Review on Heat Transfer in Boiling of Binary Mixtures

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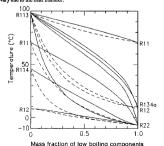
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Abstract

Heat transfer is described in nucleate pool boiling and forced convective boiling of pure Itquid and their non-accountage mixtures at low mass flow rate where the Ocean Thermal Energy Conversion system and the Kalina cycle may operate using a limited temperature difference between a hotter flouid on the sea surface and a cooler liquid at a denth of the sea.

1. Introduction

Though heat transfer in boiling of mixture was first paid attention in many chemical engineering process, it recently has received attention for improvement in a thermal efficiency in a power plant and in a coefficient of performance in a heat pump and refrigeration cycles. In the case of pure liquid, its saturation temperature is fixed at a constant value under a constant pressure, while for a binary mixture, its temperature does change along a boiling point curve on a phase equilibrium diagram with a variation of concentration as shown in Fig. 1. In addition to this, its change is responsible for the difference between corresponding saturation temperatures for the liquids. The characteristics that the saturation temperature of the binary mixture can be changed by the concentration, makes it suitable to enhance the thermal efficiency of the Lorentz cycle during which the temperatures of heat source and heat sink vary due to the heat transfer.



mass fraction of low boiling components

Figure 1 Phase equilibrium diagram of mixtures at p = 4 bar

(— Dow point curves; --Boiling point curves)

More recently, in order to improve the thermal efficiency of the Ocean Thermal Energy Conversion system which is designed to operate at a limited small temperature difference between a hotter liquid on the sea surface and a cooler liquid at a depth of the sea, one conceives a binary mixture as a candidate of test liquid. Kalina proposed the Kalina cycle with a regenerator, a separator of vapor and liquid and a absorber and succeeded in improving its thermal efficiency by using the mixture of water and ammonia. Since these systems generally make use of a small temperature difference for energy conversion, in these evaporator tubes, wall superheat or heat flux and in addition to these mass velocity(flow rate) naturally become relatively low.

Therefore, the state of knowledge on boiling heat transfer first in pure liquids and then their non-azeotropic mixtures will be described only at a low mass velocity here. It may be necessary to say that although heat transfer problems caused during the operation of the cycles have so far received considerably less attention than those of the single component systems due to a complexity of the heat transfer process, much attention becomes stressed on these problems involved.

2. Features of Flow Structure in a Heated Tube

The forced convective boiling in an evaporator tube is categorized by the development a flow configuration along the tube length because of the variation in flow state due to phase change by heat addition. Therefore, before discussing heat transfer problems we have to keep the flow situation formed in the tube in mind.

Figure 2 illustrates schematically liquid-vapor flow patterns in a vertical evaporator tube and in a horizontal evaporator tube at low and high mass flow rates, respectively. As shown in Fig.1, the features of the flow are significantly different between vertical and horizontal ones due to an effect of the gravity. This difference becomes of importance in evaluating the heat transfer rates. In the case of the vertical flow, roughly speaking, the liquid and vapor symmetrically flows along the tube length because of no gravitational effect. The flow pattern changes from bubble flow through slug flow and annular flow to mist flow. The position at which the flow pattern changes, strongly depends both on the mass flow rate and on the vapor generation rate which it is controlled by either heat flux or temperature difference. In the case of a low mass flow rate and a small temperature difference, the bubble and annular flow regions becomes relatively longer as compared with the regions of the another flow patterns.

In addition to this, in the case of the horizontal flow since an inertia

force is coupled with the gravitational force, the development of the flow pattern strongly depends on the mass flow rate. As a result, the flow pattern strongly depends on the mass flow rate, or example, the flow pattern the vertical tube. At a low mass flow rate, or example, the flow pattern changes from bubble flow through slug flow and wavy flow to mist flow. In the wavy flow, the liquid and vapor flow separately in the lower and upper parts of the tube. At a high mass flow rate, the flow pattern develops from bubble flow through slug flow and semi-annular or wavy-annular flow, amulair flow, finally to mist flow. In the annular flow, a liquid film covers the whole tube perimeter on which its thickness usually varies from the top to the bottom due to gravity.

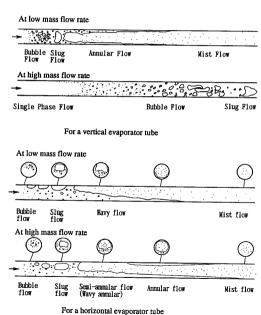


Figure 2 Flow patterns in a vertical and horizontal evaporator tubes at low and high mass flow rates

3. Characteristic of Flow Boiling Heat Transfer

Although the flow boiling heat transfer in the evaporator is strongly affected by the flow pattern, it generally consists of two different mechanisms; one is nucleate boiling of the liquid and the other is forced convection of the liquid with evaporation at the liquid-vapor interface. Since heat transfer is related closely to the flow pattern, it should be separately treated with depending on whether the surface is submerged in a large volume of liquid like a bubble flow or covered with a thin liquid film like an annular flow, and then in case of the thin liquid film on whether the direction of the surface is vertical or horizontal as shown in Fig.2. When our attention is focused on the low mass flow rate, the length of the transition region like a slug flow appearing between bubble and annular flows would be expected to be relatively short. Therefore, the heat transferred at the low mass flow rates is mainly governed by both the nucleate boiling and the forced convection. The heat transfer will be separately described here for the pure liquid and the binary mixture.

4 Correlation of heat transfer for pure liquid

4.1 Nucleate boiling heat transfer

The nucleate boiling heat transfer plays in essential role in heat transfer in the bubble flow region and then some contributions are always drawn into the another flow pattern. Therefore, it is of importance to know the characteristic of nucleate boiling even in dealing with heat transfer in flow boiling.

