### MASS TRANSFER IN SEMI-FLUIDIZED BEDS

by

Yung Chia Yang

B. S., National Taiwan University, China, 1955

-----

A THESIS

submitted in partial fulfillment of the

requirements for the degree

MASTER OF SCIENCE

Department of Chemical Engineering

KANSAS STATE UNIVERSITY
OF AGRICULTURE AND APPLIED SCIENCE

LD 2668 T4 1959 Y36 c.2 Documents

## TABLE OF CONTENTS

INTRODUCTION
LITERATURE SURVEY
EXPERIMENTAL
Apparatus
Procedure
Materials
Analysis
Physical Constants
THEORY
RESULTS OF EXPERIMENTS
CORRELATION OF RESULTS
DISCUSSION
CONCLUSIONS AND SUMMARY
TABLE OF NOMENCIATURE
ACKNOWLEDGMENT
BIBLIOGRAPHY
APPENDIX

### INTRODUCTION

'Semi-fluidization' is a new and unique fluid-solid contact operation in which the expansion of a fluidized solid particle bed is only partially allowed. The purpose of the present investigation was to study the mass transfer aspect of the process of 'Semi-fluidization' (9) (10).

The rate of mass transfer between solid particles and a flowing fluid is important in many chemical engineering operations, such as leaching, crystallization, and heterogeneously catalyzed reactions. In these applications, the solid particles may be disposed in either fixed or fluidized beds; both schemes have wide utility (8).

A state of fluidization results when the flow rate of fluid passing upward through a bed reaches a magnitude sufficient to buoy the solid particles and provide a sufficient space of free board inavailable for free upward expansion of the bed. The fluidized-bed operation has the following major advantages over the fixed bed operation: uniform exposure of particles to the fluid, ease of maintaining constant operation conditions due to the violent mixing action of solid and fluid, and more adequate removal and transport of particles from vessels than in fixed bed systems. However, the fluidization process is not immune to some serious defects of its own; such as loss of driving potential for the transport process within the bed which is largely due to the backmixing of the solid particles and fluid; attrition and elutriation of the solid particles; requirement of considerable free board above the bed; and finally erosion of the walls of the containing vessel (8).

Recently it was proposed to partially eliminate those defects of the fluidized bed and to attain a type of solid-fluid contactor which compromises the feature of both fixed and fluidized beds (9) (10). By partially restricting expansion of the fluidized bed, it was found that a definite fraction of the solid particles formed a fixed bed immediately below the top eieve plate, as shown in Plate I, and that the rest of the particles remained in a fluidized condition below the packed eection. The name of eemi-fluidization was taken from this phenomenon (9) (10).

Such possibilitiee are suggested from many of the correlations of transport process (4) (24) within the solid-particle-fluid contactore, which are claimed to be equally applicable both to the fixed and to the fluidized bed. For instance, the mass transfer factor, jd, of Chu and his co-worker (4) suggests that, irrespective of the type of operation (fixed bed or fluidized bed), the rate of mass transfer can be altered if the porosity of the bed can be changed.

In the fixed bed, the porosity is a function of geometrical configuration of the particles and in the conventional fluidized bed, porosity is a function of the characteristice of the solid particles and fluid, and the flow condition of the fluid and is not subject to arbitrary control. In other words, the porosity of a fluidized bed must be treated as a dependent variable rather than as an independent variable of the process. A typical set of available data on bed expansion is plotted in Plate II in which the porosity of the beds is plotted against the rate of fluid flow (8).

While the expansion in a conventional fluidized bed is unrestrained, in a 'Semi-fluidized bed' the bed expansion is restricted by a porous sieve plate introduced above the expanding bed thus forcing formation of a fixed bed above the fluidized bed. By adjusting the position of the top sieve plate, the overall bed porosity can be varied, and the desired heights of fixed and fluidized sections for optimum driving potential for mass, heat and momentum transfer may be obtained (9) (10). This means that the overall porosity of the semi-fluidized bed is a factor arbitrarily controllable within the limits of the rigidly fixed bed and fully fluidized bed and may be considered as an independent variable of the operation or process.

### LI TERATURE SURVEY

There is no published work available on any aspects of 'Semi-fluidization' except the one which reported the preliminary results of this investigation (9). However, the extensive theoretical and semi-empirical investigations have been done by Fan et al (10).

As mentioned in the previous section, a semi-fluidized bed contains both the fixed bed and fluidized beds simultaneously. The literature on mass transfer in fixed and fluidized beds prior 195h was extensively reviewed by Chu (25). Therefore, only the major publications appearing before 195h were reviewed, especially the ones which deal with the combined correlation of the mass transfer in both fixed and fluidized beds and the ones which contain information on the benzoic acid-water system.

### EXPLANATION OF PLATE I

A picture of semi-fluidized bed

PLATE I



# EXPLANATION OF PLATE II

Porosity of bed of benzoic acid particle fluidized with water (8) Ordinate: E , Porosity of bed, dimensionless

Abscissa: G, Liquid rate, lb/(hr.)(sq. ft.)

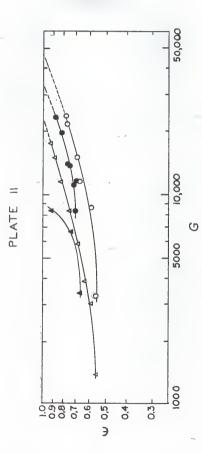
Legend:

O, 8-10 mesh benzoic acid particle

●, 12-14 mesh benzoic acid particle

 $\Delta$  , 20-24 mesh bengoic acid particle

▲, 24-28 mesh benzoic acid particle



Gamson, Thodos and Hougen (15) made psychrometric measurements for studying the rate of evaporation of water from spherical and cylindrical porous pellets into air streams. In their work, porous pellets were soaked in water usually for several hours, the excess water was drained off, the pellets were rolled in cheesecloth to remove the excess adhering water on the surface and without removal of capillary moisture. It was, therefore, expected that the interfacial surface area between the gas stream and the liquid was different from the geometric surface area of the solids on which the mass transfer coefficients calculated by Camson et al., (15) were based. Bed thicknesses not exceeding 2 1/2 in. were used with particle sizes as large as 3/4 in. In such shallow beds the entrance and exit effects and axial mixing effects might play important roles. Moreover, the height and the fractional void volume for a bed made up of a few layers of solids are quantities of questionable significance. The data were presented by plotting the jd factor against the Reynolds number,  ${
m GD}_{
m p}/\mu$  . The following equations were used to correlate their data:

$$J_{d} = 0.989 \left(\frac{D_{p}G}{\mu}\right)^{0.441} \qquad \text{for } \frac{D_{p}G}{\mu} > 350$$

$$J_{d} = 16.8 \left(\frac{D_{p}G}{\mu}\right) \qquad \text{for } \frac{D_{p}G}{\mu} < 40$$
(2)

$$J_{\rm d} = 16.8 \left(\frac{n_{\rm p} G}{\mu}\right)^{-1} \qquad \qquad \text{for } \frac{n_{\rm p} G}{\mu} < \mu_0 \tag{2}$$

Additional data to those of Gamson, Thodes and Hougen (15) were obtained by Wilke and Hougen (34) for the purpose of investigating further the region of lower gas flow rate. Equation (2) was replaced with the following equation:

$$J_{d} = 18.2 \left(\frac{D_{p}G}{\mu}\right)^{-0.51}$$
 for  $\frac{D_{p}G}{\mu} < 100$  (3)

Hurt (18) studied the following systems; (a) the adsorption of water vapor from moist air by particles of silica gel and by particles coated with phosphorous pentoxide, (b) adiabatic humidification of air over silica gel pellets wetted with water, (c) evaporation of naphthalene flakes into air and hydrogen streams. The results for (a) were considered doubtful. Data were given for case (b) and (c) together with the results obtained from calculation of Ht. The fractional void volumes were not measured. It was reported as follows:'! . . for different size particles of similar shape no satisfactory correlation of the data into a single curve could be obtained. Data are, accordingly, presented as separate curves for each size and shape investigated.'! The author apparently did not recognize that the plot of Ht/Dp against DpG/ $\mu$  would bring the data for different sizes into a single line.

Rates at which naphthalene was vaporized into air, hydrogen, and carbon dioxide streams from fixed and fluidized beds were measured by Resnick and White (27). In their fluidized bed correlation of jd factor against Reynolds number, two modes of fluidization were noted; an initial smooth bubbling type where the jd factor increased with increasing N'Re. Mesh size appeared as a parameter on a plot of jd vs. N'Re and Resnick and White correlated their data in the slugging region by replotting Jd/NSc 0.57 against N'Re/ 9. Where 6 is the dimensionless group:

$$\frac{D_{\rm p}^3 P_{\rm g} G(P_{\rm s} - P_{\rm g})}{II2}$$

Studies by Taecker and Hougen (31) were identical with those of Gamson, et al. (15), and Wilke and Hougen (34), except that Raschig rings and Berl saddles were used instead of spheres and cylinders. The effective particle diameter,  $D_p$ ', for the spherical shape packings used, was calculated by  $D_p$ ' =  $\sqrt{A_p/\pi}$ , where  $A_p$  is the dry surface area for one packing unit. Even with this modification, the jd factors for Raschig rings and partition rings were 19% lower than those for solid spheres and cylinders. Berl saddles gave valves about 30% higher than for the rings. In order to correlate their data the following equations were developed:

$$J_{\rm d} = 1.251 \left(\frac{\rm GD_p!}{\mu}\right)^{-0.41}$$
 for  $\frac{\rm GD_p!}{\mu} < 620$  (4)

$$Jd = 2.24 \left(\frac{GD_p!}{\mu}\right)^{-0.51} \qquad \qquad for \frac{GD_p!}{\mu} > 620$$
 (5)

Hobson and Thodos (17) passed water through a bed of celite spheres presoaked in a saturated solution of either water in isobutyl alcohol or of water in methyl ethyl ketone. The variation of the effluent concentration with time was measured so that the transfer rates would be calculated at "zero time" when the particles were believed to be completely wetted by saturated solution. The questions that concerned Gaffney and Drew (13) were: (1) how accurate was the extrapolation to a somewhat uncertain "zero time" for a curve of rapidly changing slope? (2) did the pellets have excess solution on their surface initially? (3) if not, how were they removed without depleting the surface? The following generalized equation, correlating also the data of Gamson, Thodos and Hougen (15) and Wilke and Hougen (3h) was recommended by Ergun (7):

$$\log J_{d} = 0.7683 - 0.9175 \log(N'_{Re}) + 0.0817 \left[\log N'_{Re}\right]$$
 (6)

