

# Investigation of Analogy between Boiling and Hydrogen Evolving System in Nucleate Bubble Regime

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## 1. Introduction

Operating a heat transfer device in boiling mode is preferable due to the high heat transfer rate compared to the single-phase heat transfer. Therefore, nucleate boiling has raised wide research interests worldwide [1]. Also in nuclear areas, there have been numerous efforts to investigate the nucleate boiling, since all the nuclear power plants have steam generator [2]. However, the boiling experiments were not performed sufficiently due to the high power density, extreme thermal conditions, measurement difficulties, etc.

The present study aimed at simulating the saturated nucleate pool boiling phenomenon using Hydrogen Evolving System (HES). The reduction of hydrogen ions by electrochemical process substituted the vaporization process in the boiling system. The basic idea is that the hydrodynamic behavior of the both systems should be analogous. And our research group previously performed the related studies [3,4]. 1.5 M of sulfuric-acid ( $H_2SO_4$ ) solution was used as working fluid and some bubble parameters such as nucleation site density ( $N_a$ ), bubble departure diameter ( $D_b$ ) and bubble frequency ( $f$ ) were measured using thin wire and vertical disk plate as cathode surface, which simulated heating surface.

## 2. Theoretical Backgrounds

### 2.1 Nucleation site density

Gaertner and Westwater [5] observed that  $N_a$  increased with the heat flux as expressed in Eq. (1)

$$N_a \sim q^{2.1}. \quad (1)$$

Paul and Abdel-Khalik [6] measured  $N_a$  using platinum wire and water at saturated condition. The individual bubble sites were counted at each heat flux step using high-speed camera. The results were fitted as

$$N_a = 1.207q'' - 1.574 \times 10^{-2}. \quad (2)$$

Yeom et al. [7] examined the influence of nanoparticle surface on the  $N_a$  using zirconium wire and water at saturated condition. The  $N_a$  was counted at each heat flux value up to the CHF. The  $N_a$  showed peak value before the CHF point, irrespective of the surface conditions.

Therefore, the  $N_a$  increases as the heat flux increases generally.

### 2.2 Bubble departure diameter

Fritz [8] developed the correlation, Eq. (3) to predict  $D_b$  introducing using contact angle of the bubble. Cole [9] developed the correlation using  $Ja$ , Eq. (4).

$$Bo_d^{0.5} = 0.0208\theta. \quad (3)$$

$$Bo_d^{0.5} = 0.04Ja,$$

$$\text{where } Bo_d = g\Delta\rho D_b^2/\sigma \text{ and } Ja = C_p\Delta T/h_{fg}. \quad (4)$$

Paul and Abdel-Khalik [6] and Yeom et al. [7] measured the  $D_b$  with respect to the heat flux with the identical apparatus measuring the  $N_a$ . Paul and Abdel-Khalik [6] found linear relationship between  $D_b$  and the heat flux. The  $D_b$  was also measured for individual bubble. However, Yeom et al. [7] reported that the bubble departure volume, which is proportional to the third power of the diameter, increased exponentially according to the heat flux up to the CHF point irrespective of the surface condition.

### 2.3 Bubble frequency

The liquid inertia carries the bubble away from the heating surface [10]. The time interval  $t_d$  is required for bubble to detach from the surface. Then the bulk liquid rushes after the bubble detachment and the time interval  $t_w$  is required for a subsequent nucleation [11]. Thus, bubble frequency can be expressed by

$$f = \frac{1}{t_w + t_d}. \quad (5)$$

Paul and Abdel-Khalik [6] calculated  $f$  based on the  $D_b$  data using frequency distribution function and obtained a linear relationship according to the heat flux. Yeom et al. [7] measured  $f$  by counting image frames for  $t_w$  and  $t_d$  and defined  $f$  as function of the heat flux up to the CHF. A peak was measured irrespective of the surface condition due to the bubble coalescence at a certain high heat flux condition.

## 3. Experimental setup

### 3.1 Test apparatus and electric circuit

Figure 1 shows the experimental apparatus and electric circuit. Two geometrical types of cathode were employed: 0.2 mm thick horizontal copper wire and vertical copper disk of 40 mm diameter. Three bubble parameters,  $N_a$ ,  $D_b$  and  $f$  were measured using high speed camera with wire cathode at high current density. And in order to observe increased  $N_a$  at low current density, vertical disk cathode was employed. The cathode and the anode were located in the glass container filled with the aqueous solution of sulfuric acid ( $H_2SO_4$ ) of 1.5 M at atmospheric pressure and room temperature, 294 K. The high-speed camera recorded the hydrogen bubbles at the cathode surface. The electric current was controlled using the power supply (N8952A, Keysight).