Many correlations are so far proposed to predict the heat transfer coefficients for the nucleate boiling as listed in Table 1. The nucleate beat transfer is subject to the properties of any particular liquid-surface combination but is hardly affected by the forced convection(the mass flow rate) and the orientation of the surface that would be easily understood from the fact that these effects are not included into any correlation in Table 1. It is very difficult to recommend which correlation is the best appropriate because each one inherits from what effects is mainly taken into account in its derivation.

4.2 Convective heat transfer in the vertical evaporator tube

There exist two ways of how to predict the heat transfer the annular flow. One is based on a relation of the form:

$$\frac{h_{TP}}{h_{TO}} = f(X_{H}, B_{O}) \tag{1}$$

where h_{LO} is the heat transfer coefficient if the pure liquid was flowing through the tube, X_{tt} is the Lockhart-Martinelli parameter, which plays an important role in determining the flow pattern:

$$X_{\mu} = \left(\frac{1-x}{x}\right)^{0.9} \left(\frac{\rho_{\nu}}{\rho_{\nu}}\right)^{0.5} \left(\frac{\mu_{L}}{\mu_{\nu}}\right)^{0.1}$$
 (2)

$$B \ o = \frac{q}{G \ H} \tag{3}$$

Table 2 gives an example of this types of correlations. The second one is

$$h_{TP} = S h_{NR} + F h_{cor} \qquad (4)$$

where has is the local heat transfer coefficient, S han is the contribution due to the nucleate boiling, F hcon the contribution due to convection with evaporation, how is the heat transfer coefficient of ordinary pool boiling and hCON is the heat transfer coefficient for the two phase flow. The mostly used correlation for saturated boiling in the annular flow is the Chen correlation[14] and later his original correlation was modified by Bennett and Chen[15] which succeeded in correlating all the forced convective boiling heat transfer data for water and organic systems. Since this correlation stands on the assumption that both mechanisms occur to some degree over the entire range of the correlation and the contributions made by the two mechanisms are additive, the multiplier, F, means the effect of the increase in liquid velocity, which is caused by the existence of vapor flow and then the other factor, S, means the effect of the suppression of nucleate boiling due to the steeping of the temperature gradient in the liquid film on the wall, which is caused by the forced convection. Chen proposes F hCON using the multiplier F and the Dittus-Boelter correlation as:

$$h_{CON} = 0.023 \frac{k_L}{D} \left(\frac{G(1-x)D}{\mu_L} \right)^{0.8} \left(\frac{c_L \mu_L}{k_L} \right)^{0.4}$$
 (5)

and

$$F = \frac{\left(\frac{dp}{dx}\right)_{Tp}}{\left(\frac{dp}{dx}\right)_L} \begin{cases} \frac{1}{2} & \text{if } X_s \le 01 \\ \frac{dp}{dx} & \text{if } X_s \ge 01 \end{cases}$$

(6)[16]

and S h_{NB} using the Forster and Zuber[10] analysis and the suppression factor as:

$$h_{NB} = 0.00122 \left(\frac{k_L^{0.79} c_P^{0.45} \rho_L^{0.45}}{\sigma^{0.5} \mu^{0.24} h^{0.24} \rho_L^{0.24}} \right) \Delta T_{sat}^{0.24} \Delta P_{sat}^{0.75}$$
(7)

where
$$\Delta T_{sst} = T_w - T_{sst}$$
 and $\Delta p_{sst} = p_{sst}(T_w) - p_{sst}(T_{sst})$ and
$$S = \left(\frac{1}{1 + 2.53 \times 10^{-6} F^{1.46} \text{ Re}_{s}^{-1.17}}\right)$$

It might be expected that the suppression factor, S, would approach unity at low flows and zero at high mass flows. Chen[14] suggests that S can be first represented as a function of the local two-phase Reynolds number, $Rep_1 = \mathbb{R}^{1.25} Re_1$, $Re_1 = \mathbb{G}(1+x)D|\mu$, and the Bennet and Chen[15] improved it to give equation(8). The functions F and S were determined empirically from experimental data using an iterative procedure to obtain the best solutions. It is necessary to say finally that the other correlation for the nucleane boiling is available though Chen employed only the Forster and Zaber correlation, while most of correlations employed the Ditms-Boelter correlation for convective heat transfer.

Recently, Oungor and Winterton[17], for example, derived a new generalized correlation using the Cooper correlation[9] in Table 1 for the nucleate boiling and the Dittus-Boelter correlation and determined the two factors F and S as:

$$F = 1 + 24000 Bo^{1.16} 1.37 (1 / X_{_{B}})^{0.86}$$
(9)

$$S = \left(\frac{1}{1 + 2.53 \times 10^{-6} F^{1.46} Re^{-1.17}}\right)$$
 (8)

It is concluded that flow boiling heat transfer, for saturated and subcooled conditions, vertical and horizontal flow, tubes and annuli, can be predicted with reasonable accuracy for over 4300 data points for waters, refrigerants and ethylene glycol and so on by their correlation.

4.3 Convective heat transfer in the horizontal evaporator tube

Figure 2 shows that the flow patterns in the horizontal tube are different from those in the vertical tube, especially in the annular flow where most of the heat exchange is done at the low mass flow raz, liquid and vapor separately flow each other. As the result, the difference in heat transfer between the upper and lower parts becomes significantly large. Taking account for this respect, Yoshida et al.[18] developed their method to predict the heat transfer in the vertical tube based on the Chen concept in which the circumferential averaged or axially local heat transfer coefficient for the separated and annular flows is treated with respectively during the flow boiling inside horizontal smooth tubes. This method shows the best precision in predicting flow boiling heat transfer open refrigerants among correlation prospeed hitherto.