McCune and Wilhelm (23) studied the mass transfer ratee from 2-naphthel pellets into water etreams. The work of these authors conetituted an excellent investigation free from most of the uncertaintiee of the earlier studies. Use of uniform size pellets assured accurate estimation of the geometric surface area of the solids which constituted both the surface exposed to flow and the interfacial surface for mass transfer. Ample bed heights were used. The fraction void volumes of the fixed beds could have been easily calculated from the bulk densities,  $\rho_{\rm b}$ , and the particle density of the pellets,  $\rho_{\rm s}$ , by the use of the relationship  $\epsilon=1-\rho_{\rm b}$ . This, however, apparently was overlooked by the authors, who determined the fractional void volume for each type of packing in a separate apparatus, although the pellets may not have been packed to the same bulk densities in the separate container. Data were presented as a plot of the j<sub>d</sub> factor against Reynolds number, and data were represented by two straight lines:

$$J_{\rm d} = 1.625 \left(\frac{D_{\rm pG}}{\mu}\right)^{-0.507}$$
 for  $\frac{D_{\rm pG}}{\mu} < 120$  (7)

$$J_{\rm d} = {}^{0.687} \left(\frac{{}^{\rm p}_{\rm p}{}^{\rm G}}{\mu}\right)^{-0.327}$$
 for  $\frac{{}^{\rm p}_{\rm p}{}^{\rm G}}{\mu}$ >120 (8)

Evans and Gerald (8) measured mass-transfer coefficients for fixed and fluidized beds of benzoic acid granules dissolving in water. Four different sizes of benzoic acid granules were used. By empirically modifying an application of the Reynolds analogy, the fluidized-bed data were all correlated by a single line, with mass velocity as the only flow

variable. To express the data the following equation was developed:

$$J_{\rm d} \frac{(1-\epsilon)({\rm Dp})^{2.6}}{f'} = 2.3 \times 10^{-10} \, ({\rm N}_{\rm Re}^{1})^{1.33} \tag{9}$$

Chu, et al. (14) presented mass-transfer data for transfer between regular chaped solids and a turbulent gas etream for both fixed and fluidized granular beds, and a simple generalized correlation was achieved. In contradiction to two previous investigations on gas-colid fluidized beds, the data were found to be in agreement with practically all the fixed bed data of widely varying systems, as well as with data for fluidized liquid-colid and gas-solid beds. The equations for mass transfer could be written as:

$$J_{\rm d} = 5.7 \left[ D_{\rm p} G/\mu \ (1-\epsilon) \right]^{-0.78}; \ 30 > \left[ D_{\rm p} G/\mu \ (1-\epsilon) \right] > 1 \ (10)$$

$$J_{\rm d} = 1.77 \Big[ D_{\rm p} G/\mu \ (1 - \epsilon) \Big]^{-0.4 \mu} 10,000 \times \Big[ D_{\rm p} G/\mu \ (1 - \epsilon) \Big] > 30 \quad (11)$$

The Chilton and Colburn (3) analogy of mass transfer and pressure drop for flow in conduits was modified for flow in granular beds. The resulting relation holds remarkably well over a wide range of Reynolds and Schmidt numbers, making possible the prediction of mass transfer data in granular beds from the more easily obtainable pressure drop data. The equation relating mass and momentum transfer is simply stated as  $j_{d} = f/10$ .

Ishine, Otake and Okada (20) studies mass transfer between solid benzoic acid particles in fixed beds with upward and downward flowing streams of waters. The range of experimental and correlative variables were as follows: particle shape and size, 3.5, 4.5 and 5.5 cm. cylinders with equal diameter and height; Reynolds number, 1 to 140; Schmidt numbers,

1180 to 1610; fixed bed depth, 1.3 to 13 cm. The correlated results using  $j_{\rm d}$  factor and Reynolds number were:

$$J_{\rm d} = 2.1 \, (N_{\rm Re})^{-0.80} \qquad \frac{D_{\rm p}G}{A} < 5$$
 (12)

$$J_{\rm d} = 1.444 \, (N_{\rm Re})^{-0.507} \qquad 5 < \frac{D_{\rm p}G}{44} < 140$$
 (13)

Shirai, (30) used the data obtained by several investigators (13) (17) (23) and obtained a single line correlation of the fixed beds, fluidized beds and single particle data. Shirai (30) plotted NSh $\in$  vs N'ReNSc $^{1/3}$  for mass transfer and similarly NNU  $\in$  vs N'ReNPr $^{1/3}$  for heat transfer. These can be expressed by equations:

For mass transfer 
$$N_{Sh} \in = 2.0 + 0.75 \, N_{Re}^{-1/2} N_{Sc}^{-1/3}$$
 (1h)

For heat transfer 
$$N_{Nu} \in = 2.0 + 0.75 \text{ N}_{Re}^{1/2} N_{Pr}^{1/3}$$
 (15)

Recently, Wakao and others (32) further modified Shirai's mass transfer equation to the expression:

$$N_{Sh} = 2.0 \quad 0.52 \quad N_{Re}^{0.54} \quad N_{Sc}^{0.35}$$
 (16)

Galleway et al., (14) studied the effect of packing configuration on mass and heat transfer in beds of stacked spheres. Experimental measurements were reported on the simultaneous rates of mass and heat transfer for the surface evaporation of water into air from two orthorhombic, one rhombohedral, and two simple cubic assemblages of uniform celite spheres, in the Reynolds number range 300-1200. The measurements were confined to the pre-determined constant rate drying period of the spheres. No appreciable differences in mass or heat transfer factors were found for the various assemblages, except for the cubics, which exhibited slightly

lower transfer factors than the other assemblages at the higher Reynolds number. The negative slope of the log. j vs. log. Re curves was designated the "index of turbulence inhibition". This index was correlated with the projected fractional free area of the packings, as well as with the wall effect. Various methods proposed by investigators of randomly packed beds for modifying the mass transfer factor and/or the Reynolds number failed to improve the overall correlation of the results. The inapplicability of a simple analogy between momentum and mass transfer, as proposed by Ergun (7) was demonstrated.

Wakao and others (32) studied mass transfer rates from spherical and cylindrical 2-naphthel particles to water in packed beds. Data at low water velocities show that Sherwood number tends to approach 2.0 with decrease of N<sup>1</sup>Re. His data and the previouely published data of other investigators (8) (23) for spherical and cylindrical particles were generalized and correlated by:

$$N_{Sh} = 2.0 + 1.45 N_{Sc}^{1/3} N_{Re}^{1/2} N_{Re}^{2} 100$$
 (17)

$$N_{Sh} = 2.0 + 0.90 N_{Sc}^{1/3} N_{Re}^{0.60} N_{Re}^{1} 100$$
 (18)

The data for flakes (23) and broken solids (8) for the experimental range of N $^{1}$ Re > 1, agree with the results obtained from above the equations.

Wakao et al., (33) obtained the axial turbulent diffusion coefficients by measuring axial concentration gradients in water-fluidized beds of cylindrical 2-naphthel particles. When the concentration in the bed is very small as compared with the saturation concentration, the dissolution rate approximates a zero-order reaction expressed as:

$$U_0 \frac{dc}{dl} = E_Z \frac{d^2c}{dl^2} + kpaG_s$$
 (19)

By using the boundary condition at the inlet and outlet of the bed, that

is inlet 
$$l = 0$$
  $U_0C = E_Z (dc/dl)$  (20)

outlet 
$$l = L dc/dl = 0$$
 (21)

Equation (22) can be derived from equation (19)

$$\frac{C_{\rm L}}{C_{\rm L}} = n + \frac{E_{\rm Z}}{U_{\rm o}L} \left[ 1 - \exp \left\{ \frac{U_{\rm o}L}{E_{\rm Z}} (1 - n) \right\} \right] \qquad \text{in which}$$
 (22)

l is axial distance, L is bed height and n = 1/0

### EXPERIMENTAL.

### Apparatus

The experimental apparatus consisted of a main test column in which granular benzoic acid particles were dissolved by a measured stream of upward flowing water, and a container for retention of samples of each run for analysis. The diagram of apparatus is shown in Plate III.

Ordinary water was supplied from a 50-gallon constant-head tank (A). Water from the tank was forced through the system by a centrifugal pump (B). The water was initially directed through a globe valve and a Brooks type lll0 flowmeter (C) as a means of controlling and measuring the rate of flow. The test column consisted of two parts, a calming section (D) and a test section (F). The calming section was a pyrex tube, h inches inside diameter, 12 inches high, and packed with Raschig rings and Berl saddles to maintain even distribution of flow. The test section was a

vertical pyrex tube, 2 inches inside diameter and 24 inches high. Two braes plates clamping a filter cloth (E) were placed between the calming and the test eections in order to establish an even distribution of the upward flow. A rigid close-fitting sieve plate (G) was attached to a long rod which could be moved up or down to a fixed position. This was installed in the test column in order to adjust bed depth and prevent the complete expansion of the eolid bed. The emall holes in this sieve plate were punched evenly in order to obtain a uniform packed section above the fluidized section. Otherwise, an inclined surface would appear between the two phasee. The outlet line consisted of half-inch tubing, which carried the product to the retention tank.

In order to carry out two series of experiments in the present investigation, the apparatue was modified.

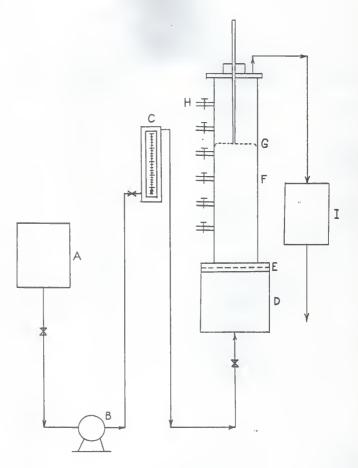
In the first series, only the inlet and outlet concentrations were measured and in the second eeries, the axial concentration gradient of the fluid was measured. The apparatus used in the later series was the same as those used for the former series, except the teet eection. This new teet section was made from a 2-inch inside diameter standard iron pipe with cocks attached for sampling, at 2 inch spacings. A sheet of filter cloth was attached by Duce cement to the inside wall over the outlet of each cock. This was intended to prevent the solid benzoic acid particles from plugging the sampling cocks. This is illustrated diagramatically in Plate III.

# EXPLANATION OF PLATE III

# Schematic diagram of experimental apparatus Legend:

- A. Water tank
- B. Pump
- C. Flowmeter
- D. Calming section
- E. Filter cloth
- F. Test section
- G. Top sieve plate
- H. Sampling outlet
- I. Retention tank

PLATE III



### Procedure

At the start of the operation, water was gradually run into the bottom of the calming section. The flow rate was regulated in such a way that air bubbles were eliminated in the inert packing material. Some difficulty was often encountered in removing air bubbles and small crystals from the test section but careful manipulation of the flow rate overcame these problems.

The solubility of the benzoic acid in water is low and in order to minimize the change of geometric shape of the particles during the experimental runs, the individual runs were not permitted to exceed an 80 seconds duration. A decrease of less than three per cent of the initial weight of the solid particles for a single run was experienced.

In a single run, a top-to-bottom sampling sequence was followed for determination of the concentration gradients. The advantage of a top-to-bottom sampling sequence was due mainly to the fact that there would be no influence in concentration gradient in the lower section of the fluid while the fluid from the upper region was being drawn off. If a reverse order were adopted, concentration gradients would vary appreciably and cause experimental errors due to the influence of the side stream sampling.