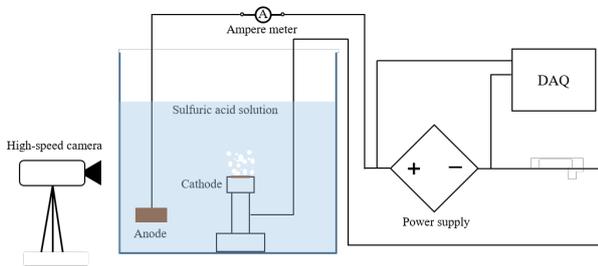


Fig. 1. Experimental apparatus and test section.

### 3.2 Test matrix

Table I sorted current density range. Low and high current densities were applied using vertical and horizontal cathode, respectively.

Table I: Range of current density for experiments

Current density scale (Cathode geometry)	Current density range ( $A/m^2$ )
Low current density (Vertical disk)	4.1–49.3
High current density (Horizontal wire)	3,900–94,700

## 4. Results and discussion

### 4.1. Nucleation site density

Figure 2 shows nucleation sites of the present hydrogen evolving system with respect to the current density at vertical disk within low current density range. The nucleation sites were randomly distributed on the cathode surface. The  $N_a$  increased as the current density increased, which is similar trend to the boiling system. However, it is difficult to quantify the  $N_a$  at the present system due to the numerous bubble sites.

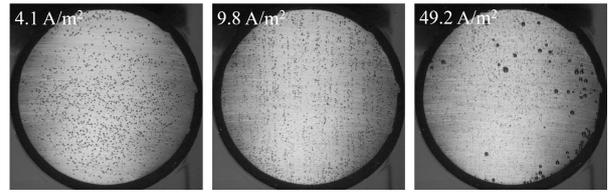


Fig. 2. Hydrogen bubble behaviors on the cylindrical ribbon.

Figure 3 compares the  $N_a$  between the boiling system and the present hydrogen evolving system at high current density range. Yeom et al. [7] measured the  $N_a$  up to the CHF point. They observed a peak value due to the bubble coalescence. Paul and Abdel-Khalik [6] also reported the similar trend. However, as they only measured the isolated bubble site at low heat flux, they could not observe the peak. Meanwhile in the present work, the bubble departure sites were measured as the  $N_a$  could not be measured due to the vigorous bubble coalescence at the high current density range. They decreased exponentially as the current density increased. Therefore, the  $N_a$  behavior and bubble departure site density in the present work are similar to that of the boiling system.

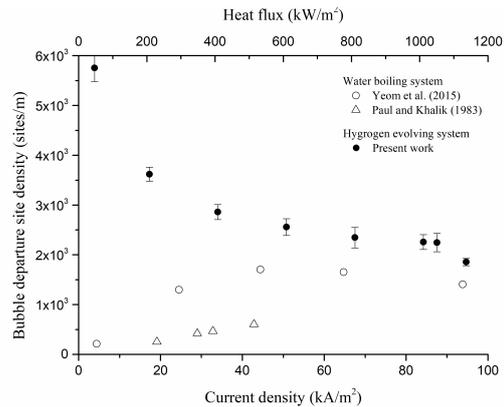


Fig. 3. Bubble departure site density of the boiling and the hydrogen evolving system.

### 4.2. Bubble departure diameter

Figure 5 shows the  $D_b$  of the boiling and the present hydrogen evolving system. Yeom et al. [7] and Paul and Abdel-Khalik [6] found the  $D_b$  increased as the heat flux increased. Similarly, the  $D_b$  of the present hydrogen evolving system increased as the current density increased as shown in Fig. 6. It is because of the vigorous bubble coalescence at the high heat flux and current density regime, which is predominated by the hydrodynamic phenomenon. However, the  $D_b$  in the hydrogen evolving system was smaller around 10% than that of the boiling system. Vogt et al. [12] insisted that the cell potential affects the wettability of the surface.