The procedure proposed by Yoshida et al.[18] is introduced as follows:

(1) Calculate a separation angle o, with equation (11)

$$\frac{\phi_{x}}{\phi_{x}} = 1 + 0.72 \left[\left(\frac{1 - x}{x} \right)^{0.5} \left(\frac{\rho_{x}}{\rho_{y}} \right)^{0.5} \right]^{2} \left(\frac{\mu_{L}}{\mu_{y}} \right)^{0.17} \tag{9}$$

where

$$n = 0.22 \left(\frac{G^2}{gD\rho_{\nu}(\rho_L - \rho_{\nu})} \right)^{0.98} \left(\frac{q}{GH_{\dot{H}}} \times 10^4 \right)^{-0.96} \left(\frac{gD}{H_{\dot{H}}} \times 10^3 \right)^{-0.27}$$
(10)

and ϕ_0 is a separation angle estimated by assuming that there is no slip between the liquid and vapor flows and that the liquid-vapor interface is a horizontal plane.

(2)When φ_s > 162 deg, set φ_s equal to 180 deg. In this case the flow regime is annular. Then, calculate the local heat transfer coefficient by equation(11) which is applicable to the annular flow.

$$h_{TP} = S h_{NB} + F h_{con}$$
 (11)

where

$$h_{CON} = 0.023 \frac{k_L}{D} \left(\frac{G(1-x)D}{\mu_L} \right)^{0.4} P_{T_L}^{0.4}$$
 (12)
 $F = 1 + 2 X_{-}^{-0.82}$ (13)

$$h_{NB} = 207 \frac{k_L}{D_b} \left(\frac{qD_b}{k_L T_b} \right)^{0.745} \left(\frac{\rho_v}{\rho_L} \right)^{0.581} Pr_L^{0.533}$$
 (14)

$$D_s \approx 0.51 \left[\frac{2\sigma}{g(\rho_L - \rho_{\gamma})} \right]^{es}$$
 (15)

$$S = \frac{1}{1 + 0.9 \left[\frac{G(1-x)D}{\mu_L} x F^{\frac{125}{104}} \right]^{0.5} \left(\frac{q}{GH_H} x 10^4 \right)^{-0.5} X_{\alpha}^{-0.5}}$$

(3)When \(\phi_i < 162\) deg, the flow regime is a separated one. In this case, the circumferential averaged heat transfer coefficient must be calculated by taking account of the circumferential heat conduction within the tube wall</p>

In the case of a material with a high thermal conductivity such as copper, the variation of the wall temperature is approximately compensated by the circumferential heat flow so that the heat transfer rate in the dry part of the tube perimeter can be ignored. The circumferential averaged or axially local heat transfer coefficient can be calculated from equation (17), where only the heat transfer in wet part is considered.

$$h_{TP} = \frac{\phi_s}{180} h_{wet} \tag{17}$$

where h_{wet} is the heat transfer coefficient for the wet part and is calculated from equation(18)

$$h_{\text{ner}} = F h_{\text{CON}} + S_{\text{ner}} h_{\text{NR}} \qquad (18)$$

$$S_{we} = \frac{1}{1 + 12 \left[\frac{G(1-x)D}{2} x^{F^{1,25}} \int_{0.01}^{0.01} \left[\frac{q}{e^{4x}} x 10^4 \right]^{-0.5} \right]}$$
(19)

$$q_{val} = \frac{180}{d}q \tag{20}$$

In the case of other material, it is needed to take into account of the heat transfer in the dry part due to the large temperature difference between the wall and vapor. The heat transfer coefficient in the dry part hdry is given as follows:

$$\frac{h_{d\gamma}}{h_{\tau}} = 1 + 153 \left[\left(\frac{GxD}{\mu_{\tau}} \right) / 10^4 \right]^{-1.02} \left[\frac{G^2}{gD\rho_{\tau}(\rho_{\tau} - \rho_{\tau})} \right]^{-0.94}$$
(21)

wher

$$h_{con} = 0.023 \frac{k_L}{D} \left(\frac{G(1-x)D}{\mu_L} \right)^{0.8} P_{\Gamma_L}^{0.4}$$
 (22)

On the other hand, Gungor and Winterton[17] revised their generalized correlation for the vertical flow by taking the flow effects into consideration using the Froud number, Fr, leading to the result that each factor for the vertical tube multiplied by $\Gamma_{hest} = \Gamma r^{(0.1-2\,R)}$ and $S_{hest} = \Gamma r^{1/2}$, respectively, is enough to predict the heat transfer coefficient in the horizontal tubes.

Jung et al. [47] also proposed their correlation using the Stephan and Abdelsalam in place of the Forster and Zuber correlation in Table 1 for the nucleate boiling and the Ditrus-Boelter correlation and determined the two factors F and S as:

$$F = 2.37 \left(0.29 + \frac{1}{V} \right)^{0.35} \tag{23}$$

(24)

$$S = 4048 \ X_{..}^{1.22} \ Bo^{1.13}$$

They reported that their correlation was better in predicting the heat transfer than either the Bennett and Chen and the Gungor and Winterton correlations

5. Correlation of heat transfer for binary mixture

It is generally known that heat transfer coefficient in nucleate

5.1 Nucleate boiling heat transfer

boiling of binary mixture is much lower than that expected from the individual components of the mixture. This is illustrated as an example in Fig.3, where the heat transfer coefficients in a boiling of R12-R113 and R22-R11 mixtures, the temperature difference between boiling and dew points on phase equilibrium at constant pressure and the difference of the mass fraction in liquid and vapor phase, are plotted over the liquid mass fraction of R12 and R22. The heat transfer coefficients are surprisingly much lower than the values, obtained when linearizing between the pure components. The same behavior is found to appear in another mixtures. The mechanism that heat transfer coefficients are sharply reduced is considered: Individual components pass from the liquid to the vapor phase in different proportions, and the faster evaporation of the more volatile component causes an enrichment of the bubble forming boundary layer with the less volatile component, so that the local boiling temperature increases. There are so far two different ways of how to estimate this effect. One is related to the temperature differences between boiling and dew points and the other one to the difference of the mass fraction in liquid and vapor phase. Many correlations proposed so far can be roughly categorized into two from the physical point of view as listed in Table 3. The reduction in the heat transfer coefficient is considered to be attributed to the difference of the mass fraction as pointed out by the most of the correlations in Table 3. Most of the correlations can be roughly given in the following form:

Most of the correlations can be roughly given in the following form
$$\frac{h}{h_{td}} = \frac{1}{C_{mix}}, \quad h_{td} = \frac{1}{\frac{x_1}{h_1} + \frac{x_2}{h_2}}$$
(25)

except for the case of Jungnickel [28] employing $h_{ic} = h_1 x_1 + h_2 x_2$. The constant C_{abc} is always larger than unity whose magnitude largely depends on the mass fraction. The constant C_{abc} means a level of the reduction in heat transfer coefficient due to the effect of mass transfer.

