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# Mass-Transfer Characteristics of a Double-Y-Type Microchannel Device

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## 1. Introduction

Microfluidic devices have been extensively applied to high-throughput chemical analyses or reactions using hazardous substances (van den Berg et al., 2000). These systems are characterized by a large area per volume and a well-defined laminar flow. As a result of their high potential for the precise manipulation of chemical reactions with respect to controlled temperature, contacting time, and molecular orientation at the interface, researches on microfluidic devices for the production of highly valuable substances are increasing in importance (Ajmera et al., 2001, Maruyama et al., 2003, Löb et al., 2004). The production system is often referred to as a desktop chemical plant and the keys to realizing the plant are understanding and successfully manipulating the flow in microchannels (Hetsroni et al., 2005).

A unique feature of the microchannels is the formation of liquid layers when multiple liquids are brought into one channel. Reactions and/or mass-transfers occur in the vicinity of the interface. Liquid-liquid extractions (Brody and Yager, 1997, Tokeshi et al., 2000) are reported for systems of aqueous and organic solutions. Even mutually miscible liquids form the layers because of the laminar-flow nature (Yamaguchi et al., 2004) and a mass-transfer occurs between the liquids under well-controlled conditions. For the purpose of chemical processing, the stacked liquid layers should be separated in each layer for further treatment. However, flow splitting in microchannel devices is rarely found in published papers. In reported works for the analytical application of microchannels, the solute's concentration profile was directly measured within the channel using a laser-based refractive index detector (Costin and Synovec, 2002) or a microscope for measuring fluorescence emission (Costin et al., 2004). A strategy for flow splitting has not yet been developed for the application of microchannel devices to analytical chemistry.

In the present study, two aqueous layers were brought into contact and separated into two streams after a mass-transfer between the layers. The amount of solute transferred was obtained with a simple analysis of liquid samples taken at the channel outlet. The mass-transfer characteristics were examined through the measurement of the diffusion coefficients for solutes. Even under carefully controlled liquid feed conditions, equal flow splitting was difficult to obtain. Proposed is a method to choose solute concentration data for equal flow splitting using the ratio of the flow rate of one liquid to the other. The mass-transfer of benzoic acid was successfully analyzed with a conventional penetration model with an additional parameter. Furthermore, the diffusion coefficients of sucrose, glycine,

tryptophan, and urea were measured using a basic equation obtained with benzoic acid. The values are in accordance with the literature or the correlated values. Since the same parameter is applied for obtaining the diffusion coefficients of the four different solutes, the additional parameter should characterize the flow behavior in the microchannel. The proposed method is also useful for determining the unknown diffusivities of solutes. Furthermore, to investigate the flow characteristics, the value of an additional parameter is determined in various channels having different lengths, depths and knife-edge angles. A Comparison of this parameter helps us to understand the effect of flow on mass-transfer.

## 2. Determination of diffusivity

### 2.1 Microchannel module and materials

Benzoic acid, sucrose, tryptophan, urea, and glycine were selected as transferring solutes. The aqueous solutions were prepared as feedstock, and the concentration of each of these was set at  $1.19 \times 10^{-3} \text{ kmol/m}^3$ ,  $1.46 \times 10^{-2} \text{ kmol/m}^3$ ,  $1.19 \times 10^{-3} \text{ kmol/m}^3$ ,  $8.7 \times 10^{-2} \text{ kmol/m}^3$ , and  $6.7 \times 10^{-2} \text{ kmol/m}^3$  respectively. The solutions were filtered prior to use to remove the suspended solid to prevent the clogging of the channel.

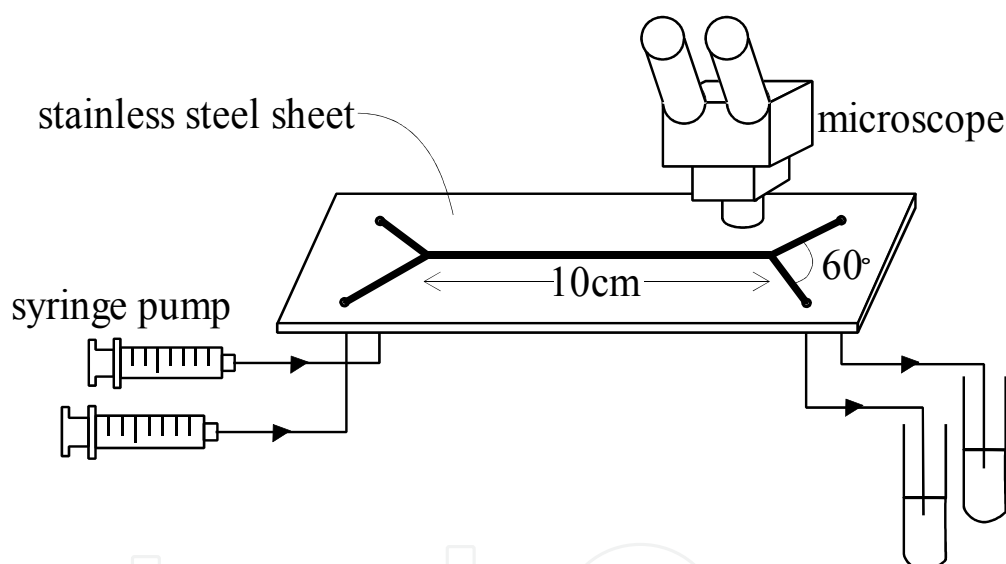


Fig. 1. Schematic representation of microchannel

The microchannel used was fabricated on a stainless steel sheet as shown in Fig. 1. The thickness of the sheet was  $200 \mu\text{m}$ . The channel width was  $200 \mu\text{m}$  for both the branches and the main portion. As shown in Fig. 2, the stainless steel sheet was set in between gaskets of a polyethylene film ( $100 \mu\text{m}$  thick) to form the flow channels of rectangular cross section. An aqueous solution of the solute and deionized water were supplied with syringe pumps at exactly the same flow rate. These two liquids were introduced to both inlets and brought into contact at the main portion of the channel. The two liquids split at a knife-edge, and each liquid flowed into a branch. The liquids were taken out from the end of the branch. The side where the feed solution flowed is called the feed-side, and the other side where the deionized water flowed is the recovery-side. Liquid flow at the main portion of the channel was monitored with a microscope to confirm that there was no flow disturbance. Figure 3 shows a photograph of the flow splitting point, which was taken at the examination of the

flow profile with the supply of a Cibacron blue solution and water in each inlet. The blue solution formed a layer and a stable interface is observed at the center of the channel. The two solutions were then split with a knife-edge at the end of the main portion of the channel.