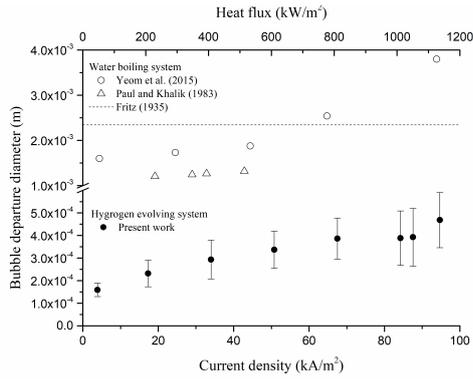


Fig. 5. Bubble departure diameter of the boiling and the hydrogen evolving system.

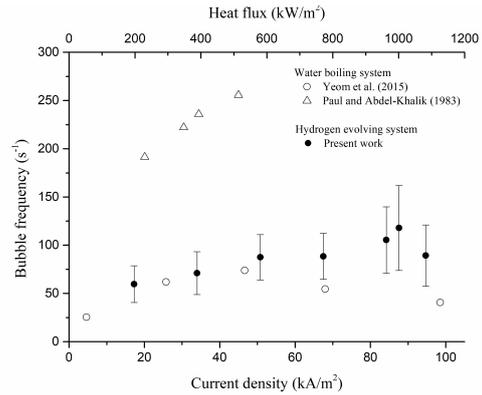


Fig. 7. Bubble frequency of the boiling and the hydrogen evolving system.

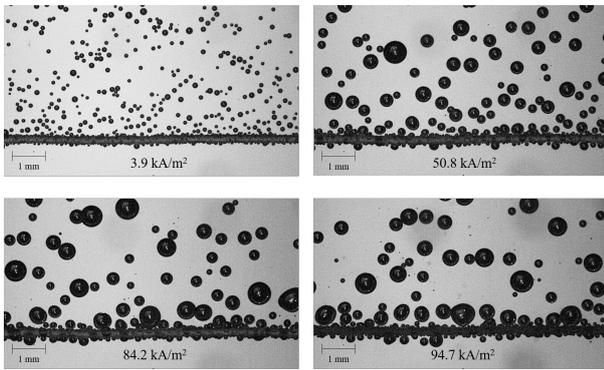


Fig. 6. Hydrogen bubble behaviors on the cylindrical ribbon.

#### 4.3. Bubble frequency

Figure 7 represents the  $f$  of the boiling and hydrogen evolving system. In the boiling system, Paul and Abdel-Khalik [6] calculated the  $f$  based on the  $D_b$  and frequency distribution function. It showed the linear relationship with the heat flux. Also Yeom et al. [7] measured a peak value of  $f$  with respect to the heat flux due to the coalescence of the bubbles at the surface. The similar tendency of the  $f$  was measured in the hydrogen evolving system having a peak value with respect to the current density. Moreover, the absolute values of the  $f$  in the present work and Yeom et al. [7] were similar. It can be deduced that the existence of the peak value would be affected by the vigorous bubble interaction near the surface, similar to the boiling system. And thus it implies the hydrodynamic similarity between the two systems.

### 5. Conclusions

Saturated nucleate pool boiling phenomenon was simulated by the hydrogen evolving system using 1.5 M of sulfuric-acid solution and copper electrodes. The nucleation site density, the bubble departure diameter and the bubble frequency were measured by taking photographic image using high-speed camera. The similarities and the differences of the bubble behaviors between the hydrogen evolving and boiling system were discussed.

The nucleation site density according to the current density showed similar trend at the low heat flux and current density, since both the hydrogen and bubble generation mechanisms are dominated by the heterogeneous nucleation. The nucleation sites were increased as the heat flux or current density increased. However at the high heat flux and current density, the coalescence of bubbles reduced the bubble departure sites in both systems.

The tendency of the bubble departure diameter was similar between the two systems, meanwhile the diameters of the hydrogen bubble are small, around 10% of the vapor bubbles. The bubble frequency of the two systems were also similar, which showed peak values.

It is concluded that there is a certain analogous phenomenon between the boiling and the hydrogen evolving system in the hydrodynamic perspectives. The authors expect that the present work may contribute to establish the analogy research between the hydrogen evolving and the boiling system. However, the further investigation should be performed to establish improved analogy concept regarding the cell potential affecting the surface wettability and the surface tension.

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