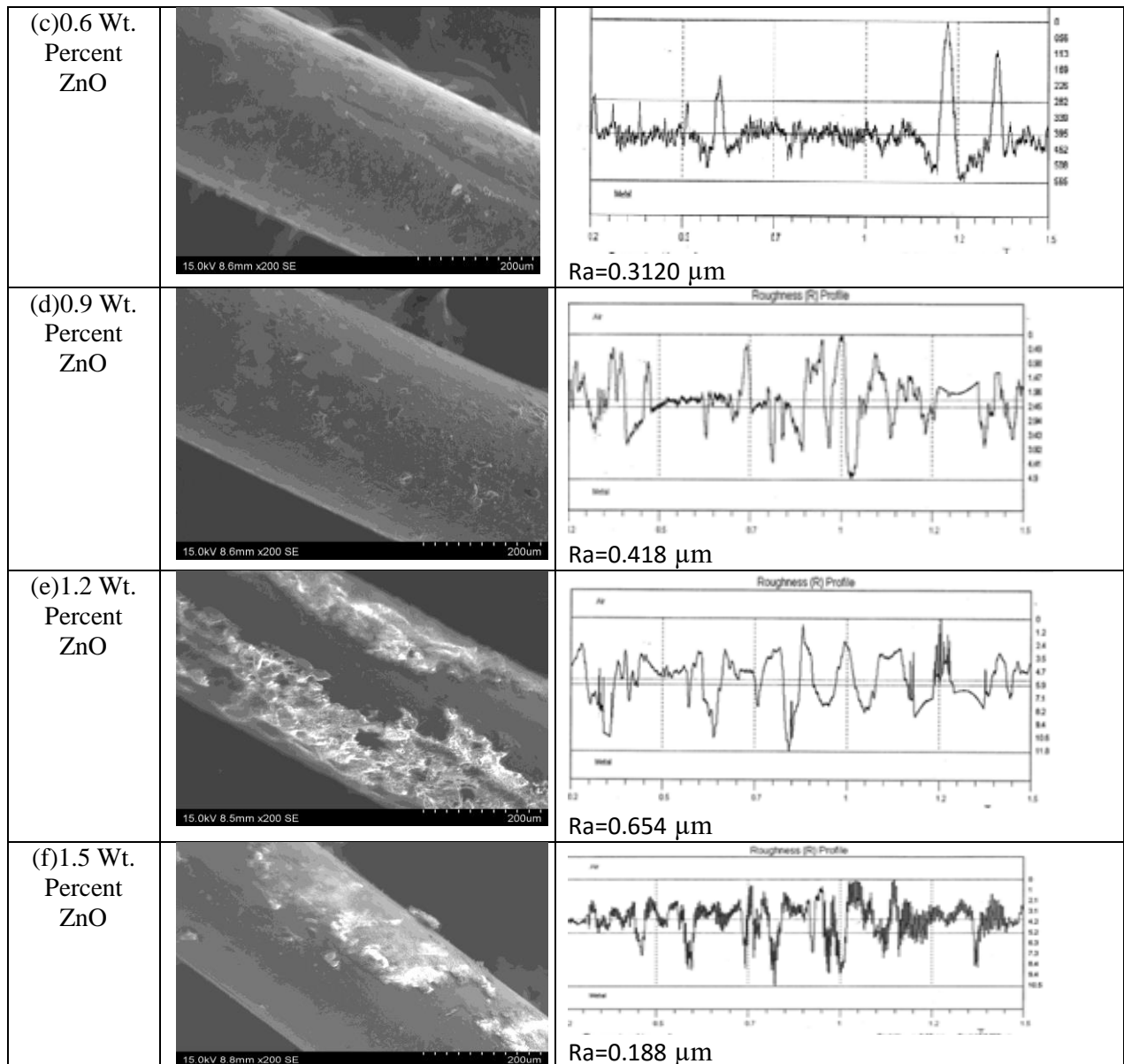


Figure 8 : (a-f) SEM image showing nanoparticles deposition on heater surface with different wt. Concentration of nanofluid and surface roughness of heater surface

Conclusion:

In this study, pool boiling CHF behaviours on electrically heated NiCr wire with nano-porous deposits were investigated at saturated temperature under atmospheric pressure. Various deposit structures were obtained by varying the increasing rate and maximum value of heat flux during the pre-boiling in different weight concentration of water-ZnO nanofluid.

Surface properties of the deposit wires were characterized to identify the major surface parameters associated with the significant CHF increases on a heater with nanoparticles deposits. It was found that heat capacity of heater surface in the present study changes due to the nanoparticles deposit so that it is not a major parameter in interpreting the increased CHF. Pool boiling characteristics in ZnO nanofluid were investigated with five weight concentrations 0.3, 0.6, 0.9, 1.2 and 1.5 weight percent of ZnO with deionised water. The enhancement in CHF for every concentration is studied; the maximum enhancement of 47.16 % CHF takes place at 1.5 weight percent of ZnO and for 0.3 weight per cent 13.88%.

During experimentation it is observed that roughness on heater surface increases up to 1.2 wt. percent after that it suddenly decreases, even CHF increases slowly from 1.2. to 1.5 wt. percent , rate of enhancement in CHF decreases from 1.2 to 1.5 wt. Percent of ZnO nanofluid.

CHF enhancement is directly related with surface microstructure and enhanced topography resulting from the deposition of nanoparticles SEM image of the heater surface shows porous layer build up due to boiling induced precipitation of nanofluid. Decrease in surface roughness can be attributed to nanoparticles filling the micro cavities formed over the heater surface.

Nomenclature:

Abbreviations

h_{fg}	latent heat of vaporization, [kJ/kg]
T_s	temperature of heater surface, [°C]
T_{sat}	saturation temperature, [°C]
q''_{CHF}	Critical heat flux, [MW/m ²]
A	area of heater surface, [m ²]
D	diameter of heater, [m]
L	length of heater, [m]
ρ_l	density of liquid
ρ_v	density of vapour
CHF	Critical Heat Flux
SEM	Scanning Electron Microscopy
I	current [ampere]
V	voltage [volts]
R_a	average roughness[μm]

Greek symbols

Δ	difference
σ	surface tension, [N/m]
λ	X- ray wavelength of vapour jets, [m]
θ	peak, [radian]
β	full width half maxima of the peak, [radian]
ν	frequency, [Hz]
ρ	density, [kg/m ³]

Subscripts

C	Critical
HF	Heat flux
fg	Vaporization
sat	Saturation
l:	Liquid
v	Vapour

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