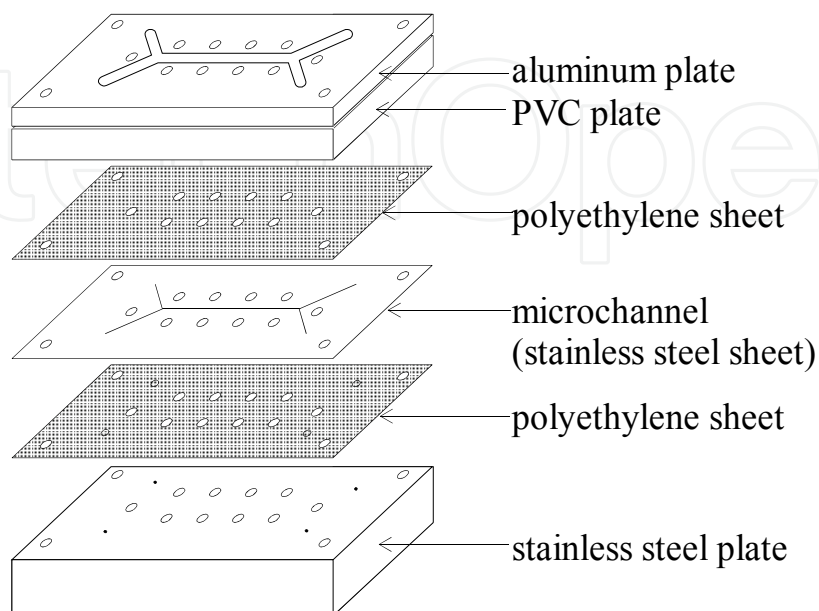


Fig. 2. Expanded view of microchannel module

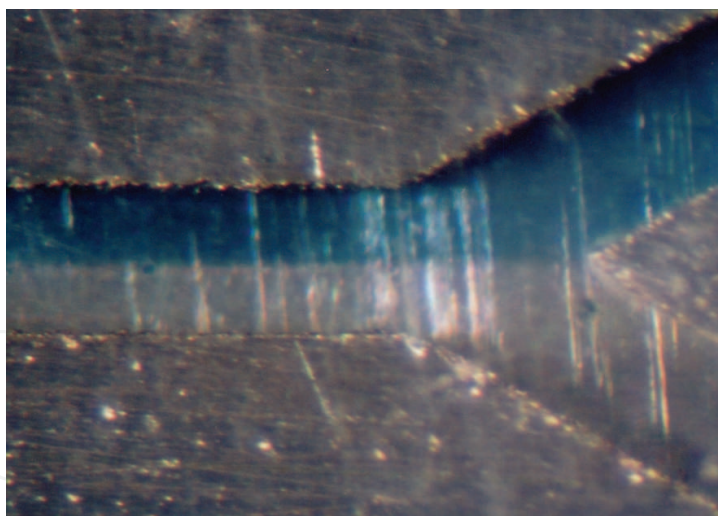


Fig. 3. Flow profile within microchannel (Cibacron blue solution and water)

The solute in the feed solution began to diffuse into the deionized water right after the contact. A predetermined amount of samples was taken at the channel outlet, and the samples were analyzed with a UV spectrophotometer (UV-1600, Shimadzu Co., Ltd.) or a refractive index meter (830-RI, Japan Spectroscopic Co., Ltd.). The liquid flow rates on both sides were determined by weighing the liquid samples collected for a defined time. Although the inlet liquids were fed at the same rate, unequal flow rates were observed at the outlets. In order to adjust the flow rate of each side, capillaries were installed at the effluent tubing. All experiments were carried out at a temperature of  $298 \pm 1$  K.

## 2.2 Mass-transfer model

Within the range of liquid velocities examined, the Reynolds number varies from 17 to 172. The liquid flow examined is laminar. Even when two kinds of aqueous solutions are brought into contact at the same velocity in one microchannel, a contacting plane is formed. If a concentration difference exists between the two liquids, the solute will diffuse from the concentrated to the less concentrated side across the contacting plane. Under conditions of short contact time for a mass-transfer and a considerably larger solute concentration in the feed-side than the recovery-side, the penetration model (Higbie, 1935) can be applied for solute diffusion.

$$N_A = 2\sqrt{\frac{D}{\pi\theta}}(C_i - C_b) \quad (1)$$

where  $N_A$  is the mass-transfer rate at the interface;  $D$ , the diffusion coefficient;  $\theta$ , the contact time;  $C_i$  the concentration at the contacting plane;  $C_b$ , the concentration of bulk liquid. The contact time;  $\theta$ , is calculated from the channel length of main portion of the channel,  $l$  divided by the average velocity of the contacting plane,  $v_{\text{int,av}}$ . Eq. (1) then reduces to

$$N_A = 2\sqrt{\frac{D v_{\text{int,av}}}{\pi l}}(C_i - C_b) \quad (2)$$

The value of  $v_{\text{int,av}}$  is determined from the simulated velocity profile within the microchannel. Details will be presented later. At the contacting plane, a parabolic profile is obtained by solving the Navier-Stokes equation with a finite element method.