All the experimente were carried out at room temperature and the fluid temperature was measured at the inlet and outlet of the column.

### Materials

Granular particles of benzoic acid were prepared in three Tyler screen ranges; 8-10 mesh, 12-14 mesh and 2-24 mesh. U.S.P. benzoic acid

was melted in large cast, broken up in small lumps, and finally ground in a Braun Fulverizer type crusher. The particles were ecreened in a Rotap ehaker. Undesired screen fractions were remelted and reground. Adhering dust was removed and surface wetability improved by contacting the bulk preparation of each size with a large volume of distilled water. The particles were subsequently air-dried to constant weight (1).

### Analyeis

The determination of the concentration of the water sampler was made by direct titration using 0.04 N. sodium hydroxide with phenolphthalein indicator. No difficulty was encountered in securing a reproducible endpoints. The acid concentration varied from 0.0004 gm. acid per gm. of water to 0.0030 gms. acid per gm. of water.

### Physical Constants

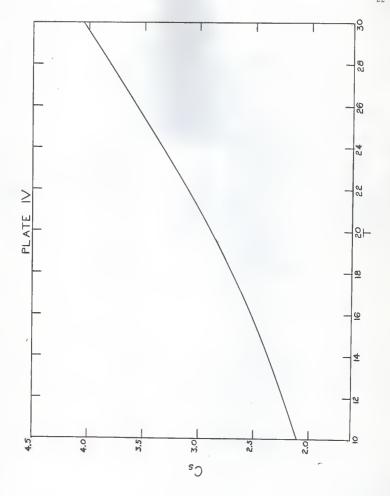
The physical constants used in the present investigation are briefly summarized below:

### Solubility, C<sub>S</sub>:

The saturation concentration of benzoic acid in water for use in evaluating driving force was obtained by reviewing all the available data in the literature (29) (19). Although there was a fair amount of disagreement, a sufficient number of substantiated values were found for the required temperature range to insure sufficient accuracy for this work. Solution saturated with the benzoic acid used in the present investigation was found by titration to be in agreement with the literature value (29). A plot of solubility related with temperature is shown in Plate IV. Also

# EXPLANATION OF PLATE IV

Solubility of benzoic acid in water Ordinate:  $C_{\rm B}$ , Solubility, gm./liter Abscissa: I, temperature,  $^{\rm O}{\rm C}$ 



solubility can be represented by an equation reported by Ishine (20) as function of temperature:

$$C_s = 0.17 + 0.00197t + 0.000202t^2$$
 (t: 0° - 30°C) (23)

# 2. Diffusivity, $D_v$ : and Schmidt Number $\mu/PD_v$

The Diffusivity and Schmidt number were taken directly from the data of Chang (2), quoted by Linton and Sherwood (22). Values of the Schmidt number at four temperatures were reported. A smooth reference curve, Plate V, VI, was plotted from these values.

### 3. Viscosity, A:

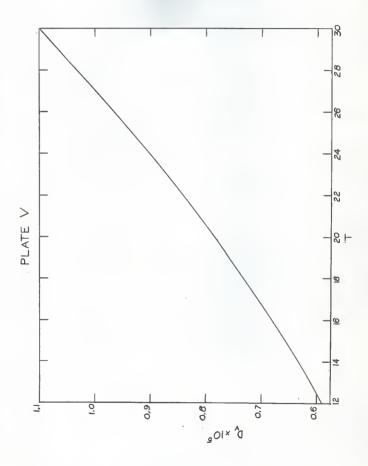
Since the concentrations of benzoic acid solution obtained in this work were very low, water viscosity was employed to calculate Reynolds numbers (26).

## 4. Density of particle, $\rho_s$ :

The density of particles was measured by weighing a sample of particles and adding it to a graduated volumetric cylinder. A known volume of saturated benzoic acid solution was added. The total volume of the particles plus added liquid was read and the volume of particles was determined by the difference to an accuracy sufficient for this work. The value determined was 1.293 gm./ml., somewhat above the literature value of 1.266 (19).

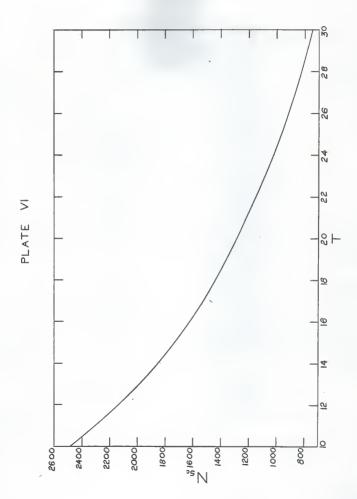
# EXPLANATION OF PLATE V

Diffusivity of benzoic acid in water Ordinate: Drx 105, Diffusivity, cm²/sec. Abscissa: T, Temperature, °C



# EXPLANATION OF PLATE VI

Schmidt number of benzolc acid in water Ordinate:  $N_{S_{\mathcal{O}}}$ , Schmidt number, dimensionless Abscissa: T, Temperature,  $^{\circ}$ C



# 5. Bulk density of particles, Pb:

The bulk density of particles was measured by weighing a sample of particles and adding it to a graduate volumetric cylinder. The volume of particles was directly readed. Plate VII showed the relationships between bulk density of particles and particle diameter for benzoic acid particles.

$$\rho_{b} = \frac{\text{Weight of benzoic acid}}{\text{Volume occupied by benzoic acid particles}}$$
(25)

# 6. Particle size, Dp:

The particle sizes reported here were the diameter of an equivalent sphere, having the same volume as the particles.

$$D_{p} = \left(\frac{6}{\tilde{\Pi}} \quad \frac{\text{Average weight of particle}}{\tilde{P}_{S}}\right)^{-1/3}$$
(26)

# 7. Shape factor, ø:

The shape factor of the particles, defined as the ratio of the surface area of the equivolume sphere to the surface area of the actual particle, varied from 0.50 to 0.65. A plot of shape factor against particle diameter was shown in Plate VIII.

$$\phi = \frac{\pi D_{\rm D}^2}{\text{Surface area of particle}}$$
(27)

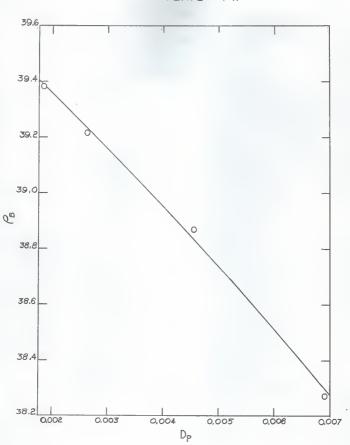
# 8. Specific surface, So:

The specific surface is defined as the ratio of surface area of the equivolume sphere to the volume of the particle, and was determined by measuring the pressure drop for a measured flow rate of saturated benzoic

### EXPLANATION OF PLATE VII

Bulk density of benzoic acid related to the particle diameter Ordinate:  $\rho_{\rm b}$ , Bulk density, lb./cu.ft. Abscissa:  $p_{\rm p}$ , Particle diameter, ft.

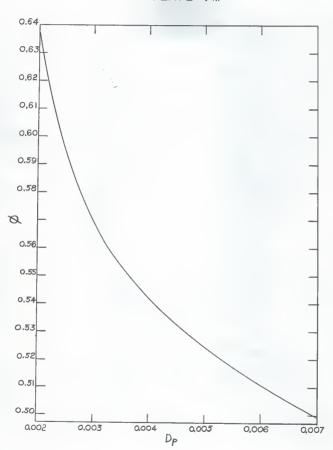




### EXPLANATION OF PLATE VIII

Shape factor of benzoic acid related to the particle diameter Ordinate:  $\phi$ , Shape factor, dimensionless Abscissa: Dp, Particle diameter, ft.





acid solution through a bed of particles. The equipment used for this purpose was similar to that of Carman (1) and his equation relating surface area to pressure drop was used.

$$S_0 = \frac{6}{D_0 \delta} \tag{28}$$

# 9. Porosity of solid particle bed, €:

The porosity of solid particle bed is defined as the ratio of the volume of voids in the bed to the total volume of the bed. It may be evaluated as one minus the ratio of bulk density to its solid density as:

$$\epsilon_0 = 1 - \frac{\rho_b}{\rho_s}$$
 (29)

When the bed expands from ho to h, the porosity becomes:

$$\epsilon = 1 - \frac{1 - \epsilon_0}{R} \tag{30}$$

where R is the ratio of the original bed height to the expanded bed height.

In the present work, the bulk density of the dumped particle bed was used to calculate the porosity of the packed bed. This porosity may be defined as the dumped porosity or dumped void, and can be considered as the porosity of a most loosely and randomly packed bed. It is shown in Plate IX. The properties of benzoic acid particles used in this investigation are listed in Table I.

### THEORY

To derive the rate equation for the mass transfer between the solid particles and fluid, the following assumptions were made:

### EXPLANATION OF PLATE IX

Dumped void related to the particle diameter Ordinate:  $\epsilon_0$ , Dumped void of particle, dimensionless Abscissa: Dp, particle diameter, ft.

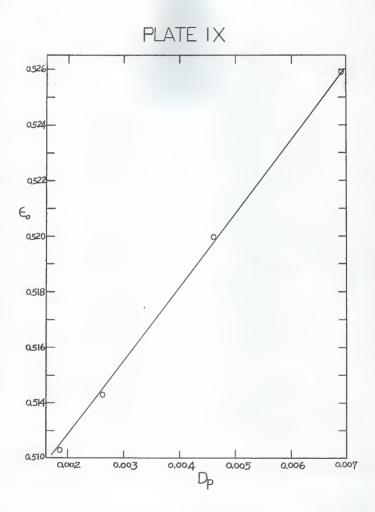


Table 1. Properties of benzoic acid particle

yler Screen Mesh	No. per	 Bulk density lb/ft3	. So, ft-1	Dp. ft.	Shap	Shape factor	e e
8-12	156.5	38.2757	1720	0.00693	0	0.500	0.5259
12-14	545	38.7128	2480	0.00456	Ó	0,531	0.5205
20-24	2870	39.2123	3890	0,00262	Ó	0.588	0.5143

- (1) The concentration of the fluid in contact with the particle surface is equal to the saturated concentration and the controlling resistance to the dissolution of solid is the diffusion through the liquid film surrounding the solid particles.
- (2) The velocity distribution and concentration of fluid are uniform at any cross-section of the test section.
  - (3) The specific surface of the particles in each bed is uniform.
- (4) Axial turbulent diffusion coefficient, Ez, in each bed rsmains constant under definite flow conditions.
  - (5) The process is isothermal.

Under those assumptions and the boundary conditions, the following material balance could be obtained for any bed height, dl:

$$U_0 \frac{dC}{dL} = E_z \frac{d^2C}{dL^2} + k_p a(C_s - C)$$
 (31)

whers Uo = superficial fluid velocity based on empty column.

