<table>
<thead>
<tr>
<th>項目</th>
<th>内容</th>
</tr>
</thead>
</table>
| タイトル               | ガス-液体の気液混合の前線流れる気液混合装置における気液の移動特性

著者                  | 坂井 竹彦；伊地知 真也；上村 文雄；米原 信裕          |
|                       |                                                      |
| 引用                  | ISIJ International, 42(1), 112-114                 |
|                       | https://doi.org/10.2355/isijinternational.42.112 |

日時                  | 2002-01-15                                              |

権利                  | 著作権は日本鉄鋼協会にある |

ファイル               | 16-T. Kumagai-4.01.pdf                                   |

Hokkaido University Collection of Scholarly and Academic Papers: HUSCAP
Gas–Liquid Mass Transfer in Simulated Turbulent Wake Flow

Takehiko KUMAGAI, Manabu IGUCHI, Tomomasa UEMURA1) and Noriyoshi YONEHARA1)

Graduate School of Engineering, Hokkaido University, North 13, West 8, Kita-ku, Sapporo 060-8628 Japan.
1) Faculty of Engineering, Kansai University, 3 Chou-me, 3 Ban-chi, Yamate-chou, Suita-shi, Osaka-fu 564-8680 Japan.

(Received on July 26, 2001; accepted in final form on September 29, 2001)

1. Introduction

In materials engineering, gas injection processes are frequently adopted to remove impurities such as oxygen and carbon.1–3) There are two types of gas dispersion patterns formed above the nozzle; bubbling and jetting. The former is realized when the gas velocity at the nozzle exit, \( v_n \), is lower than the speed of sound, \( c \), while the latter is realized for \( v_n^c \) and a gas column is formed on the nozzle. This column disintegrates into many bubbles with different diameters above a certain distance from the nozzle exit. Many bubbles are therefore generated in the molten metal bath regardless of the bubbling and jetting. The removal of impurities is closely associated with the dissolving rate of the gas into the bath. However, under such a highly turbulent condition it is difficult to evaluate the mass transfer coefficient between the bubbles and molten metal because the precise evaluation of the interfacial area is actually impossible.4–6)

As the gas flow rate increases, mass transfer at the gas–liquid interface increases. However, it is not clear whether the mass flux at the gas–liquid interface is attributed to the enhancement of the interfacial area \( A \) or to the enhancement of the mass transfer coefficient \( k \). Accordingly, previous researchers7–13) introduced the volumetric mass transfer coefficient defined as \( \frac{kA}{V} \), where \( V \) is the bath volume.

In a series of studies14–18) on gas injection, the authors have tried to elucidate the contributions of the interfacial area and the mass transfer coefficient to the volumetric mass transfer coefficient. The mass transfer coefficient at a gas–liquid interface exposed to two types of liquid jets shown in Figs. 1(a) and 1(b) were measured.18) CO2 gas was supplied with a syringe so that a gas–liquid interface was formed at the lance exit.

As time elapses, the CO2 gas dissolves into the water, and, as a result, the interface rises upward if the CO2 gas is not fed. The CO2 gas was therefore supplied continuously to keep the gas–liquid interface at the lance exit anytime. The gas–liquid interfacial area is controlled by changing the inner pipe diameter. Three pipes of different diameters were prepared. The inner cross-sectional areas are 2.50 \( \times \) 10\(^{-4} \) m\(^2\), 0.498 \( \times \) 10\(^{-4} \) m\(^2\) and 0.249 \( \times \) 10\(^{-4} \) m\(^2\).

The velocity of water flow at the exit plane of the top lance was measured with a two-channel laser Doppler velocimeter. The mass transfer coefficient \( k \) was calculated from the relation\(^{19–21} \):

\[
m = k \cdot \Delta C \cdot (t \cdot A)
\]

where \( m \) is the dissolved mass of CO2, \( \Delta C \) is the concentration difference, \( t \) is time and \( A \) is the gas–liquid interfacial area. The mass flux was determined by measuring the volume of the supplied CO2 gas.

2. Experimental Apparatus and Procedure

Figure 2 shows a schematic of the experimental apparatus. Distilled water was circulated with a pump. The water flow rate was controlled with a flow control valve. A top lance made of a glass pipe was enclosed with a transparent glass pipe with a larger diameter. The water passed between the two pipes. Glass wool was used as a flow straightener. CO2 gas was supplied with a syringe so that a gas–liquid interface was formed at the lance exit.

As time elapses, the CO2 gas dissolves into the water, and, as a result, the interface rises upward if the CO2 gas is not fed. The CO2 gas was therefore supplied continuously to keep the gas–liquid interface at the lance exit anytime. The gas–liquid interfacial area is controlled by changing the inner pipe diameter. Three pipes of different diameters were prepared. The inner cross-sectional areas are 2.50 \( \times \) 10\(^{-4} \) m\(^2\), 0.498 \( \times \) 10\(^{-4} \) m\(^2\) and 0.249 \( \times \) 10\(^{-4} \) m\(^2\).