Some experiments for the nucleate boiling are, for reference, summarized in Table 4.

Recently, Incue and Monde[32] and Inoue et al. [57] lawe measured boiling heat transfer coefficients for binary mixtures with a large between the corresponding pure substances(see Fig.1) and pointed out that the Jungnuckel et al. [28] and the Schlunder[29] correlations which stand on the difference of the mass fraction are rather different in advancer from the measured data as shown in Fig.8, while the Thome correlation[26] has a similar trend, but underpredicts the data as shown in Fig.5. Therefore, they have devised the Thome correlation by taking into account of the effect of the heat flux on the hasts of observation of

boiling behavior(ref. see Fig.6 in Inoue and Monde[32]) on a horizontal wire and succeeded in correlating their data well.

Figures 6 and 7 show a comparison of their new correlation[57] and the measured data for mixtures with a large difference in saturation temperatures and with a small difference in saturation temperatures, respectively.

Fujira and Tsutsui(33) separately propose the correlation from the similar concept as estimated from comparison of both equations listed in Table 3. Their correlation can predict their data for methanol-water, methanol-ethanol, ethanol-a-butanol, and methanol-benzene mixtures with a good agreement. It is worth to say that the correlations in which only the values determined from phase equilibrium diagram are used, are easier for calculating the heat transfer coefficient than those using the physical properties depending on the mass fraction.

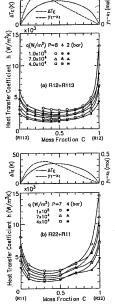


Figure 3 Effect of the mass fraction on the heat transfer coefficients together with relations of DTE and $(y_1 - x_1)$

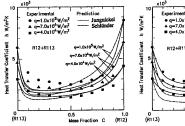


Figure 4 Comparison of the the Jungnuckel et al. [28] and the Schlünder[29] correlations and their data

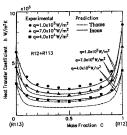
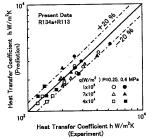
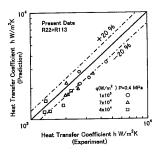


Figure 5 Comparison of the Thome et al. [26] correlation and their data





(a) R134a + R113

(b) R22 + R113

Figure 6 Comparison of the Inoue and Monde correlation with their measured data

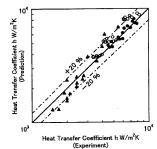


Figure 7 Comparison of the Inoue and Monde correlation with existing

- Data from Jungnickel et al.: R22+R12; C = 0.1 0.9; p = 2, 4, 10 bar; q = 4 x 104 W/m2
- Data from Gorenilo et al.: R22+R114; C = 1 0.9; p/pc = 0.2, 0.4, 0.6, q = 104 W/m2
- Data from Fujita et al.: R11+R113; C = 0.2 0.8; p = 1, 2, 4, 8 bar,
 q = 105 W/m2

5.2 Convective heat transfer in the vertical evaporator tube

The heat transfer in forced convective boiling is also much lower for the mixtures than that for the pure liquid as estimated from the result obtained in the nucleate boiling. Bennett and Chen[15] developed the Chen correlation for the pure liquids by taking into account of mass transfer effects, so as to extend to the aqueous mixtures of ethylene glycol and then gave the following equation for the heat transfer of the mixtures.

$$h_{mix} = h_{CON} F_{mix} + h_{NB} S_{mix}$$
(26)

$$F_{mix} = F f (Pr_L) \left[\frac{\Delta T}{\Delta T_s} \right]_{CON}$$
 (27)

$$f(Pr_L) = \left(\frac{Pr_L + 1}{2}\right)^{0.444}$$
(28)

$$\left[\frac{\Delta T}{\Delta T_s}\right]_{CON} = 1 - \frac{(1 - y_1)q}{\rho_L H_{fs} h_{min} \Delta T_s} \left[\frac{dT_{mis}}{dx}\right]_{p, bmin}$$
(29)

$$h_{mx} = 0.023 \frac{D_{dY.L}}{D} \text{Re}_L^{0.8} [f(Pr_L)F] \left(\frac{\mu_L}{D_{dY.L}}\right)^{0.4}$$
 (30)

$$S = \left(\frac{1}{1 + 2.53 \times 10^{-6} \left[f \left(\text{Pr}_L \right) F \right]^{1.46} \text{Re}_L^{1.17}} \right) / C, \tag{31}$$

$$C = \begin{bmatrix} \frac{1}{1 - \frac{c_{\mu L}(y_1 - x_1)}{H_{\theta_0}} \left[\frac{dT_{\mu L}}{dx} \right] \left(\frac{a_L}{D_{\mu L}} \right)^{1/2}} \end{bmatrix}$$

Recently, Celata et al.[42] show that the Bannett and Chen correlation can applicable for the binary mixtures of R12-R114 and is superior to the type of correlation such as proposed by Mishra et al.[43].

Table 5, for reference, listed some recent experimental studies of heat transfer for mixtures in flow boiling.

5.3 Convective heat transfer in the horizontal evaporator tube

A significant reduction in the heat transfer coefficient appears for binary mixtures attributable to mass transfer effects.