C = concentration

Cs : saturated concentration

L = axial distance

 $\mathbf{E}_{\mathbf{Z}}$  = turbulent diffusion coefficient in axial direction, based on bed cross section

kp = particle to fluid mass transfer coefficient

a = specific surface of particles in bed

Assuming that longitudinal or axial mixing in the bed can be negligible:

$$E_{z} \frac{d^{2}C}{dl^{2}} = 0$$
 (32)

Then, equation (31) can be expressed as:

$$U_0 \frac{dC}{dL} = k_p a(C_s - C)$$
 (33)

By integration of the equation (33)

$${\tt U_0} \int_{{\tt C_0}}^{{\tt C_1}} \frac{{\tt d}\mathcal{C}}{{\tt C_s-c}} + {\tt U_0} \!\! \int_{{\tt C_1}}^{{\tt C_2}} \frac{{\tt d}\mathcal{C}}{{\tt C_s-c}} = ({\tt k_p})_{\tt f} \quad {\tt a_f} \int_{{\tt O}}^{{\tt L_1}} {\tt d} {\tt L} + ({\tt k_p})_{\tt pa} \; {\tt a_pa} \!\! \int_{{\tt L_1}}^{{\tt L_2}} {\tt d} {\tt L}$$

or 
$$U_0 \ln \frac{c_s - c_0}{c_s - c_1} + U_0 \ln \frac{c_s - c_1}{c_s - c_2} = (k_p)_f \text{ af } h_f + (kp)_{pa} \text{ a}_{pa} h_{pa}$$
 (35)

where  $Co = inlet concentration [M/L^3]$ 

 $C_{1}$  = concentration of the interface between the fixed and fluidized bed  $M/L^{3}$ 

 $C_2 = \text{outlet concentration} \left( M/L^3 \right)$ 

 $(k_p)f$  mass transfer coefficient in fluidized bed  $(L/\theta)$ 

 $(k_p)_{pa}$  = mass transfer coefficient in packed bed  $(L/\theta)$ 

 $a_f$  = specific surface of particles in fluidized bed  $\left( L^{-1} \right)$ 

apa = specific surface of particles in packed bed ( L )

hf = height of fluidized bed (L)

 $h_{pa}$  = height of fixed bed (L)

 $h = total height = h_f + h_{pa} ( \ \ )$ 

Equation (35) can be further simplified as:

$$U_{0}\ln \frac{C_{s}-C_{0}}{C_{s}-C_{2}} = (kp)_{f} a_{f} h_{f} + (kp)_{P_{a}} a_{P_{a}} hpa$$
 (36)

If the weighted mean value of  $\mathbf{k}_{p}\mathbf{\bar{a}}$  based on the overall bed height is defined as:

$$(k_p)_m a_m = (k_p)_f a_f \frac{h_f}{h} + (k_p)_{pa} a_{pa} \frac{h_{pa}}{h}.$$
 (37)

Then equation (36) may be transformed into the following form:

$$U_0 \ln \frac{C_S - C_0}{U_S - C_2} = (k_p)_m a_m h$$
 (38)

where (a)  $_{\rm m}$  = overall mean specific surface of particles in bed  ${\rm (kp)}_{\rm m}$  = overall mean mass transfer coefficient

Since

$$(\Delta C)_{lm} = \frac{\frac{C_2 - C_0}{ln \frac{C_8 - C_0}{C_8 - C_2}}}{(39)}$$

or 
$$\ln \frac{C_{s} - C_{o}}{C_{s} - C_{c}} = \frac{C_{2} - C_{o}}{(\Delta C)_{1m}}$$
 (40)

$$\frac{U_o (c_2 - c_o)}{(\Delta C)_{\downarrow m}} = (k_p)_m a_m h$$
 (41)

Then, the overall mean mass transfer coefficient can be calculated as

$$(kp)_{m} = \frac{U_{0}(C_{2} - C_{0})}{a_{m} h (\triangle C)_{\downarrow m_{o}}}$$
 (42)

or, by multiplying both sides of the equation by ho , the above equation can be further transformed into the form:

$$(k_L)_m = \frac{G'(c_2 - C_0)}{A(AC)_{1m}}$$
 (43)

where  $(k_L)_m$  = overall mean mass transfer coefficient  $(M/L^2\theta)$ 

G' = flow rate through bed  $(M/\theta)$ 

 $C_2$ ,  $C_0$  = inlet and outlet concentration (M/M)

A \* total particle surface in bed  $(L^2)$ 

In order to account for the effect of axial mixing, which was neglected previously, a correction factor (6), F, to modify the logarithmic mean driving potential can be introduced as:

$$(k_L)_m = \frac{G'(c_2 - c_0)}{A(\Delta c)_{1 \cdot mF}}$$
 (44)

or 
$$(k_L)_m F = \frac{C'(c_2 - c_o)}{A(\Delta C)_{lm}}$$
 (45)

In case the axial concentration gradient is available, the value of  $^{k}$ L in each bed can be directly calculated from the definition of  $^{k}$ L by use of the integrated mean driving force as:

$$(k_{\rm L})_{\rm f.} = \frac{G'(c_{\rm L} - c_{\rm o})}{A_{\rm f}(a_{\rm C})^{\dagger}t_{\rm c}} \tag{46}$$

$$(k_L)_{Pa} = \frac{G'(c_2 - c_1)}{A_P(ac)_{tp}}$$
 (47)

or the overall mean mass transfer coefficient may be evaluated as

$$(k_{L})_{m} = \frac{G^{1}(C_{2} - C_{0})}{A(\Delta C)_{t,m}}$$

$$= \frac{G^{1}(C_{2} - C_{1}) + G(C_{1} - C_{0})}{A(\Delta C)_{t,m}}$$

$$= \frac{(k_{L})_{p_{d}}A_{p_{d}}(\Delta C)_{t,m} + (k_{L})_{f}A_{f}(\Delta C)_{tf}}{A(\Delta C)_{t,m}}$$

$$= X(k_{L})_{p_{d}} \frac{(\Delta C)_{t,m}}{(\Delta C)_{t,m}} + (1 - X)(k_{L})_{f} \frac{(\Delta C)_{t,f}}{(\Delta C)_{t,m}}$$
(48)

where X  $_{\Xi}$   $\frac{A_{\underline{p}}}{A}$  , surface fraction of fixed bed particles or weight

fraction of packed bed.

A dimensionless parameter, the  $j_d$  factor, was first defined by Chilton and Colburn (3) for the purpose of correlating the transfer coefficient  $k_{\underline{l}}$  to the physical properties of the system. For dilute solution,  $j_d$  factor is defined as (3):

$$J_{d} = \frac{k_{L}}{G} \left( N_{Sc} \right)^{\frac{2}{3}}. \tag{49}$$

Another dimensionless parameter, the Sherwood number, has also been frequently utilized to correlate the mass transfer data. It is defined as (30) (32) (33):

$$N_{\rm sh} = \frac{k_{\rm p} p_{\rm p}}{n_{\rm r}} \tag{50}$$

As described already, in semi-fluidization, the overall porosity,  $\epsilon$ , can be arbitrary controlled by adjusting the moveable top sieve plate, and, therefore, it should be considered as an independent variable rather than dependent variable.

In general

$$J_{d} = f(N_{R_{o}}, \epsilon)$$
 (51)

or 
$$N_{sh} = f(N_{Re}, N_{Sc}, \epsilon)$$
 (52)

for the mass transfer between the solid and fluid phases flowing through the bed of solid particles.

### RESULTS OF EXPERIMENT

In the first series of experiments, measurements included the following variables: exit concentration, water temperature, water flow rate, bed depth, top sieve plate height, and particle and bed

characteristics. Computed variables derived from these measurements included  $k_L$ ,  $J_d$ ,  $N^{\dagger}_{Re}$ ,  $N_{Sh}$ ,  $N_{Sc}$ ,  $\in$  and  $J^{\dagger}_d$ .

The range of major variables were as follows:

Particle diameter: 8-10 mesh, 12-14 mesh, and 20-24 mesh.

Water temperature: 16.8 C to 24 C.

Bed depth: 3 in. to 6 in.

Voidage: 0.65 to 0.90

Modified Reynolds number: 5 to 130

Sherwood number: 21 to 116

Schmidt number: 1020 to 15h0

They are tabulated in Table 2 in the appendix.

Table 3 in the appendix contains the measured and derived data for the second series of experiments in which the axial concentration gradients of fluid were measured in addition to those variables mentioned above.

All the recalculated values for Evan and Gerald's (8) data are tabulated in Tables 4 and 5 in the appendix.

Concentration gradients measured in the second series of experiments are all graphically illustrated in the appendix.

### CORRELATION OF RESULTS

First, the mass transfer data were correlated in terms of the  $J_{\rm d}$  factor and modified Reynolds number as is illustrated in Plate X. The mass transfer coefficients were calculated on the basis of the overall logarithmic mean driving force, as was done by other investigators (h) (8) for the packed and fluidized bed mass transfer.

As defined previously, the  $J_{\rm d}$  factor for the dilute solutions is written as (28):

$$J_{d} = \frac{k_{L}}{G} (N_{Sc})^{2/3} = (N_{Sh})(N_{Re})^{-1}(N_{Sc})^{-1/3}$$
 (53)

As discussed previously in this investigation, instead of  $(k_L)_m$ ,  $(k_L)_m$ F was employed not only to show the existence of the axial mixing effect but also to indicate the use of the overall logarithmic mean driving concentration including both the packed and fluidized bed.

The broken line in Plate X represents the generalized correlation used by Chu, et al., (4) to correlate the mass transfer data in both packed and fluidized beds. The solid line correlates all the semi-fluidized data with the magnitude of the deviations comparable to the correlation used by Chu, et. al. (4). The equation for the solid line was found to be:

$$J_{\rm d} = 1.865 \left(\frac{N' \, {\rm Re}}{1 - \epsilon}\right)^{-0.48}$$
 (54)

Under convective conditions, either natural or forced, a relationship for mass transfer similar to that obtained for heat transfer may be expected to have a form (16):

$$N_{sh} = \int (N_{Re}, N_{Sc}, N_{Gr})$$
 (55)

where NSh, N'Re, NSc, and N'Gr are respectively the Sherwood, Reynolds, Schmidt and Grashof number for the mass transfer. Such relationship has been obtained theoretically by Eckert (5) (16) from a consideration of the boundary condition.