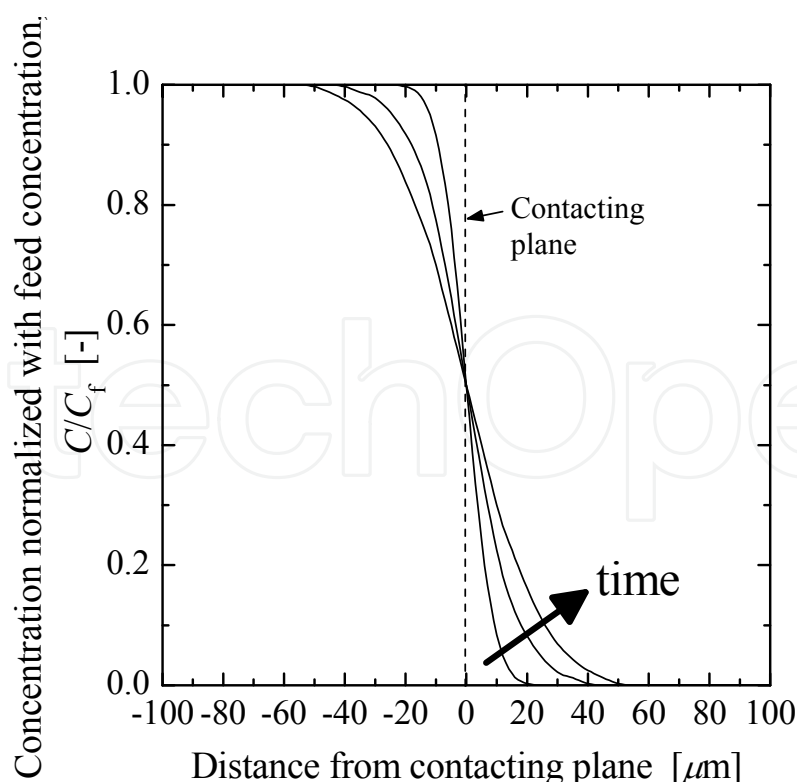


Fig. 4. Schematic representation of concentration profile within channel

The mass-transfer across the contacting plane is modeled on the basis of the diffusion in a semifinite slab. For unsteady diffusion in a semifinite slab, the concentration profile at the cross section of the main portion is schematically represented in Fig. 4. Since the two liquids are aqueous solutions, a symmetrical profile is formed in the vicinity of the contacting plane. The profile decays with the progress of contact time with maintaining a constant concentration at contacting plane,  $C_i$ . When deionized water is used as the recovery liquid,  $C_i$  equals to  $0.5C_f$ , and the bulk concentration,  $C_b$ , can be assumed to be zero. Then Eq. (2) is simplified to

$$N_A = \sqrt{\frac{D v_{\text{int,av}}}{\pi l}} C_f \quad (3)$$

The mass balance around the recovery-side of the channel can be expressed as

$$N_A A = v_{\text{av}} S (C_{\text{out}} - C_{\text{in}}) \quad (4)$$

where  $A$  is the area of the contacting plane;  $v_{\text{av}}$ , the average liquid velocity in the main portion of the channel, which is defined as the total volumetric flow rate divided by the cross-sectional area ( $200\mu\text{m} \times 200\mu\text{m}$ );  $S$ , the cross-sectional area of the recovery-side, which is half the cross-sectional area of the main portion;  $C_{\text{out}}$ , the outlet concentration of the recovery liquid; and  $C_{\text{in}}$ , the concentration at the inlet of the recovery liquid. The area,  $A$ , is expressed as the channel height,  $h$ , multiplied by the length of the main portion of the channel,  $l$ , and the area  $S$  is written as  $hL$ , where  $L$  is half the channel width. Hence, Eq. (4) becomes,

$$N_A = \frac{L v_{\text{av}} (C_{\text{out}} - C_{\text{in}})}{l} \quad (5)$$

For the supply of deionized water to the recovery-side,  $C_{\text{in}}$ , is set to zero. Therefore, the mass balance around the recovery-side is written as

$$N_A = \frac{L v_{\text{av}} C_{\text{out}}}{l} \quad (6)$$

Attention should be paid to the difference between  $v_{\text{av}}$  and  $v_{\text{int,av}}$ . The former is a three-dimensional average, while the latter is two dimensional. Velocities calculated at the contacting plane,  $v_{\text{int}}$ , are normalized with  $v_{\text{av}}$ , and the values of  $v_{\text{int}}/v_{\text{av}}$  are plotted in Fig. 5. A parabolic profile is obtained and the velocity of the contacting plane is averaged to determine the value of  $v_{\text{int,av}}/v_{\text{av}}$ . As is presented in the figure, the value of  $v_{\text{int,av}}/v_{\text{av}}$  is 1.4. The result suggests that the two-dimensional average of  $v_{\text{int,av}}$  is 1.4 times bigger than the three-dimensional average,  $v_{\text{av}}$ . Combining Eqs. (3) and (6) gives the outlet concentration of the recovery-side,

$$C_{\text{out}} = \frac{\sqrt{D}}{L} \sqrt{\frac{1.4l}{\pi v_{\text{av}}}} C_f \quad (7)$$

This is the fundamental equation for determining the solute diffusion coefficient from the measurement of the solute concentration at the outlet of the recovery-side.

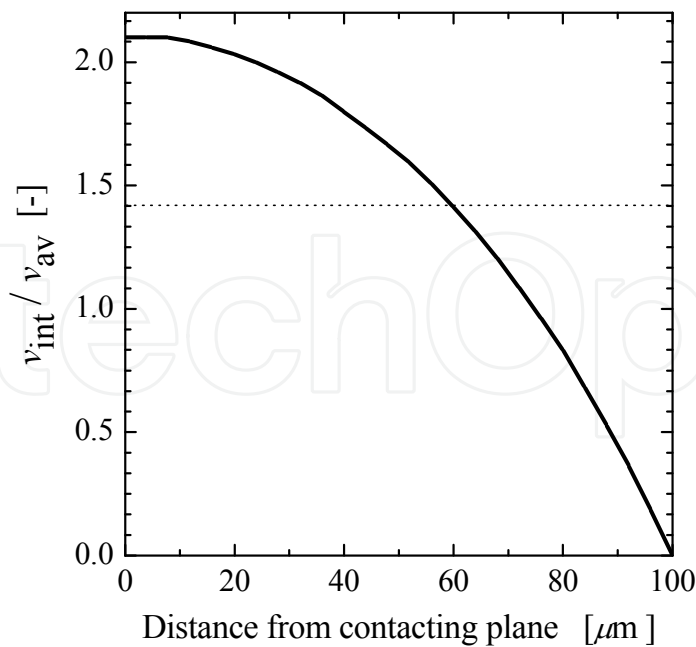


Fig. 5. Simulated velocity profile at contacting plane

2.3 Effect of contacting and splitting flow on mass-transfer

A typical experimental result for the benzoic acid concentration at the outlet of the recovery-side is presented in Fig. 6. The horizontal axis is taken as  $v_{\text{av}}^{-0.5}$  in order to examine the validity of the penetration model. The observed concentration scatters at a given liquid velocity. Although the two liquids were supplied at the same velocity, the outlet liquid velocities on each side were unequal. The results clearly show the difficulty in flow splitting.