Jung et al.[49] proposed their following correlation which is derived based on the Chen concept but different for the Bannett and Chen correlation, and reported that it can predict their data for the mixture of R22-R114 with a mean deviation of 9.6 %.

$$h_{\text{mix}} = h_{\text{CON}} F_{\text{mix}} + h_{\text{NR} \text{mix}} S_{\text{mix}}$$
(32)

where hem is the Unal correlation, hoon is also the Dittus-Boelter equation, Smix = S/Ctin, Ctin is the constant in the Unal correlation, S is given by equation (24) and $F_{mix} = F C_{max}$, F is given by equation (23) and Cm is proposed as:

$$C_{me} = (1 - 0.35|y_1 - x_1|^{1.56})$$
 (33)

Jung et al.[49] recommend to use the Stephan and Abdelsalam correlation for the heat transfer coefficients for the individual components needed when calculating how min. They concluded that their correlation is the best one in correlating their data among the other correlations[15][17][46].

It may be noticed finally that most of the correlations predicting heat transfer for the annular flow derived based on the Chen criterion in which the Dittus-Boelter equation is commonly employed for heat transfer in forced convective flow, while the correlation for the nucleate heat transfer seems to be arbitrary chosen from any correlation in Table 3. The reduction in the heat transfer for the mixture due to the mass transfer effects seems to be treated with from the two different points of view not only in nucleate boiling but also in convective boiling; one is focused on the temperature difference, and the other one the concentration difference.

6.Concluding Remarks

The interest in flow boiling heat transfer in mixtures has been increased. Further experimental works with different mixtures must be done on a wide range of compositions to verify the known correlations and to improve them. It would be hoped to make it clear how the mass transfer influences the heat transfer, especially in the field of nucleate boiling.

Nomenclature

- a :Thermal diffusivity
- B_a : Boiling number (= $q/\rho_L H_{fg}$)
- C :Mass fraction of a more volatile component in a system
- Cp : Specific heat at constant pressure
- d. :Diameter of bubble departure
- D :Tube diameter of Mass diffusivity
- :Pressure factor
- f. :Surface factor
- f. :Nucleation factor
- :Enhancement factor
- Fr :Froude number $\{=G^2/(\rho_L^2gD)\}$
- g :Acceleration of gravity
- G :Mass flux
- h :Heat transfer coefficient
- h_{kl} : Ideal heat transfer { = 1/(x₁/h₁ + 1/(x₂/h₂) }
- His: Latent heat in vaporization
- Representative length of heated surface
- M :Molecular weight
 - P :Pressure
- P. :Critical pressure
- Pr ·Prandtl number
- q :Heat flux
- R. :Smoothness degree of heated surface Re_L :Reynolds number (= $G(1-x)D/u_r$)
- S :Suppression factor
 - T :Temperature

T. :Critical temperature

T_s:Saturation temperature

 T_{wid} : Ideal wall temperature { = $(T_{bulk} + \Delta T_{id})$ }

 ΔT_B :Temperature difference between dew and bubble point

{ =(T_{local,max} - T_{bulk})

 ΔT_{id} : ideal wall superheat { = ($\Delta T_{int,1} x_1 + \Delta T_{int,2} x_2$)}

ΔT_{set} :Wall superheat

x, y:Quality or Mole fractions of liquid and vapor

 $\mathbf{x}_{\mathbf{m}}, \mathbf{y}_{\mathbf{m}}$: Mass fraction of liquid and vapor