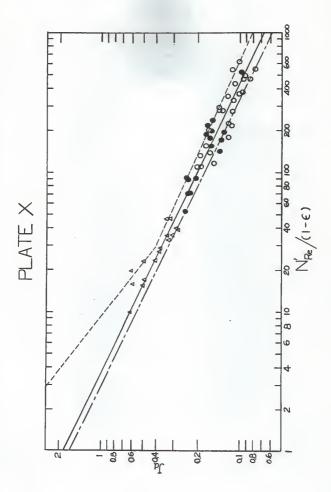
For mass transfer under the condition of forced convection where the Grashof number is unimportant, the general expression becomes:

### EXPLANATION OF PLATE X

Jd factor ws. N'Re/(l.	$J_{\rm d}$ factor vs. $N^{1}_{\rm Re}/(1-\varepsilon$ ) correlation of first series data by use
of logarithmic mean driving force.	riving force.
Ordinate: Jd, dimensionless	lonless
Abscissa: $N^{1}Re/(1-\varepsilon)$ , dimensionless	), dimensionless
Legend:	
	Correlation line of author's data
	Chu and other $(\mu)$ correlation line
	Correlation line of Evan and Gerald's data (8)

12-1μ mesh particle
 Δ 20-2μ mesh particle

O 8-10 mesh particle



$$N_{Sh} = f (N'_{Re}, N_{Sc})$$
 (56)

which boundary layer theory suggests, should take the form: (5)

$$N_{\rm Sh} \propto N_{\rm Re}^{1/2} N_{\rm So}^{1/3} \tag{57}$$

Froseling (11) and Garner and Suckling (16) from the study of the mass transfer from a single epherical particle in fluids derived a relationship which may be written in the form:

$$N_{Sh} = 2.0 + aN^{1}R_{e}^{1/2} N_{Sc}^{1/3}$$
 (58)

Fuchs (12) showed theoretically that the limiting value of the dimensionless mass transfer Sherwood number is 2.0, which appears in the right hand side of the equation, when the sphere become infinitely small.

As already reviewed in the previous chapter, Shirai (30) derived a similar equation for a generalized correlation of mass transfer including packed beds, fluidized beds, and eingle spheres into one single line.

Shirai's correlation is written as:

$$N_{Sh} \in = 2.0 + 0.75 \, \text{N'}_{Re}^{1/2} N_{Sc}^{1/3}$$
 (59)

Plate XI showe the eimilar correlation for the present data and a comparison with Shirai's equation (30). The best line through the data is expressed as:

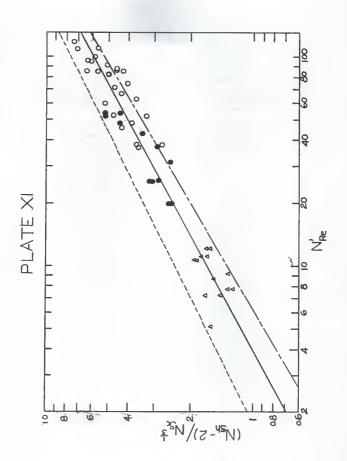
$$N_{Sh} \in = 2.0 + 0.46N^{1}_{Re}^{0.543} N_{Se}^{0.33}$$
 (60)

Since the  $J_{\rm d}$  factor correlation was derived primarily for turbulent condition, no allowance is made for the effect of transfer by natural convection and the molecular diffusion. According to the Chu et al., (4)

# EXPLANATION OF PLATE XI

Nsh6 -2 NSc1/3	$vs.\ N^{}_{Re}$ correlation of first series data by use of
logari thmi	logarithmic mean driving force.
Ordinate:	$\frac{NSNG}{NScL/3}$ , dimensionless
Abscissa:	$^{\mathrm{N}^{\prime}}\mathrm{Re}$ , dimensionless
Legend:	
	Correlation line of author's data
i	Shirai's (30) correlation line
	Correlation line of Evan and Gerald's data
	O 8-10 mesh particle
	• 12-14 mesh particle
	Δ 20-24 mesh particle

(8)



the  $J_{\mbox{\scriptsize d}}$  factor correlation can be generally expressed as a function of the Reynolds number as:

$$J_{d} = \frac{k_{L}}{G} (N_{SC})^{2/3} = a(\frac{N \cdot Re}{1 - \epsilon})^{-b}$$
 (61)

or, in terms of the Sherwood number it is written as:

$$N_{sh} = a(N_{Re})^{1-b} (N_{Sc})^{1/3} (1-b)^b$$
 (61)

The equation above can be modified by an analogical method used by previous investigators (12) (30) (32) (33) to account for the molecular diffusion at low Reynolds number as follows:

$$N_{\rm sh} = 2.0 \text{ a } (N_{\rm Re})^{1-b} (N_{\rm Sc})^{1/3} (1 - e)^{b}$$
 (62)

Then, a modified  $J_d$  factor,  $J^i_{d^i}$ , may be defined as:

$$Jd' = \frac{N_{sh} - 2}{(N'Re)(N_{Sc})I/3}$$

Plate XII shows the plot of jd' vs  $\frac{N'Re}{1-}$  , which yields the relationship:

$$J_{\rm d}' = 1.51 \left( \frac{N' \, {\rm Re}}{1 - \epsilon} \right)^{1/2}$$
 (64)

or 
$$N_{sh} = 2.0 + 1.51 \left( N_{Re}(1 - \epsilon) \right) \frac{1}{2} \left( N_{Sc} \right) \frac{1}{3}$$
 (65)

The correlations for the second series of the experiments, which are similar to the ones described above, are plotted in Plates XIII, XIV, and XV.

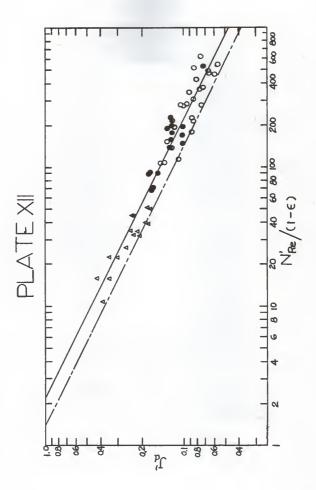
### EXPLANATION OF PLATE XII

J'd vs. N'Re/(1- E ) correlation of first series data by use of logarithmic mean driving force Ordinate: J'd, modified Jd factor, dimensionless Abscissa: N'Re/(1- E ), dimensionless Legend:

Correlation line of author's data Correlation line of Evan and Gerald's data (8)

O 8-10 mesh particle • 12-14 mesh particle

A 20-24 mesh particle



# EXPIANATION OF PLATE XIII

Jd factor vs.  $N^{\rm I}_{\rm Re}/(1-\xi$  ) correlation of second series data by use of integrated mean driving force

Ordinate: Jd, dimensionless

Abscissa: N'Re/(l- E ), dimensionless

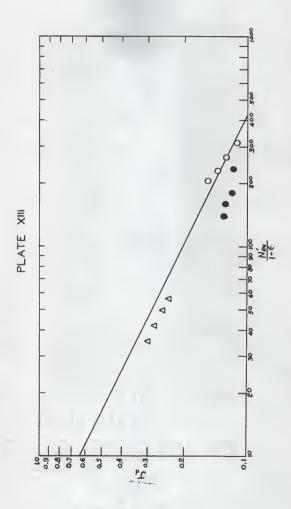
Legend:

Line of equation (54)

O 8-10 mesh particle

● 12-14 mesh particle

Δ 20-24 mesh particle



## EXPLANATION OF PLATE XIV

(Nsh6 -2)/NSc1/3 vs. N'Re correlation of second series data by use of integrated mean driving force

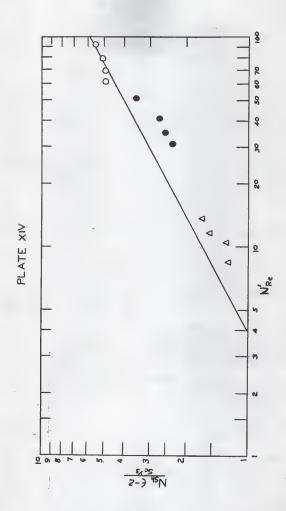
Ordinate:  $(N_{\rm sh}\xi-2)/N_{\rm Sc}1/3$ , dimensionless

Absclssa: N'Re, dimensionless

Legend:

Line of equation (60)

- O 8-10 mesh particle
- 12-14 mesh particle
- △ 20-24 mesh particle



### EXPLANATION OF PLATE XV

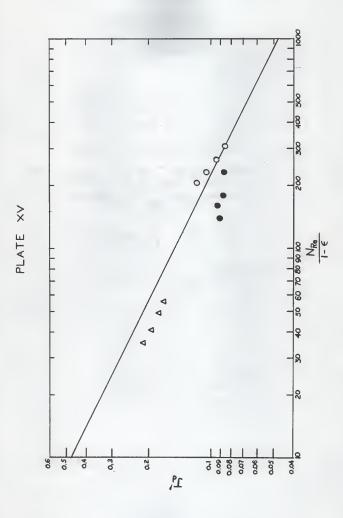
Jid vs.  $N^i_{Ro}/(1-\xi$ ) correlation of second series data by use of integrated mean driving force Ordinate: Jid, modified Jd factor, dimensionless Abscissa:  $N^i_{Ro}/(1-\xi)$ , dimensionless Legend:

Line of equation (65)

O 8-10 mesh particle

• 12-14 mesh particle

A 20-24 mesh particle



### DISCUSSION

In order to intelligently understand the effect of the packed bed formation on the mass transfer in the fluidized bed, the mechanism of the packed bed formation should be understood first. At least the size of the packed bed should be quantitatively correlated with the independent variables of the experiment and the characteristice of ordinary fluidization under eimilar flow conditions.

The experimental data of the present investigation, as shown in Plates XVI, XVII and XVIII, definitely indicate that the segregation of the particles depends on factors such as the particle and fluid characteristics, flow conditions and the expansion of the bed.

Based on a simple material balance, the height of the packed bed section in a semi-fluidized bed can be derived as follows (9) (10). It is assumed that the liquid is incompressible, the particles in a column do not interact with each other and the particle distribution is uniform. Referring to Plate XIX:

$$h_{pa}\rho_{s} (1 - \epsilon_{pa}) = (h_{pa} + (h_{fo} - h)) \rho_{s} (1 - \epsilon_{f})$$
 (66)

or 
$$h_{pa} = (h_{fo} - h) \left(\frac{1 - \epsilon_f}{\epsilon_{f} - \epsilon_{pa}}\right)$$
 (67)

where hpa = height of packed bed section

€pa .= porceity of the packed bed eection

hfo = height of the bed under complete fluidization conditions

 $\epsilon_{\rm f}$  = porceity of the fluidized bed section

h = height of the top eieve plate

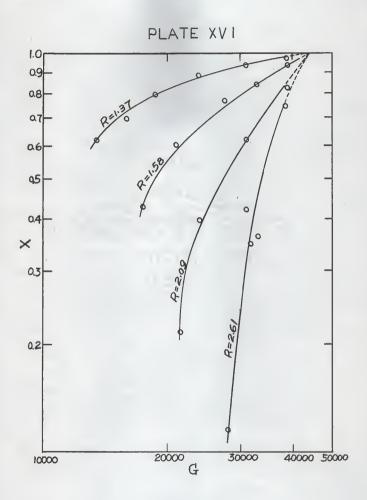
### EXPLANATION OF PLATE XVI

Weight fraction of packed particles

Particle size: 8-10 mesh

Ordinate: X, Weight fraction in packed bed, dimensionless

Abscissa: G, Fluid rate, lb/(hr.) (sq. ft.)