The velocity of water flow at the exit plane of the top lance was measured with a two-channel laser Doppler velocimeter. The mass transfer coefficient \( k \) was calculated from the relation\(^{19–21} \):

\[
m = k \cdot \Delta C \cdot (t \cdot A)
\]

where \( m \) is the dissolved mass of CO2, \( \Delta C \) is the concentration difference, \( t \) is time and \( A \) is the gas–liquid interfacial area. The mass flux was determined by measuring the volume of the supplied CO2 gas.
3. Experimental Results and Discussion

3.1. Brief Summary of Previous Experimental Results of Mass Transfer Coefficient

According to the previous study of the present authors, the mass transfer coefficient in the flow systems which are shown in Figs. 1(a) and 1(b) can be correlated in terms of the Sherwood number similitude.

\[ \text{Sh} = 0.664 \text{Sc}^{1/3} \text{Re}^{0.5} \text{Tu}^{0.1} \]  

\[ \text{Re} = \bar{u}d/D \]  

\[ \text{Sc} = v/D \]  

\[ \text{Tu} = u_{rms}/\bar{u} \]

where \( \text{Re} \) is the Reynolds number, \( \text{Sc} \) is the Schmidt number, \( \bar{u} \) is the mean flow velocity, \( v \) is the kinematic viscosity of liquid, \( d \) is the lance diameter, and \( D \) is the diffusion coefficient, \( \text{Tu} \) is the turbulence intensity of the parallel and cross flows and \( u'_{rms} \) is the root-mean-square value of the turbulence component. The measured values of the mass flux were approximated by Eq. (2) within a scatter of \( \pm 30\% \).

It is interesting to note that Eq. (2) approaches an empirical relation for the mass transfer coefficient at a liquid–solid interface as the Reynolds number increases, although the evidence is not shown here. This fact means that the water is fully contaminated during the measurement, and, hence, the gas–liquid interface is dirty. Considering these circumstances, experiments were carried out also in this study after much time had passed from the start of water circulation. Whether the water is contaminated or not can be judged by measuring the rising velocity of a single bubble in a still water bath.

3.2. Correlation of Mass Transfer Coefficient

Figure 3 shows the relationship between the mass flux and the Reynolds number \( \text{Re} \). The gas–liquid interfacial area is used as a parameter. The mass flux increased with an increase in the Reynolds number but decreased with an increase in the interfacial area. The mean velocity around the gas–liquid interface is shown in Fig. 4. The radial distribution of the mean water velocity is nearly uniform outside the gas–liquid interface. The turbulence intensity was 0.26.

The data on the mass flux for \( A = 2.508 \times 10^{-4} \text{ m}^2 \) are replotted in Fig. 5 together with the calculated values from Eq. (2). The measured values are smaller than the calculated values at every Reynolds number. Therefore, the coefficient in Eq. (2), 0.664, was replaced by 0.500.

\[ \text{Sh} = 0.500 \text{Sc}^{1/3} \text{Re}^{0.5} \text{Tu}^{0.1} \]

All the measured values of the mass flux obtained in this
study are compared with the calculated values from Eq. (7) in Fig. 6. The measured values can be predicted by Eq. (7) within a scatter of ±30%.

4. Conclusions

Model experiments were carried out to determine the mass transfer coefficient at the rear part of a bubble rising in a highly turbulent water flow. The gas–liquid interface was contaminated by surfactant. The mass transfer coefficient was nearly approximated by the empirical relation proposed in this study, Eq. (7).

Nomenclature

\[ A: \text{Gas–liquid interfacial area (m}^2\text{)} \]
\[ c: \text{Speed of sound (m/s)} \]
\[ \Delta C: \text{Concentration difference (g/m}^3\text{)} \]
\[ D: \text{Diffusion coefficient (m}^2\text{/s)} \]
\[ d: \text{Lance diameter (m)} \]
\[ k: \text{Mass transfer coefficient (m/s)} \]
\[ m: \text{Dissolved mass of CO}_2\text{ gas (g)} \]
\[ Sh: \text{Sherwood number (-)} \]
\[ Re: \text{Reynolds number (-)} \]
\[ Sc: \text{Schmidt number (-)} \]
\[ t: \text{Time (s)} \]
\[ Tu: \text{Turbulence intensity (-)} \]
\[ \bar{u}: \text{Mean flow velocity (m/s)} \]
\[ u_{rms}': \text{Root-mean-square value of turbulence component (m/s)} \]
\[ v_n: \text{Gas velocity at nozzle exit (m/s)} \]
\[ v: \text{Kinematic viscosity of liquid (m}^2\text{/s)} \]

REFERENCES