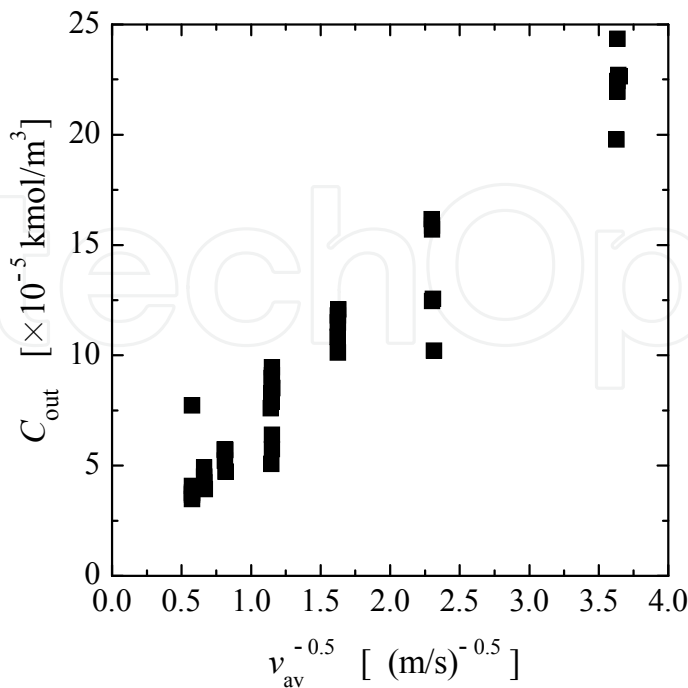


Fig. 6. Typical experimental results of benzoic acid concentration at recovery-side outlet



The concentration profile shown in Fig. 4 suggests a very steep curve around the contacting plane. When the liquid split equally, the separation was right at the contacting plane. Because of the big concentration change around the contacting plane, a small disturbance results in a large variation in the concentration. One of the possible reasons for the disturbance is the lack of precision in the machining of knife-edge fabrication or the attachment of fine air bubbles to the walls of the branch channels.

Splitting flows in microchannel devices seems to be a key technique for the practical application to chemical processing. Our results suggest that special attention should be paid to the fabrication of the knife-edge and the channel wall. This implies that the success of microchannel devices depends strongly on the accuracy of fabrication. This limitation act as a barrier to the development of microchannel devices. We seek a method that can use rather simple devices for exploiting the characteristics of microchannel devices.

The measurement of diffusion coefficients was carried out to understand the mass-transfer characteristics of microchannels. To acquire the solute concentration for equal flow splitting, an intentional change was given to the ratio of the flow rate of the feed-side to the recovery-side, and the solute-concentration data in the recovery-side was recorded. The flow rate were changed by setting capillaries on tubing to obtain a ratio of around unity. The solute concentration corresponding to the ratio of unity was taken to be as the value for equal splitting. Hence, this value is referred to as the relevant concentration.

Figure 7 shows a typical example of determining the relevant concentration. The ratio,  $R_{out}$ , is defined by  $Q_r / Q_f$ , where  $Q_r$  and  $Q_f$  are the flow rates of the recovery-side and the feed-side, respectively. The ratio varied within the range of  $\pm 10\%$  around unity. The concentration value for the ratio unity is found from the curve fitting of the observed data. The measurements were repeated for each liquid velocity in order to determine the relevant concentration. The values are plotted in Fig. 8. It can be seen that  $C_{out}$  increases linearly with  $v_{av}^{-0.5}$ , and that this dependence demonstrates the validity of the penetration model.

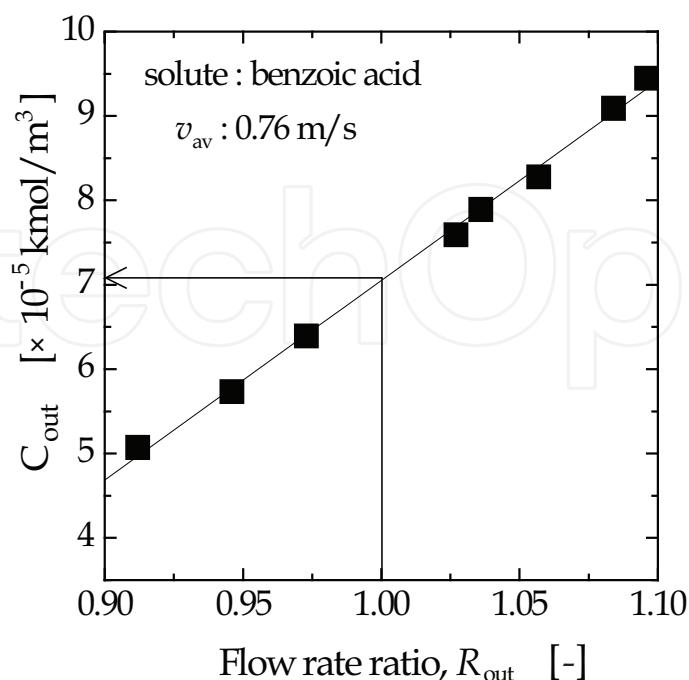


Fig. 7. Determination of relevant concentration at recovery-side outlet for equal flow splitting



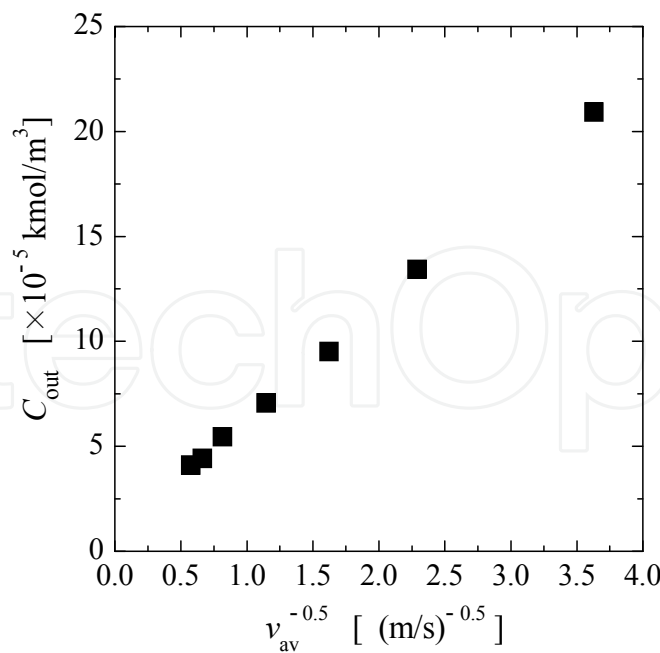


Fig. 8. Relevant concentration of benzoic acid at recovery-side outlet  
According to Eq. (7), the diffusion coefficient of the solute can be determined from the slope of  $C_{out}$  plotted against  $v_{av}^{-0.5}$ .

$$\text{The slope} = \frac{\sqrt{D}}{L} \sqrt{\frac{1.4l}{\pi}} C_f \tag{8}$$

Since the diffusion coefficient of benzoic acid has been well investigated, it is used as a standard material for determining the diffusion coefficients. The slope calculated from the reported diffusion coefficient,  $9.0 \times 10^{-10} \text{ m}^2/\text{s}$  at 298K (Yang and Matthews, 2000), is  $7.54 \times 10^{-5}$ .