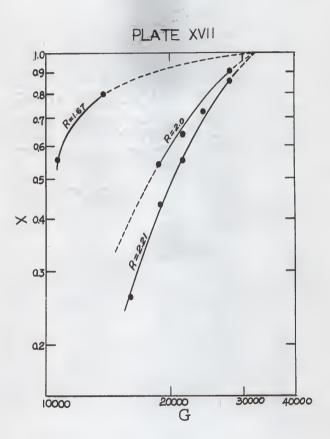
### EXPLANATION OF PLATE XVII

Weight fraction of packed particles

Particle size: 12-14 mesh

Ordinate: X, Weight fraction in packed bed, dimensionless

Abscissa: G, Fluid rate, lb./(hr.)(sq. ft.)



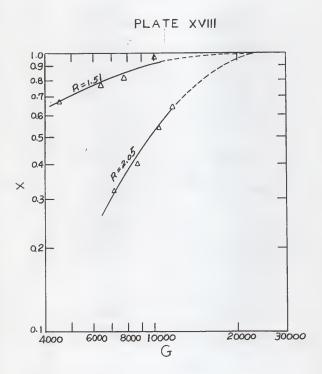
### EXPLANATION OF PLATE XVIII

Weight fraction of packed particles

Particle size: 20-24 mesh

Ordinate: X, Weight fraction in packed bed, dimensionless

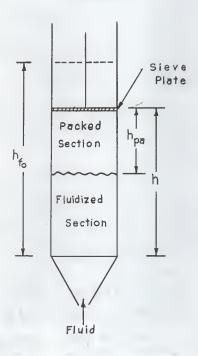
Abscissa: G, Fluid rate, lb./(hr.)(sq. ft.)



### EXPLANATION OF PLATE XIX

Schematic diagram of semi-fluidized bed

PLATE XIX



In the right hand side of the above equations, h and  $\epsilon$  pa are independent variables of the experiment. The relationship between  $h_f$  or  $\epsilon_f$  and flow conditions can be obtained from the Evan and Gerald'e data (8). The height of the packed bed section calculated from equation (67) and that obtained from the experimente are compared as shown in Plate IX.

The weight fraction of solid in the packed bed aection, I, then, can be calculated as:

$$I = \rho_{a} \cdot A \cdot h_{pa} (1 - \epsilon_{pa}) / W$$

$$= \frac{\rho_{e} A (1 - \epsilon_{pa}) (h_{fo} - h) (1 - \epsilon_{f})}{W}$$

$$(68)$$

The experimental data are compared with the calculated values based on the equation (68) as shown in Plate XXI.

The agreement indicated in Plate XX and Plate XXI is good considering the fact that the large and irregular eize particles of benzoic acid were employed and that the data in the range of  $\epsilon > 0.8$  were not accurate due to the difficulty of measurement of bed height (8) (9) (10).

As indicated in the figuree the deviation for the larger particlea ie greater than that for the smaller ones. The errore were probably due to the difficulty in measuring the expanded heights of fluidized beds of the coarse and the irregular particlee, and the assumption of uniform bed density.

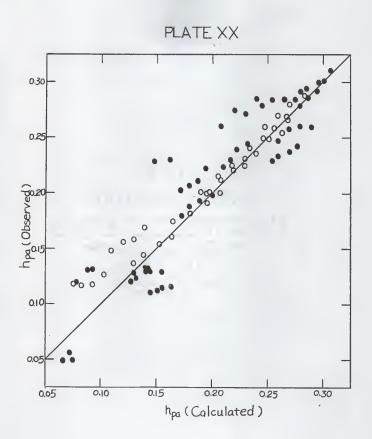
It is also interesting to note that the extrapolated curves of X vs. G for each particle sizes all converge to one value of G at X = 1 and these values of G coincide with the value of G at  $\in$  f = 1 obtained from

### EXPLANATION OF PLATE XX

Heights of packed bed section calculated based on equation (67) compared with that experimentally observed

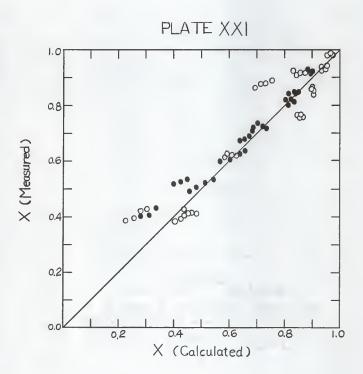
Ordinate: hpa, Observed height, ft.

Abscissa: hpa, Calculated height, ft.



### EXPLANATION OF PLATE XXI

Weight fraction of packed bed section calculated based on equation (68) compared with that experimentally measured Ordinate: X, Weight fraction measured, dimensionless Abscissa: X, Weight fraction calculated, dimensionless



the extrapolation of the curve of  $\epsilon_f$  vs. G as shown in Plate II. Theoretically, all the particles in the fluidized bed disappear, or in the other words, when X becomes unity the fluid flow rate reaches the maximum fluid velocity at which  $\epsilon_f = 1$ .

Chu and co-workers (h) have pointed out that the experimental error is effected by the degree of saturation of the outlet solution. The experimental errors are magnified by a factor of h at 90% saturation when used to compute the mass transfer coefficient and after 90% eaturation the degree of multiplication of experimental errore increase sharply, which makes precise measurements in this region quite difficult. Now, referring to Plate X, it may be seen that Chu's original  $J_{\rm d}$  factor correlation yields jd values consistently higher than the present experimental data. Chu's correlation was derived only for the ordinary fluidized and packed beds and the maximum deviation of his correlation appears to be 22% in the range of Reynolds numbers used in the present experiments. Furthermore, Evan and Gerald'e (8) data for benzoic acidwater system were not included in the original correlation by Chu et. al. (h). Evan and Gerald's (8) data were recalculated in terms of  $J_{\rm d}$  vs.  $N^{+}_{\rm L-f}$ 

It clearly indicated that Evan and Gerald's (8) data also deviate consistently from Chu'e et al., (4) original correlation with approximately the same magnitudes as those of the semi-fluidized bed data. Their packed bed data are correlated as:

$$J_{\rm d} = 2.132 \, \left( \frac{\rm N^{1}Re}{1 - \epsilon} \right) 0.5122$$
 (69)

and the fluidized bed data are correlated as:

$$J_{\rm d} = 1.340 \, \left( \frac{N^{1} \, \rm Re}{1 - \epsilon} \right) 0.4675$$
 (70)

Consequently, the overall correlation of Evan and Gerald's (8) data may be expressed as:

$$J_{\rm d} = 1.68 \, \left( \frac{N^{\, \rm l}_{\rm Re}}{1 - \epsilon} \right)^{0.49}$$
 (71)

Comparison of equations (71) with (54) reveals that, within the range of the variables of the present investigation and the errors of experimentation the semi-fluidized data may be correlated as well as the data for ordinary fluidized and packed beds by use of  $J_{\rm d}$  factor vs.  $\frac{N^{1}{\rm Re}}{1-\epsilon}$ , which is based on the overall mean mass transfer coefficient,  $(k_{\rm L})_{\rm me}$ . As mentioned previously, the  $J_{\rm d}$  factor used in Plate X should include the F factor to indicate that the overall logarithmic mean driving force was used in spite of the existence of axial mixing of the fluid and the abrupt change in the axial concentration gradient.

The data for the benzoic acid-water system were also not included in Shirai's (30) original generalized correlation including the fluidized bed, fixed bed, and a single particle. Evan and Gerald's (8) data were again recorrelated by the method suggested by Shirai (30). It is illustrated in Flate XI and the calculations are summarized in Tables 4 and 5, Appendix.

Plate XI indicates that the experimental data were consistently lower than the calculated values by approximately 38%. However, this is within the range of accuracy expected from the generalized correlation of Shirai (30) for any specific system. Evan and Gerald's (8) data are expressed as follows:

For packed bed:

$$N_{Sh} \in 2 + 0.424N_{Re}^{0.558}N_{Sc}^{1/3}$$
 (72)

For fluidized bed:

$$N_{Sh} \in = 2 + 0.230N_{Re}^{0.659} N_{Sc}^{1/3}$$
 (73)

For overall correlation:

$$N_{Sh} \in = 2 + 0.312N^{\dagger}Re^{0.608}N_{Sc}^{1/3}$$
 (74)

The overall correlation expressed by equation (7h) is again very similar to that for the semi-fluidized data as expressed by equation (65) and Plate XII. These similarities of the correlation between the ordinary packed beds and semi-fluidized beds by use of the methods suggested by Chu et.al.(h0) and Shirai (30) would indicate that it is possible, at least for the purpose of approximation to consider the semi-fluidized bed to possess uniform solid density distribution. When correlations are not available for any specific system for semi-fluidization, some form of the generalized mass transfer correlation of the ordinary packed and fluidized beds may be employed to estimate the mean mass transfer coefficients.

Equation (65) which was derived by modification of the  $J_d$  factor correlation to the present data yields a mean deviation of 11.7% while equations (54) and (60) give a deviation of 9.87% and 11.73% respectively. This implies that the use of the modified correlation as expressed by equation (65) does not improve the correlation. However, the form of this equation would agree with the expression predicted by the boundary layer theory which is:

$$N_{\rm Sh} \propto N_{\rm Re}^{1/2} N_{\rm Se}^{1/3} \tag{57}$$

In addition, the use of this expression offers much simpler means of calculation.

Evan and Gerald's (8) data were recorrelated in form of equation (65). The packed bed data are correlated as:

$$N_{Sh} = 2 + 1.483 \left[ (N_{Re})(1 - \epsilon) \right]^{1/2} \left( N_{Sc} \right)^{1/3}$$
 (75)

and the fluidized bed data are correlated as:

$$N_{Sh} = 2 + 1.031 \left[ (N_{Re})(1 - \epsilon) \right]^{1/2} \left[ N_{Sc} \right]^{1/3}$$
 (76)

and overall correlation equation may be expressed as:

$$N_{Sh} = 2 + 1.202 \left[ (N_{Re})(1 - \epsilon) \right]^{1/2} \left[ N_{Sc} \right]^{1/3}$$
 (77)

or as Plate XII indicates:

$$N_{Sh} \simeq 2 + 1.51 \left( (N_{Re})(1 - \epsilon) \right)^{1/2} \left( N_{Sc} \right)^{1/3}$$
 (77')

It is interesting to note that their data of ordinary packed and fluidized beds are well correlated by the same equation as equation (65). However, equation (65) or any other similar mass transfer equation has different significance when applied to the semi-fluidized bed that when applied to the ordinary packed and fluidized beds. As already stated, the overall porosity of the semi-fluidized bed is an independent variable of the experiment, i.e. it can be arbitrarily changed or fixed. It can be expressed as:

$$\epsilon = 1 - \frac{1 - \epsilon_0}{R} = \frac{V - W/\rho_s}{V} = 1 - \frac{W}{\rho_s h S}$$
 (78)

Where  $\epsilon_0$  and  $\rho_s$  are characteristics of the solid particle and V, h, W, and S are the variables which can be independently determined. Substitution of equation (78) into the equation (65) leads to the following relationship:

$$(k_{\rm L})_{\rm m} \ {\rm F} \propto (\frac{1}{\rm R})^{1/2} = {\rm R}^{-1/2}$$
 (79)

or

$$(k_L)_m F_{oc}(\frac{1}{hs})^{1/2} = (hs)^{-1/2}$$
 (80)

Or for the column with definite cross-section:

$$(k_L)_m F \propto (\frac{1}{h})^{1/2} = (h)^{-1/2}$$
 (81)

These characteristics of the semi-fluidized bed are illustrated in Plate XXII. In Plate XXII the data at flow rate  $G=28,000\ lb/(hr)(ft^2)$  are used.