 X_{tt} :Lockhart-Martinelli parameter

Greek symbol

α :Volume expansion coefficient

B :Mass transfer coefficient

κ :Thermal conductivity
λ :Thermal conductivity

μ :Viscosity

v :Kinematic viscosity

ρ :Density

σ :Surface tension

σ

Suffix

:More volatile component

2 :Less volatile component

bulk :Bulk

CON : Convective flow

dry:Dry surface

id :Ideal

L :Liquid

local :Local

mix:Mixture NB :Nucleate

Two phase flow

v :Vapor

wet:Wet surface

Table 1 Correlations for pool boiling in pure liquid

Author	Correlation
Kutateladze[1]	$N_{n,b} = 7.0 \times 10^{-4} R_{n,b}^{0.7} P_{r,L}^{0.35} f_{p}^{0.7}$
	$N_{n,b} = \frac{hb}{\lambda_L} R_{n,b} = \frac{qb}{H_{\ell,n} \rho_{\tau, \nu_L}}$
	$f_{P} = \frac{P}{\sqrt{\sigma g(\rho_{\nu} - \rho_{*})}} , b = \sqrt{\frac{\sigma}{g(\rho_{\nu} - \rho_{*})}}$
McNelly[2]	$\tilde{N}_{u} = 0.225 \left(\frac{qd}{H_{f_{\#} \mu_{L}}} \right)^{0.58} \left(\frac{pd}{\sigma} \right)^{0.31} \left(\frac{\rho_{L} - \rho_{v}}{\rho_{v}} \right)^{0.33} P_{r_{L}^{0.58}}$
Rohsenow[3]	$N_{ub} = \frac{P_{rL}^{-0.7}}{C_{sf}} \left(R_{eb} \frac{\rho_{v}}{\rho_{L}} \right)^{0.887}, C_{sf} = 0.00225 \sim 0.0154$
Forster and Greif[4]	N_{*} = 0.0346 $\sqrt{R_{*}}$ $\left(\frac{\sigma}{\rho_{\nu}\nu_{L}}\frac{b^{*}}{\nu_{L}}\right)^{0.128} P_{rL}^{0.6}$
	$\mathtt{N_{ub}^*} = \frac{\mathtt{hb}}{\lambda_\mathtt{L}} \ , \ \mathtt{R_{ub}^*} = \frac{\mathtt{qb^*}}{\mathtt{H_{te}} \rho_\mathtt{L} \nu_\mathtt{L}} \ , \ \mathtt{b^*} = \frac{\rho_\mathtt{L} \mathtt{C}_\mathtt{FL} \mathtt{T_u}}{\rho_\mathtt{v} \mathtt{H_{te}}} \frac{\sigma}{\rho_\mathtt{v} \mathtt{H_{te}}}$
Zuber[5]	$N_{s} = \text{const.} \left[\frac{g l^{3}}{\nu_{L} a_{L}} \left(\alpha \Delta T_{**} + \frac{nb}{\Lambda} \frac{\pi}{6} d_{o}^{2} \frac{\rho_{L} - \rho_{v}}{\rho_{L}} \right) \right]^{1/3}$
	$d_0 = 0.021 \theta \left[\frac{\sigma}{g(\rho_L - \rho_V)} \right]^{1/2}$
Borishanskii[6]	$N_{ub} = 8.7 \times 10^{-4} R_{e_0}^{0.7} P_r_{L}^{0.7} f_{P_0}^{0.7}$
Nishikawa and Fujita[7]	Laminer Flow: $Y=6.24(f_c f_P X)^{2/3}$
	Turburent Flow: $Y=0.661^{-2/5}(f_cf_pX)^{4/5}$
	$Y = \frac{hl}{\lambda_1}$, $X = \left\{ \left(\frac{1}{W^2 N} \right) \frac{C_{PL} \rho_L^2 g}{2 \sigma_1 H_{PL} \rho_L} \right\}^{1/2} ql^{3/2}$
	N=1.976 W , M=900 m ⁻¹ , $f_p = \left(\frac{P}{P_e}\right)^{0.7} \left[1+3\left(\frac{P}{P_e}\right)^3\right]$
	Low Heat Flux : $h_L = 1.140 \frac{P_c^{1/3}}{\mathfrak{M}^{1/8}T_c^{5/6}} f_{PL} (f_c q)^{3/3}$
	$f_{PL} = \left(\frac{P}{P_c}\right)^{0.19} \left[1 + 2\left(\frac{P}{P_c}\right)^2 + \left(\frac{P}{P_c}\right)^6\right]$
	High Heat Flux : $h_u=492\frac{P_c^{1/s}}{2\overline{N}^{1/s} \circ T_c^{s/10}} f_{PM}(f_cq)^{4/s}$
	$f_{PH} = \left(\frac{P}{P_{c}}\right)^{c. \ 2a} \left[1 + 2\left(\frac{P}{P_{c}}\right)^{a} + 8\left(\frac{P}{P_{c}}\right)^{a}\right]$

Table 1 Correlations for pool boiling in pure liquid(Continued)

Author

Correlation

Stephan and Abdelsalam[8]

Water:
$$N_{\kappa} = 0.246 \times 10^{7} \left(\frac{qd}{\lambda_{c}T_{***}}\right)^{0.678} \left(\frac{C_{\sigma}d^{2}T_{***}}{a_{c}^{2}}\right)^{1.86} \times \left(\frac{H_{c}d^{2}}{a_{c}^{3}}\right)^{-1.68} \left(\frac{\rho_{L}-\rho_{*}}{\rho_{L}}\right)^{6.23}$$

$$10^{-4} < \frac{P}{P_{*}} < 0.886 , \theta = 45$$

$$\begin{split} \text{Hydrocarbons}: N_s = & 0.0546 \bigg\{ \frac{qd}{\lambda L T_{s.s.}} \bigg\}^{0.57} \bigg(\frac{H_{14}d^3}{a_L^3} \bigg)^{0.2485} \\ & \times \bigg(\frac{\rho_*}{\rho_L} \bigg)^{0.3335} \bigg(\frac{\rho_* - \rho_*}{\rho_L} \bigg)^{-4.33} \\ & 5.7 \times 10^{-8} < \frac{\rho_*}{P_*} < 0.9 \ , \ \theta = 35 \ . \end{split}$$

Cryogenic fluids :
$$N_* = 4.82 \left(\frac{qd}{\lambda_L T_{t+1}}\right)^{0.884} \left(\frac{C_{\nu} d^2 T_{t+1}}{a_L^2}\right)^{0.874} \left(\frac{H_{t} d^2}{a_L^2}\right)^{-0.826} \times \left(\frac{\rho}{\rho_L}\right)^{0.287} \left(\frac{\rho v C_{\nu \nu} \lambda_{\nu}}{\rho_L C_{\nu L} \lambda_L}\right)^{0.117} \times 10^{-8} < \frac{\rho}{\rho} < 0.97$$
, $\theta = 1$

Refrigerants:
$$N_s = 207 \left(\frac{qd}{\lambda_L T_{sec}} \right)^{0.748} \left(\frac{\rho_T}{\rho_L} \right)^{0.881} P_{rL}^{0.833}$$

 $3 \times 10^{-3} < \frac{P}{P_s} < 0.78$, $\theta = 35^{\circ}$, $d = 0.16 \theta \sqrt{\frac{2\sigma}{K(\rho_L - \rho_A)}}$

Cooper[9]

Forster and Zuber[10]
$$\frac{q}{\rho_* H_{fg}} \left(\frac{\pi}{\alpha_L}\right)^{1/2} \left(\frac{\rho_L r^{*3}}{2\sigma}\right)^{1/4} = 0.0015 \left[\frac{\rho_L}{\mu_L} \left(\frac{\Delta T_{int} \lambda_L}{\rho_* H_{fg}}\right)^2 \frac{\pi}{\alpha_L}\right]^{2/3} \left(\frac{C_p \, \mu}{k_B}\right)^{1/2} \\ r^* = \frac{2 \, \sigma \, R T_{int}^2}{H_{fg} \text{Np}_L \Delta T_{int}}$$