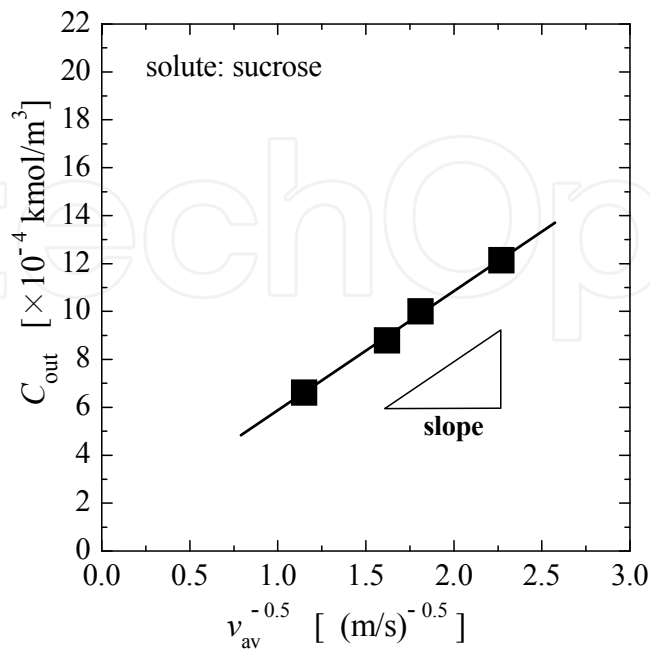


Fig. 9. Determination of diffusion coefficient of sucrose

For experiments with benzoic acid, the observed slope determined from the least mean squares approximation is  $5.35 \times 10^{-5}$ . The agreement of this value with this predicted one is poor. Hence, a correction factor,  $P$ , should be multiplied with the observed slope value to fit the predicted slope value. The value of  $P$  is found to be 1.4. To verify the applicability of the basic equation (8), diffusion coefficients of other solutes are determined with this method and the values are compared with the reported ones. Four solutes - sucrose, glycine, tryptophan, and urea - are selected. The results of  $C_{out}$  plotted against  $v_{av}^{-0.5}$  are shown in Figs. 9 to 12.

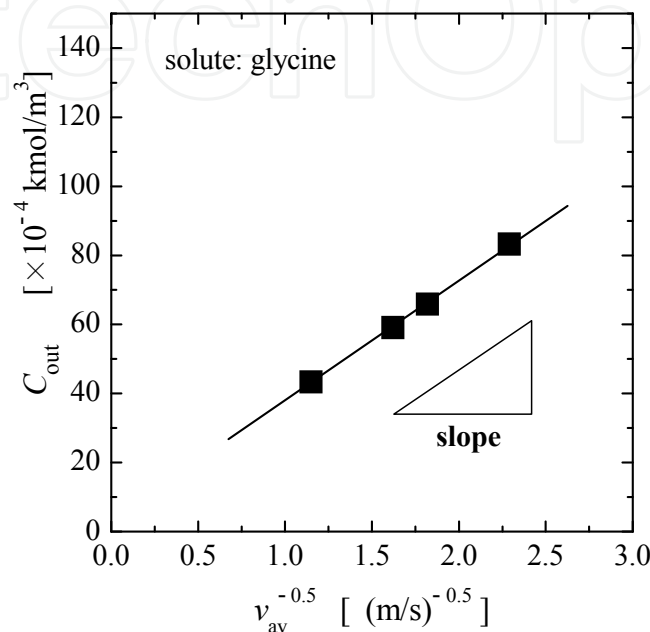


Fig. 10. Determination of diffusion coefficient of glycine

Solute	Diffusion coefficient (observed)[m <sup>2</sup> /s] at 298K	Diffusion coefficient (literature)[m <sup>2</sup> /s] at 298K
Benzoic acid	$9.0 \times 10^{-10}$	* $9.0 \times 10^{-10}$
Sucrose	$5.2 \pm 0.2 \times 10^{-10}$	** $5.2 \times 10^{-10}$
Glycine	$1.2 \pm 0.1 \times 10^{-9}$	** $1.1 \times 10^{-9}$
Urea	$1.4 \times 10^{-9}$	** $1.4 \times 10^{-9}$
Tryptophan	$5.6 \times 10^{-10}$	*** $5.7 \times 10^{-10}$

\*(Yang and Matthews, 2000),  
\*\*(Jost, 1952),  
\*\*\*Calculated value from the correlation (Wilke and Chang, 1955)

Table 1. Comparison of the diffusion coefficient.

The diffusion coefficients are calculated from the corrected slope of the plots, and the results are compared with the reported diffusion coefficients in Table 1. The observed diffusion coefficients agree well with the reported values or the correlated values. The validity of the basic equation is thus confirmed for the determination of the unknown diffusion coefficient of solutes. The correction factor,  $P$ , to be applied to Eq. (8) expresses the characteristics of the