On the other hand, there is no freedom to alter the porosity of the packed bed in the operation of a particular system, i.e.

$$(k_{\rm L})_{\rm m}F \neq f(\in) \tag{82}$$

or

$$(k_L)_m F \neq f (R \text{ or } h)$$
 (83)

Equations (54) (60) and (65), when applied to the fluidized bed, cannot be considered as explicit equations, since  $\varepsilon$  is again a function of the flow condition. In order to change  $\varepsilon$ , the flow rate has to be changed.

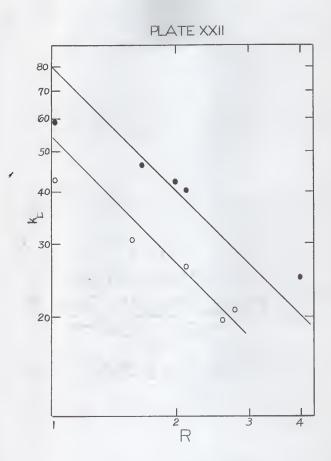
### EXPLANATION OF PLATE XXII

Expansion ratio vs. mass-transfer coefficient

Ordinate: kL, Mass-transfer coefficient, lb/(hr.)

(sq.ft.)

Abscissa: R, expansion ratio, dimension



This implies that the F-factors included in the correlation of the data from the first series of the experiments were close to unity, in other words, the effects of axial fluid mixing and the exietence of abrupt change in the axial concentration profile were emall. The valuee of F are plotted against X and  $\frac{N^*Re}{1-\epsilon}$  in Platee XXIII and XXIV respectively. The conditions of no axial mixing of fluid and existence of smooth concentration gradient profile lead to the value of F = 1. The maximum deviation as indicated in Plate XXIII and XXIV was 7.65% and no abrupt change in the axial concentration gradient.

Finally equation ( $\mu\beta$ ) developed earlier was tested by use of the experimental data. In Plate XXV, the calculated values are plotted against the experiment values. The validity of equation ( $\mu\beta$ ) indicates the internal mechanism of mase transfer in the semi-fluidized bed. It can be described as follows: the overall mass transfer coefficiente of eemi-fluidized beds represents the linear combination of the effective packed bed mass transfer coefficient and the effective fluidized bed mass transfer coefficient which are defined as  $(k_L)_{pa}$   $(\Delta C)_{tp}$  and  $(k_L)_f$   $(\Delta C)_{tof}$  respectively.

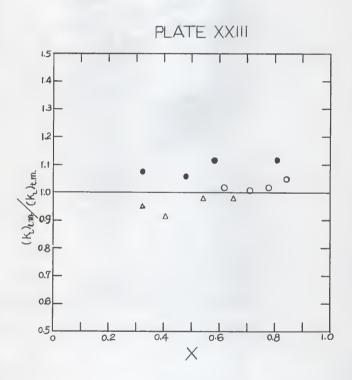
In the first series of experimente the axial concentration was not measured and in the correlation of the data, the F-factor, which was originally introduced by Epetein (6) was employed to indicate the use of the logarithmic mean driving force. The importance of the axial mixing in the solid-liquid contact process has been emphasized by various investigatore recently (8) (9). However, it appears that, so far, there is no way to predict exactly the axial mixing coefficient of a particular system

### EXPLANATION OF PLATE XXIII

 $(k_{\rm L})$ 1.m./ $(k_{\rm L})$ t.m. vs. weight fraction in packed bed

Ordinate: (kL)1.m./(kL)t.m., dimensionless

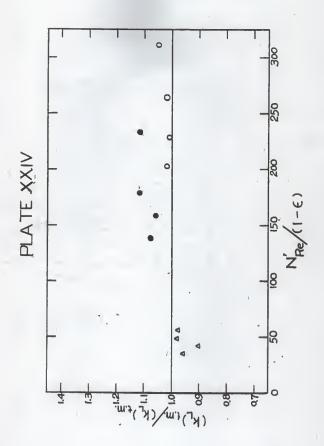
Abscissa: X, Weight fraction in packed bed, dimensionless



### EXPLANATION OF PLATE XXIV

 $(k_L)_{1,m_*}/(k_L)_{t,m_*} \quad vs. \quad N^1Re/(1-\varepsilon \ )$  Ordinate:  $(k_L)_{1,m_*}/(k_L)_{t,m_*}, \ dimensionless$ 

Abscissa: N'Re/(1 - € )



### EXPLANATION OF PLATE XXV

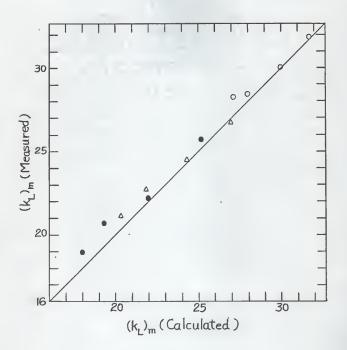
Mass-transfer coefficient calculated based on equation (48)

Compared with that experimentally measured

Ordinate: (kL)m, Measured value, lb./(hr.)(sq.ft.)

Abscissa: (k<sub>L</sub>)m, Calculated value, lb./(hr.)(sq.ft.)

### PLATE XXV



under certain operating conditions with sufficient accuracy for mass transfer calculations. But there is good indication that axial mixing in the liquid-solid systems is much less extensive than in the gas-solid systems especially with large particles such as used in this investigation. Richardson and Mitson (28) studied heat transfer in liquid-solid fluidizing systems and found that the longitudinal temperature gradient throughout the system was only slightly affected by the presence of the solid, indicating that the amount of back mixing of the liquid was negligible.

In Plates XIII, XIV, and XV, the mass transfer coefficients calculated by use of the integrated mean driving force are compared with the correlation obtained in the first series of experiments. They indicate that the deviation of the second series experimental data from the correlation are within the range of the first series of experiments. The use of the integrate of mean driving force eliminated the necessity of F-factor.

### CONCLUSION AND SUMMARY

The following significant conclusions could be drawn from the results of the present investigation:

(1) The segregation of the solid particles into two distinct sections, the fixed bed (upper section) and the fluidized bed (lower section) is caused within a single vessel containing the solid particles by denying its full expansion. In other words, it is possible to carry out the fixed bed and fluidized operations simultaneously within a single vessel.

(2) The depth of the packed section, and consequently also the fluidized section, are functions of the flow conditions, particle and fluid characteristics, and the expansion of the bed allowed.

The ratio of the depth of the fixed bed to the depth of the fluidized section can be fixed arbitrarily under certain flow conditions by controlling the expansion of the bed with the upper sieve plate.

- (3) The dimensionless equation which correlates the overall mass transfer coefficient of semi-fluidized beds and which is applicable to ordinary packed and fluidized beds was determined.
- (4) The rate of mass transfer is also affected, not only by the characteristics of the particles, fluide, and flow rate, but also by the amount of expansion of the bed allowed. The magnitudes of mass transfer coefficients can be controlled in an inverse proportionality to and within the limite of completely fixed bed and a fully fluidized bed by means of bed expansion alone.
- (5) The overall mass transfer coefficient of semi-fluidized beds can be evaluated as the linear combination of effective packed bed and fluidized bed mass transfer coefficients.

The expansion of the bed can be introduced as an independent factor, in addition to such factors as temperature, pressure, and flow rate, in controlling the rate of mass transfer for any solid particles-fluid system.

The great possibility of the practical application of this new operation of semi-fluidization is beyond the need of much description from the fact that both fixed beds and fluidized beds are extensively employed in industry at catalystic reactors, ion exchange columns, heat exchangers, driers, solvent extractors, and other process equipment. It is difficult to predict the occurrence of semi-fluidization for a large diameter column (of the order of 2-3 ft.) based on the evidence from the small laboratory scale column. However, it is believed that such phenomena are quite possible in a particular fluidization, but may not be possible in an aggregative fluidization.

### TABLE OF NOMENCIATURE

- A = total particle surface area, sq. ft.
- C = concentration of benzoic acid in water stream, lb/lb
- CS = concentration of benzoic acid in saturated solution, lb/lb
- Dp = equivalent spherical particle diameter, ft.
- D<sub>v</sub> = diffusivity of benzoic acid in water, (sq.ft.)/hr
- F = correction factor, dimensionlees
- G = superficial mass velocity, lb/(hr)(sq.ft.)(unit C)
- G' = mass velocity, lb/hr
- h = height of the top sieve plate or overall depth of bed, ft.
- ho = depth of height of initial bed, ft
- hf = height of fluidized bed section, ft
- hpa = height of packed bed section, ft
- Jd = mass transfer factor, dimensionless
- J'd = modified mass transfer factor, dimensionless
- kL = mass transfer coefficient, lb/(hr)(sq.ft.)(unit C)
- N'Gr = graph of number for mass transfer, dimensionless
- N'Re = modified Reynolde number, dimensionless
- NSc = Schmidt number, dimensionlese
- NSh = Sherwood number, dimensionless
- R bed expansion ratio, dimensionless
- S = cross-eection area of the column, sq. ft.
- T = temperature, °C
- V = overall volume of the bed, cu.ft.
- W = weight of the particle bed, lb.
- X = weight fraction of particles in the packed bed eection, dimensionlese

### Greek Letters

 $\mathcal{E}$  = void fraction or porosity in the bed, dimensionlese

 $\rho$  = fluid density, lb/cu. ft.

 $P_{\rm B}$  = bulk deneity of the solid, lb/cu. ft.

 $P_S = \text{solid density, lb/cu. ft.}$ 

μ = viscosity, lb/(hr)(ft)

 $\phi$  = particle shape factor, dimensionless

### Subscript

f = fluidized bed

p = particle

pa = packed bed

0 = at bed inlet

1 - at interface between packed and fluidized beds

2 = at bed outlet

m = mean value

1.m. = logarithmic mean value

t.m. = integrated mean value

### ACKNOWLEDGMEN T

The author takes this opportunity to express his appreciation to: Dr. Henry T. Ward, Head of the Department, for his advice and encouragement.

Dr. L. T. Fan, who originated this study, for his interest and guidance during the work. This appreciation is acknowledged with utmost sincerity for his enthusiastic spirit and wide scope of knowledge in this field have been major factors in the successful fulfillment of this contribution.

The author also wishes to express his gratitude to the Kansas State University Engineering Experiment Station for the financial support.