Table 2 Heat transfer correlation for forced convective boiling

Authors	Correlations	Condition	
Schrock and Grossman[11]	$\frac{h_{TP}}{h_{LO'}} = 0.739 \left[Bo \times 10^4 + 1.5 \left(\frac{1}{X_{tt}} \right)^{2/3} \right]$	Water, p = 3 - 35 ata Upward flow	
Wright[12]	$\frac{h_{TP}}{h_{LO'}} = 0.67 \left[Bo \times 10^4 + 3.5 \left(\frac{1}{X_{tt}} \right)^{2/3} \right]$	Water, p = 1 - 5 ata Downward flow	
Pujol and Stenning[13]	$\frac{h_{TP}}{h_{LO}} = 0.90 \left[Bo \times 10^4 + 4.45 \left(\frac{1}{X_{tt}} \right)^{0.37} \right]$ $\frac{h_{TP}}{h_{LO}} = 0.53 \left[Bo \times 10^4 + 7.75 \left(\frac{1}{X_{tt}} \right)^{0.37} \right]$	R113 Upward flow R113 Downward flow	

Table 3 Correlations for pool boiling in mixture

Authors	Correlations
Van Stralen[19]	$h = \frac{1}{2} \rho_2 H_{tz} k_i \sqrt{\tau}$
	$k_1\!=\!\!\sqrt{\frac{12}{\pi}}\;\frac{\sqrt{a_L}}{\frac{\rho_{\text{\tiny S}}\left(\frac{1}{C_{PL}}\!-\!C\!\left(\frac{y_{a_L}\!-\!x_{a_L}}{x_{a_L}}\right)\!\sqrt{\frac{s_L}{D}}\left(\frac{dT}{dx_{a_L}}\right)_{x=c}\right.}$
Stephan and Körner[20]	$\frac{h}{h_{1d}} = \frac{1}{1 + A_0 (0.88 + 0.12P[bar]) (y - x) \Delta T_{1d}}$
Tolubinsky and Ostrobsky[21]	$N_{u} = \left[\frac{q}{\rho_{v} H_{rg} \{x_{n1} (fd_{0})_{1} + x_{n2} (fd_{0})_{2}\}}\right]^{0.7}$
	$\times \left\{ 1 - \frac{(y_{n1} - x_{n1})^2}{y_{n1}(1 - x_{n1})} \right\}^{1 \cdot 0} P_{rL}^{2 \cdot 2}$
Calus and Rice[22]	$N_{n} = f_{n} \left[\frac{P_{n}}{1 + (y_{n,1} - x_{n,1}) \sqrt{\frac{a_{n}}{D}}} \right]^{0.7} \left[\frac{T_{n}}{T_{n,m}} \right]^{4} f_{p} \circ .7$
	$f_{p} = \frac{P}{\sqrt{g \sigma (\rho_{L} - \rho_{\tau})}}$
Calus and Leonidopoulos[23]	$\Delta T_{\text{est}} = (x_{\text{mi}} \Delta T_{\text{sat}, 1} + x_{\text{m2}} \Delta T_{\text{est}, 2})$
	$\times \left\{ 1^{-\left(y_{m1}-x_{m1}\right)} \sqrt{\frac{\underline{a_{t_{c}}}}{D}} \left(\frac{\underline{C_{P}}}{H_{fg}}\right) \left(\frac{d\overline{I}}{dx_{m1}}\right) \right\}$
Happel and Stephan[24]	$\frac{h}{h_{id}} = 1 - k_{i} (y_{1} - x_{i})^{-1}$
Stephan and Preusser[25]	$\frac{hd_o}{\lambda_L} = 0.10 \left(\frac{qd_o}{\lambda_L T_{eat}} \right)^{0.874} \left(\frac{\rho_v}{\rho_L} \right)^{0.188}$
	$\times \left(\frac{\mathbb{E}_{\text{rg}} \mathbf{d}_{0}^{2}}{\mathbf{a}_{L}^{2}}\right)^{0.37 \text{t}} \left(\frac{\mathbf{a}_{L}^{2}}{\sigma \mathbf{d}_{0}}\right)^{0.380} \left(\frac{\mu_{L} C_{P}}{\lambda_{L}}\right)^{-0.182}$
	$\times \left\{ 1 + \left \sum_{i} (y_i - x_i) \left(\frac{\partial y_i}{\partial x_i} \right)_{x_{d, p}} \right \right\}^{-0.0733}$
	$d_0 = 0.0146 \theta \left\{ \frac{2\sigma}{g(\rho_L - \rho_T)} \right\}^{0.8}$

Table 3 Correlations for pool boiling in mixture(Continue)

Author	Correlations			
Thome[26]	$\frac{h}{h_{1d}} = N_{\pi\pi}^{7/5}$			
	$N_{nn} = \left\{1 - (y_1 - x_1) \sqrt{\frac{a_L}{D}} \left(\frac{C_P}{H_{f,g}}\right) \left(\frac{dT}{dx}\right)\right\}^{-1}$			
Thome[27]	$\frac{h}{h_{1d}} = \frac{1}{1 + \frac{\Delta T_x}{\Delta T_{1d}}}$			
Junguickel et al.[28]	$\frac{h}{h_{1d}} = \frac{1}{1 + k_0 \frac{\rho_v}{\rho_L} y_1 - x_1 q^{(0.48+0.1x1)}}$			
Schländer[29]	$\frac{h}{h_{1d}} = \frac{1}{1 + \frac{h_{1d}}{q} (T_{n2} - T_{n1})(y_1 - x_1) \left\{ 1 - exp \left(-\frac{B_0 q}{\rho_L \beta_L H_{1c}} \right) \right\}}$			
Bier et al.[30]	$\frac{h}{h_{t,d}} = 1 - \left\{ A + B \left(\frac{P}{P_c} \right) + \frac{C}{1 - \left(\frac{P}{P_c} \right)} \right\} (y_1 - x_1)^{n}$			
Ünal[31]	$\frac{h}{h_{1:d}} = \frac{1}{C_{uu}}$ $C_{uu} = \{1 + (b_u + b_u)(1 + b_u)\}(1 + b_u)$			
	$b_{a} = (1 - X_{t}) ln \left(\frac{1.01 - X_{t}}{1.01 - Y_{t}} \right) + X_{t} ln \left(\frac{X_{t}}{Y_{t}} \right) + Y_{t} - X_{t} ^{1.8}$			
	b ₂ =0 for X ₁ ≥0.01			
	$b_0 = (Y_1/X_1)^{0.1} - 1$ for $X_1 < 0.01$			
	b,=152(p/p.=v.) * *			
	$b_0 = 0.92 Y_1 - X_1 ^{\alpha \cos(p/p_{maxe})^{\alpha}}$ se			
	$X_i/Y_i=1 \text{for} X_i=Y_i=0$			
Inoue and Monde[32]	$\frac{h}{h_{id}} = \frac{1}{\Delta T_E}, k = 0.45 \times 10^{-5} q + 0.25$			
	$\frac{\Delta T_{E}}{1 + k \Delta T_{id}} , k = 0.43 \times 10^{-4} q + 0.25$ $10^{4} \le q [W/m^{2}] \le 10^{5}$			
Fujita and Tsutsui[33]	$\frac{h}{h_{1d}} = \frac{1}{1 + k \frac{\Delta T_E}{\Delta T_{1d}}}, k = 1 - 0.8 exp(-10^{-5}q)$			
Inoue et al.[57]	$\frac{h}{h_{1d}} = \frac{1}{1 + k \frac{\Delta T_E}{\Delta T_{1d}}}, k = 1 - 0.75 \text{ exp}[-0.75 \text{x} 10 - 5 \text{ q}]$ $10^4 \le q [\text{W/m}^2] \le 10^5$			