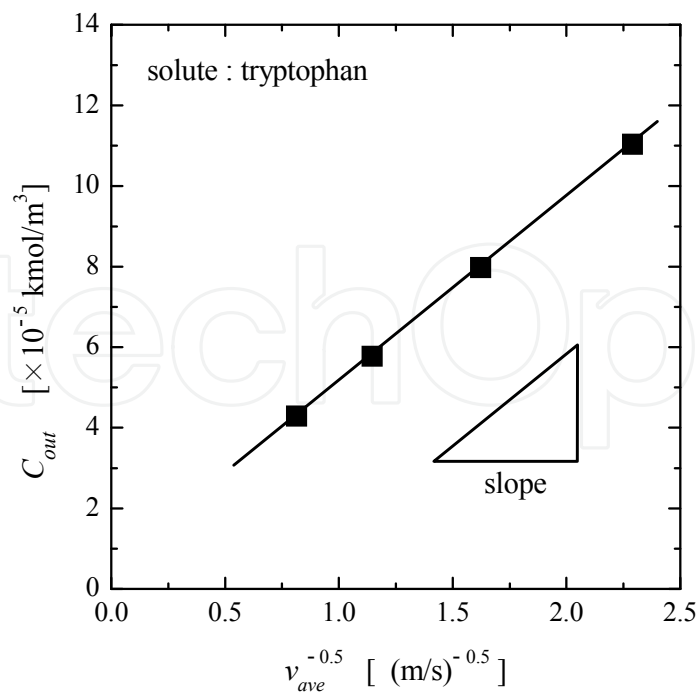


Fig. 11. Determination of diffusion coefficient of tryptophan

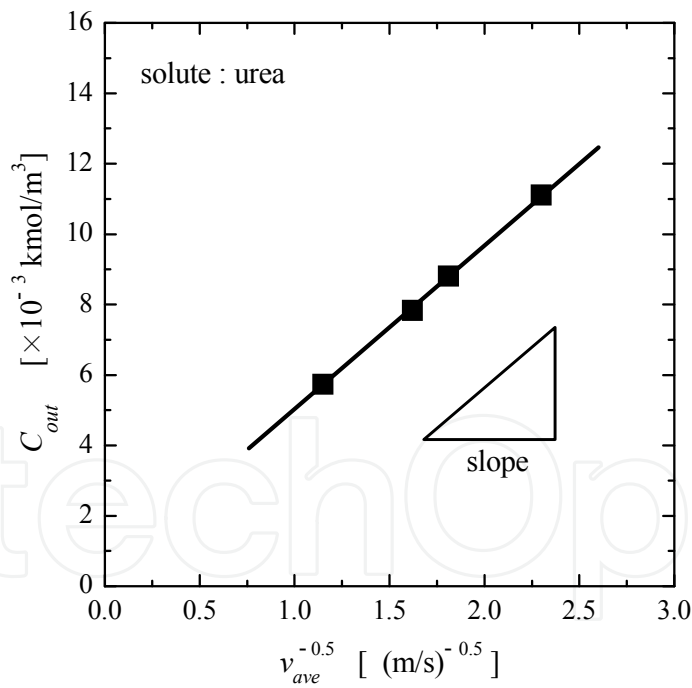


Fig. 12. Determination of diffusion coefficient of urea

mass-transfer in this microchannel. Since the  $P$  value is independent of the diffusing species, it can be influenced by the flow behavior in the microchannel. The value of the correction factor is bigger than one, which implies that the observed amount of the solute transferred is smaller than the theoretical amount. In the following chapter, the correction factor is determined by using microchannels of various configurations. The values are compared to discuss the flow behavior in a double-Y-type microchannel.

### 3. Flow characteristics in double-Y-type microchannel

#### 3.1 Channel configurations and materials

Four types of microchannels were fabricated on the stainless sheet as shown in Fig. 11. The type(a) channel is used as a benchmark; it was used in the previous chapter. For type(b), (c) and(d) channels, the depth of the channels, which is determined from the thickness of the stainless sheet is set at  $200\mu\text{m}$ . There are three channels of different depths for type (a); depths are 100, 200, and  $300\mu\text{m}$ . Type (b) and (c) have different angles of knife-edge. Type (d) has a short contacting length for two liquid flows. For all types, the channel width is  $200\mu\text{m}$ . As was described in the section 2.1, the channel was set in the module, and mass-transfer experiments were carried out using benzoic acid and sucrose as transferring material. On the feed side, an aqueous solution of each material was supplied, and deionized water flowed on the recovery side.

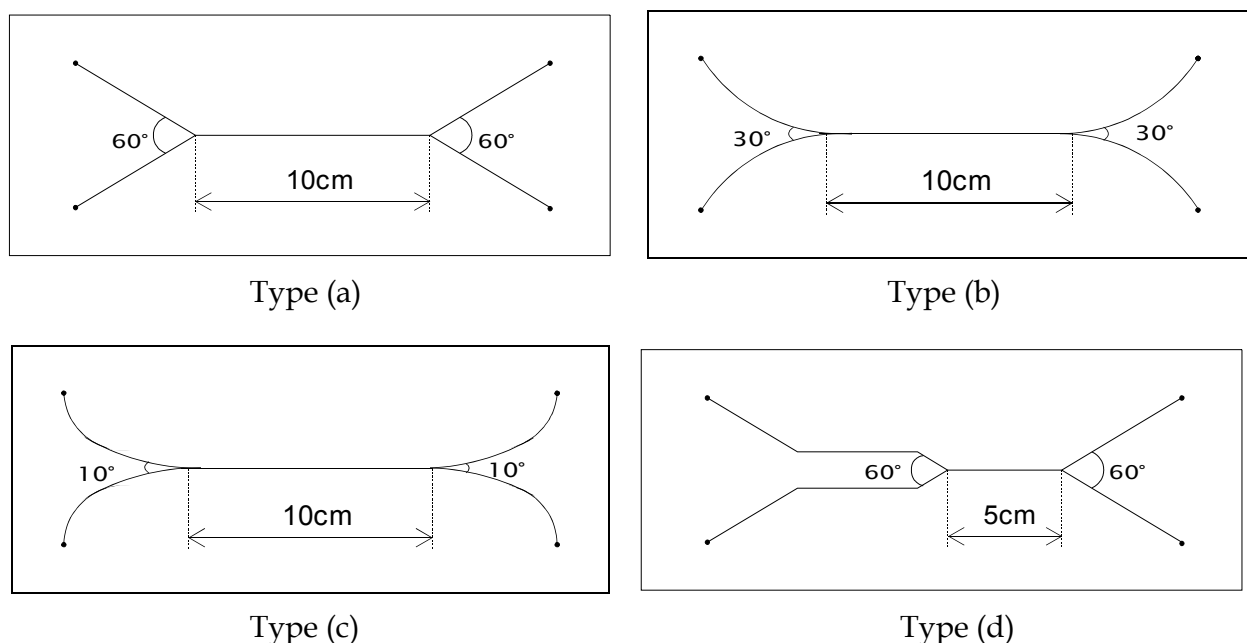


Fig. 11. Configuration of microchannels

#### 3.2 Effect of channel thickness

Sucrose was transferred in three type(a) microchannels of different depths (100, 200, and  $300\mu\text{m}$ ). The concentration values at the outlet of the recovery side were compared to examine the effect of the channel depth on the mass-transfer. Since the channel width,  $200\mu\text{m}$ , is constant, mass-transfer area increases linearly with the depth. The volumetric liquid flow rate is also proportional to the depth. Therefore, the amount of solute transferred is doubled when the depth doubled. It is expected that there will be no influence of channel depth on the outlet concentration. This trend is also predicted by the simulation. The experimental result is shown in Fig. 12. All the data located on a single line and the mass-transfer behavior was the same as the one expected.

The correction factor,  $P$ , was determined for the three channels by using the literature value of the diffusion coefficient for sucrose. For all the channels examined, the  $P$  value of 1.4 was to obtain the appropriate diffusion coefficient of sucrose. This fact suggests that the flow behaviour is similar for all microchannels used.

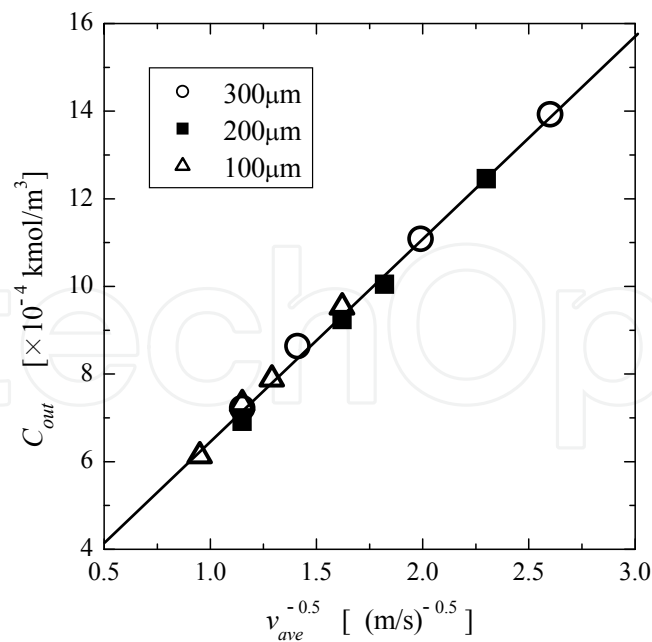


Fig. 12. Effect of channel thickness on outlet concentration of sucrose

3.3 Effect of channel length

The residence time of the liquid in contact changes with a change in the channel length. Therefore, without any disturbance in flow, the amount of solute transferred is unaffected by the channel length but by the residence time of the liquid. Sucrose was transferred

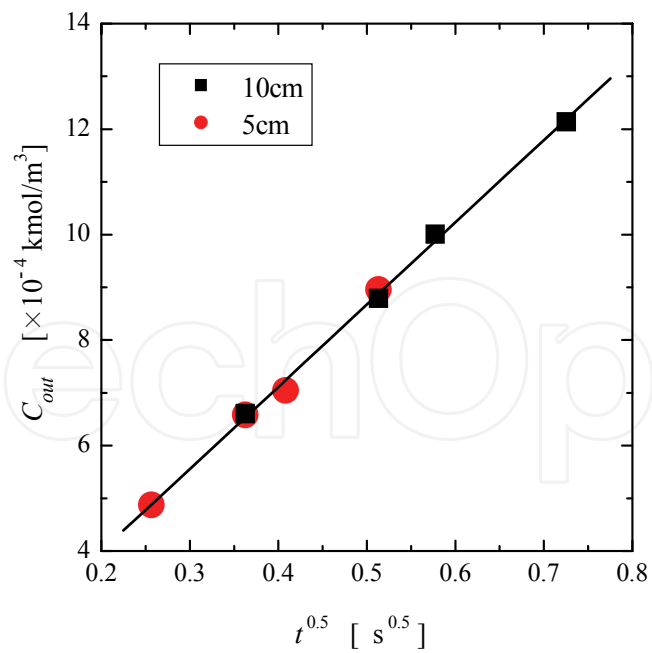


Fig. 13. Effect of channel length on outlet concentration of sucrose

in the type (d) channel having a channel length of 5cm, and the result was compared with that for the type (a). The outlet concentration was plotted against the square root of average residence time of the liquid,  $t^{0.5}$  in Fig. 13. Equation (7) is rewritten as

$$C_{\text{out}} = \frac{\sqrt{D}}{L} \sqrt{\frac{1.4l}{\pi v_{\text{av}}}} C_f = \left( \frac{C_f}{L} \sqrt{\frac{1.4D}{\pi}} \right) t^{0.5} \quad (9)$$

The factor,  $P$  for the type (d) channel was 1.4, which was the same for the type (a) and also as channels of different thickness of type (a). The accordance demonstrates that the factor is not an intrinsic value of a specific channel, and characterizes a general flow behavior found in the double-Y-type channels.

### 3.4 Effect of knife-edge angle

Using type (b) and (c) channels, we carried out sucrose transfer experiments were carried out to determine the each  $P$  value. The simulation predicts no effect of the knife-edge angle on the mass-transfer. In contrast with the simulation, the experimental result suggested that the mass-transfer was affected by the angle. The sucrose concentration at the outlet was plotted against  $v_{\text{av}}^{-0.5}$  in Fig. 14. Since each curve had a different slope, the correction factor,  $P$ , had to be modified to fit the diffusion coefficient of sucrose.

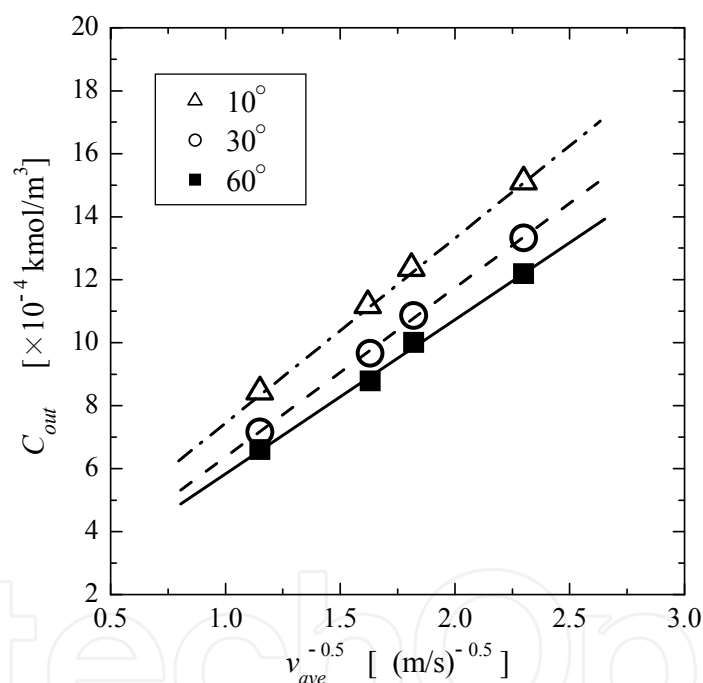


Fig. 14. Effect of knife-edge angle on outlet concentration of sucrose

Figure 15 shows a comparison of the  $P$  values for the three channels, type (a), (b), and (c). The  $P$  value changed with the knife-edge angle, and the result for benzoic acid showed a tendency similar to that in the case of the sucrose. With decreasing the knife-edge angle, the  $P$  value approaches unity, which corresponds with the pure penetration model. Now, it becomes clear that the factor,  $P$ , illustrates the characteristics of the contacting and the splitting flow in the double-Y-type channels.