Table 4 Some experimental studis of pool boiling heat transfer in mixtures

Authors	Heated Surface	Test Substances Press	ures(MPa)
Bonilla et al.	Horizontal Plate	Acetone+Ethanol	0.1
[34]		Ethanol+Butanol	
		Acetone+Water	191
		the second of th	
		Ethanol+Water	
		Isobutanol+Water	
Van Wijk et al.	Horizontal Wire	Ethanoi+Toluene	0.1
[35]		Isopropanol+Cyclo Hexane	
		Acetone+Water	
		Butanol+Water	
		Pentanol+Water	
		rentanoi+water	
Van Stralen[36]		Acetone+Butanol	_ =
Sternling et al.		Ethanol+Benzene	
[37]		Isopropanol+Ethylene Glycol	
		Isopropanol+Cyclo Hexane	
		Methanol+Ethylene Glycol	
		Methanoi+Eurylene Grycol	
Grigor'ev et al. I	Horizontal Tube	Ethanol+Butanol	0.1
[38]		Acetone+Butanol	
		Ethanol+Water	
	Iorizontal Wire	Methyl-Ethyl-Ketone+Water	0.1
[19]		Methanol+Water	
Tolubinsky et al. I	Iorizontal Tube	Ethanol+Benzene	0.1
[39]		Ethanol+Water	0.1
(1		Editator Water	
Körner		Acetone+Butanol	
[40]		n-Heptane+Methyl-Cyclo-Hexane	
		Methyl Ethyl Ketone+Toluene	
		Acetone+Water	
		Propanol+Water	
Stephan et al.		Acetone+Butanol	0.1-1.0
[20]		Heptane+Methyl-Cyclo-Hexane	
		Methanol+Benzol	
Tolubinsky et al. V	Vertical Tube	Ethanol+Water	0.1
[21]	Oldon 1000		0.1
[41]		Methanoi+Water	
		Ethanol+Butanol	
		Ethanol+Benzene	

Table 4 Some experimental studis of pool boiling heat transfer in mixtures(Continued)

Authors	Heated Surface	Test Substances	Pressures(MPa)
Stephan et al. [41]	Horizontal Tube	Ethanol+Cyclo-Hexane Methanol+Benzol	0.1-0.3
Calus et al. [23]	Horizontal Wire	Isopropanol+Water Acetone+Water Ethylene-Glycol+Water	0.1
Calus et al.	Horizontal Wire	Propanol+Water	0.1
Happel et al. [24]	Horizontal Tube	Benzene+Toluene Ethanol+Benzene Isobutanol+Water	0.05-0.2
Stephan et al. [25]	Horizontal Tube	Acetone+Methanol Acetone+Water	0.1
Jungnickel et al. [28]	Horizontal Plate	R22+R12 R12+R13	0.1-2.0
Bier et al. [30]	Horizontal	TubeR12+SF6 R22+SF6 R13B1+SF6	0.56-0.97 (Reduced Pressure
Fujita et al. [33]	Horizontal Plate	Methanol+Water Ethanol+Water Methanol+Ethanol Ethanol+n-Butanol Methanol+Benzene	0.1

Table 5 Some recent Experimental studies of heat transfer for mixtures in flow boiling

Authors	year	mixture			
Bennett and Chen[15]	1980	water/ethylene	t	٧	e
		glycol			
Toral et al.[44]	1982	ethanol/	t	v	е ,
		cychlohexan			
Nakanishi et al.[45]	1986	R12/R13	t	h	e
Ross et al.[46]	1987	R152a/R13B1	t	h	e .
Hihara et al.[46]	1989	R12/R22	t	h	e
		R114/R22			
Jung et al.[47]	1989	R22/R113	t	h	e
Jung et al.[48]	1989	R12/R152a	t	h	e
Jung et al.[49]	1989	refrigerants	t	h	ė
Murata and Hashizume[50]	1990	R11/R114	t	h	e
Takamatsu et al.[51]	1990	R22/R114	t*	h	w
Fujita et al. [52]	1991	R11/R113	t	v	e
Yoshida et al.[18]	1991	R22/R114	t	h	e
Sami and Duong[53]	1992	R22/R114	a*	h	w
Sami and Schnotale[54]	1992	R22/R114	ť	h	w
		R22/R152a			
Celata et al.[42]	1993	R12/R114	t	v	e
Takamatsu et al.[55]	1993	R22/R114	t	h	w

t: tube, a: annulus,

*:enhanced surface

h: horizontal, v: vertical

e: electrically heated, w: water heated

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