How does the flow affect mass-transfer? This is a very important question; however, the answer to it is still difficult to find. One possible explanation is the effect of a secondary flow occurring in curved channels. At the knife-edge of the double-Y-type channels, two bent flows come together, or one flow is bended to split into two streams. It seems that the

phenomena that occur at the contacting point have relatively strong impact on the mass-transfer. If the secondary flow induced a rotating motion whose direction was normal to the flow axis, the mass-transfer across the interface could be influenced. In addition to the flow in the bulk region, there can be some wall effects on the flow such as the slip of the liquid or surface roughness, which were pointed out by Tretheway (2002) and Mala (1999).

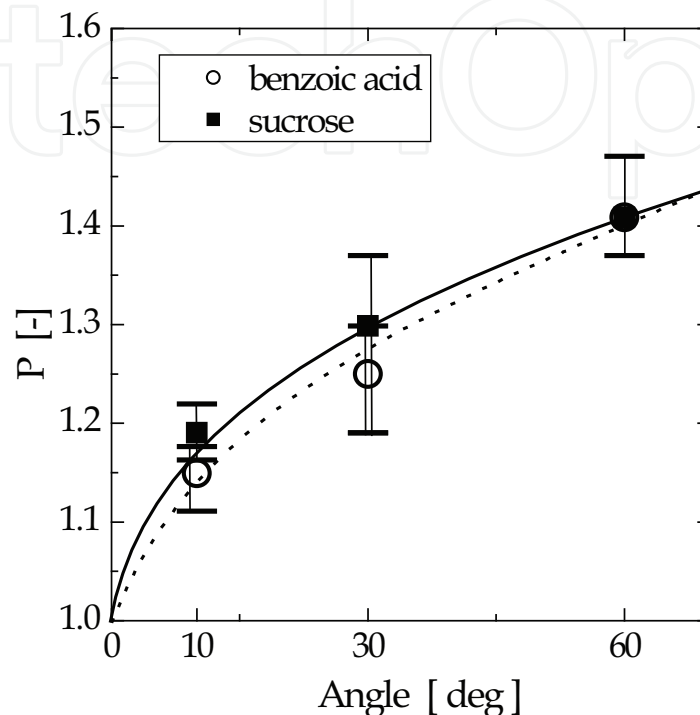


Fig. 15. Effect of knife-edge angle on the correction parameter,  $P$

#### 4. Conclusion

A double-Y-type microchannel device has been used for the mass-transfer from aqueous solutions to deionized water. Although the two liquids were carefully supplied at the same rate, the effluent flow rate for each side was sometimes unequal. Because of the unequal flow splitting, the observed values of the solute concentration of the recovery side were scattered. A method using the ratio of the flow rates has been proposed for determining a relevant concentration for equal flow splitting. On the basis of the relevant concentration, the diffusion of benzoic acid in water was successfully analyzed with the penetration model with a correction factor. Furthermore, the basic equation obtained with benzoic acid was successfully applied to measure the diffusion coefficients for sucrose, glycine, tryptophan, and urea. The observed values agreed fairly well with the reported or correlated values. This fact demonstrates that the presented method is also suitable for determining the unknown diffusivities of solutes.

To pursue the physical meaning of the correction factor, various channels with different knife-edge angles, channel lengths and depths were used for the mass-transfer experiment. The value of the correction factor is affected only by the knife-edge angle, and is independent of the channel length and depth. This factor probably characterizes the flow behavior at the knife-edge.



## 5. Nomenclature

$A$	area of contacting plane	[m <sup>2</sup> ]
$C_b$	concentration of bulk liquid on the recovery-side	[kmol/m <sup>3</sup> ]
$C_f$	concentration of liquid on the feed-side	[kmol/m <sup>3</sup> ]
$C_i$	concentration at the contacting plane	[kmol/m <sup>3</sup> ]
$C_{in}$	concentration of recovery liquid at the inlet	[kmol/m <sup>3</sup> ]
$C_{out}$	concentration of recovery liquid at the outlet	[kmol/m <sup>3</sup> ]
$D$	diffusion coefficient	[m <sup>2</sup> /s]
$D_{obs}$	observed value of diffusion coefficient	[m <sup>2</sup> /s]
$D_{lit}$	literature value of diffusion coefficient	[m <sup>2</sup> /s]
$L$	half width of the channel	[m]
$l$	length of the main portion of the channel	[m]
$N_A$	mass-transfer rate	[mol/(m <sup>2</sup> s)]
$S$	cross sectional area of the recovery-side	[m <sup>2</sup> ]
$t$	average contact time defined by $l/v_{av}$	[s]
$P$	correction parameter for Eq/(8)	[-]
$R_{out}$	flow rate ratio defined as $Q_r/Q_f$	[-]
$Q_f$	flow rate for the feed-side	[m <sup>3</sup> /s]
$Q_r$	flow rate for the recovery-side	[m <sup>3</sup> /s]
$v_{av}$	average liquid velocity	[m/s]
$v_{int, av}$	average liquid velocity of the contacting plane	[m/s]

### Greek symbol

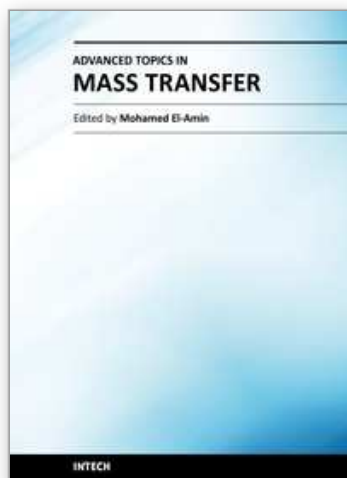
$\theta$	contact time defined by $l/v_{int, av}$	[s]
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