### BIBLIOGRAPHY

- Carman, P. C.
   Soc. Chem. Ind. <u>57</u>, 225 (1938)
- Chang, S.
   "Determination of Diffusion Coefficients in Aqueous Solutions"
   M. S. Thesis, Chem. Engg., M.I.T. (1948)
- Chilton, T. H. and A. P. Colburn Ind. Eng. Chem., 26, 1183, (1934)
- 4. Chu, J. C., J. Kalil, and W. A. Wettereth Chem. Eng. Progr. <u>49</u>, 141, (1953)
- Eckert, E. R. G.
   'Introduction to Heat and Mass Transfer',
   McGraw-Hill Book Co. Inc., New York, 1950
- Epstein, N. The Canadian Journal of Chem. Eng., 36, 210, (1958)
- Ergun, E.
   Chem. Eng. Progr., <u>18</u>, 228 (1952)
- 8. Evans, G. C. and C. F. Gerald Chem. Eng. Progr. <u>49</u>, 135 (1953)
- Fan, L. T. Unpublished note on Semi-fluidization (1953)
- Fan, L. T., Y. C. Yang and C. Y. Wen Communication submitted to A.I.Ch.E. Journal (Accepted for publication (1959)
- 11. Frossling, N. Gerl. Beit. Zur. Geophysik., <u>52</u>, 170 (1938)
- Fuchs, N.
   J. Phys. (U. S. S. R.), 6, 224 (1934)
- 13. Gaffney, B. J. and T. B. Drew Ind. Eng. Chem. <u>42</u>, 1120 (1950)
- Galloway, L. R., W. Komarnicky, and N. Epstein The Canadian Journal of Chem. Eng., 35, 139 (1957)
- Gamson, B. W., G. Thodos, and O. A. Hougen Trans. Am. Inst. Chem. Eng., 39, 1, (1943)

- 16. Garner, F. H. and R. D. Suckling A.I.Ch.E. Journal, Vol. 4, No. 1, 115 (1958)
- 17. Hobson, M. and G. Thodos Chem. Eng. Progr., <u>45</u>, 517 (1949)
- 18. Hurt, D. M. Ind. Eng. Chem., 35, 522, (1943)
- 19. International Critical table
  McGraw-Hill Book Co. Inc. (1926)
- Ishine, T., T. Otake, and T. Okada Chem. Eng. (Japan), <u>15</u>, 255 (1951)
- Lemlich, R. and I. Caldes
   A.I.Ch.E. Journal, Vol. 4, No. 3, 376 (1958)
- 22. Linten, W. H. Jr. and T. K. Sherwood Chem. Eng. Progr. 46, 259 (19)
- 23. McCune, L. K. and R. H. Welhelm Ind. Eng. Chem. <u>11</u>, 1121 (1919)
- 24. Morse, R. D. Ind. Eng. Chem. <u>41</u>, 1117, (1949)
- 25. Othmer, D. F. "Fluidization" Reinhold Publishing Corporation, New York, (1956)
- 26. Perry, J. H. Chemical Engineers handbook, 3rd edition New York, McGraw-Hill Co. (1950)
- 27. Resnick, W. and R. R. White Chem. Eng. Progr., 45, 377 (1949)
- 28. Richerdson, J. F. and A. E. Mitson Trans. Inst. Chem. Eng. 36, 270 (1958)
- Seidell, A.
   'Solubility of Organic Compounds' 3rd Ed., New York, D. Van Nostrand Co. (1941)
- Shirai, T.
   Doctor thesis, Tokyo Inst. Tech. (1955)
- 31. Taecker, R. G. and O. A. Hougen Chem. Eng. Progr., <u>45</u>, 188 (1949)
- Wakao, N., Oshima and Yagi
   Chem. Eng. (Japan), <u>22</u>, 24 (1958)

- 33. Wakao, N., Oshima and Yagi Chem. Eng. (Japan), <u>22</u>, 30 (1958)
- 34. Wilke, C. R. and O. A. Hougen Trans. Am. Inst. Chem. Eng., <u>41</u>, 445 (1945)

APPENDIX

### EXPLANATION OF PLATE XXVI

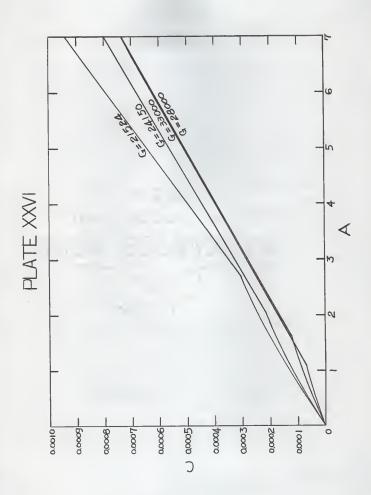
Concentration gradient in a semi-fluidized bed of benzoic acid-

water system

Particle size: 8-10 mesh

Ordinate: C, Concentration, lb./lb.

Abscissa: A, Total particle surface, sq. ft.



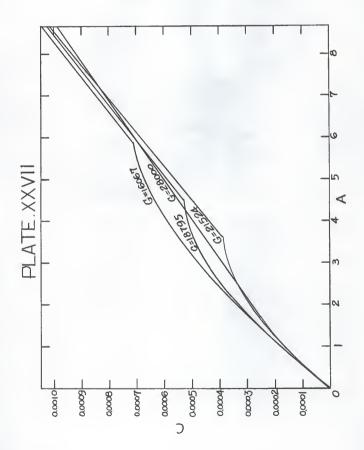
## EXPLANATION OF PLATE XXVII

Concentration gradient in semi-fluidized bed of benzoic acid-water system

Particle size: 12-14 mesh

Ordinate: C, Concentration, lb./lb.

Abscissa: A, Total particle surface, sq. ft.



# EXPLANATION OF PLATE XXVIII

Concentration gradient in semi-fluidized bed of benzoic

acid-water system

Particle size: 20-24 mesh

Ordinate: C, Concentration, lb./lb.

Abscissa: A, Total particle surface, sq. ft.

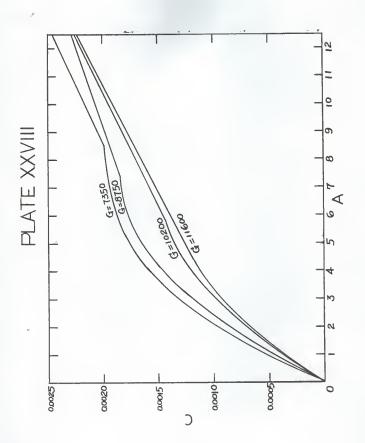


Table 2. Calculation of first series of semi-fluidized data.

	T	W	ho	h	hf
Run I	No. Temperature	Weight of	Height of	Overall	Height of Fluid-
		Particle Bed	Initial Bed	Depth of Bed	ized Bed Section
	°C	, 1b.	ft.	ft.	ft.
A-1	22.3	0.4916	0.5888	0.8091	0.4008
2	20.0	0.4916	0.5888	0.8091	0.3508
	20.0	0.4916	0.5888	0.8091	0.3258
3 4 5 6 7 8	20.0	0.4916	0.5888	0.8091	0.3758
4	21.2	0.4679	0.5603	0.8091	0.14125
6	21.5	0.4916	0.5888	1.2300	1.0925
2	21.5	0.4916	0.5888	1.2300	0.9925
ή (	20.1	0.2645	0.3168	None	0.9508
0			0.3168	0.8258	0.7091
9 10	19.0	0.2645 0.2645	0.3168	0.8258	0.7841
	17.0				
11	16.8	0.2645	0.3168	0.5008	0.2341
12	17.0	0.2645	0.3168	0.5008	0.2550
13	19.4	0.2645	0.3168	0.5008	0.2758
14	19.8	0.2645	0.3168	0.5008	0.3050
15	22.1	0.2645	0.3168	0.6675	0.5383
16	24.0	0.2645	0.3168	0.4350	0.1508
17	24.0	0.2645	0.3168	0.4350	0.1425
18	24.0	0.2645	0.3168	0.4350	0.1250
19	20.5	0.2645	0.3168	0.5008	0.3633
20	20.5	0.2645	0.3168	0.5008	0.2008
21	24.0	0.2645	0.3168	0.6617	0.4633
22	24.0	0.2645	0.3168	0.6617	0.3966
23	20.0	0.2645	0.3168	0.8258	0.6925
24	20.0	0.2645	0.3168	0.8258	0.5842
25	20.0	0.4864	0.5825	0.8091	0.3591
B1	21.00	0.2533	0.2999	0.5008	0.3342
2	21.20	0.2533	0.2999	0.5008	0.2592
3	22.10	0.2533	0.2999	0.6633	0.5842
3 4 5 6 7 8 9	22.40	0.2533	0.2999	0.6633	0.5342
5	23.00	0.2533	0.2999	0.6633	0.5008
6	18.00	0.2646	0.3131	0.4783	1.4425
7	23.00	0.2646	0.3131	0.6258	0.4550
8	23.00	0.2646	0.3131	0.6258	0.4258
9	23.00	0.2646	0.3131	0.6258	0.4008
10	21.30	0.2646	0.3131	0.6258	0.3591
11	21.90	0.2646	0.3131	0.6633	0.4091
12	21.90	0.2646	0.3131	0.5225	0.2341
13	20.50	0.2646	0.3131	0.5225	0.2967
14	21.00	0.2493	0.2952	0.5008	0.3375
15	21.20	0.2487	0.2945	0.5008	0.2592
16	21.20	0.2441	0.2890	0.5008	0.2675

C-1	22.60	0.2645	0.3093	0.4675	0.2550
2	22.60	0.2645 0.2645	0.3093	0.4675	0.2300 0.2175
۶	22.60		0.3093	0.4675	
4	22.60	0.2645	0.3093	0.4675	0.1675
5	19.80	0.2645	0.3093	0.6342	0.5342
6	19.80	0.2645	0.3093	0.6342	0.5092
7	19.80	0.2645	0.3093	0.6342	0.4675
8	19.80	0.2645	0.3093	0.6342	0.4342
9	22.60	0.2585	0.3022	0.4675	0.2633
10	22.60	0.2572	0.3007	0.4675	0.2383
11	22.60	0.2566	0.3000	0.4675	0.2258
12	22.60	0.2552	0.2983	0.4675	0.1800
13	19.80	0.2576	0.3011	0.6342	0.5425
14	19.80	0.2570	0.3004	0.6342	0.5175
15	19.80	0.2559	0.2991	0.6342	0.4758
16	19.80	0.2556	0.2988	0.6342	0.4425

h <sub>pa</sub>	X	R	€	Dp	M-1-2 P11-2-
neight of rack-	Weight Fraction				
ed Bed Section	in Packed Bed	sion Ratio		Diameter ft.	Surface Area ft2
ft.		el ca	48.49	16.	172
-0-4083	0.6935	3 2720	0.6548	0.00690	10.40
0.4583	0.7785	1.3732 1.3732	0.6548	0.00690	10.42
0.4833	0.8208	1.3732	0.6548	0.00690	10.39
0.4333	0.7360	1.3732	0.6548	0.00690	10.40
0.3666	0.65h2	1.4440	0.6717	0.00682	10.15
0.3000	0.0342	2.0902	0.7732	0.00690	10.41
0.1375	0.4034	2.0902	0.7732	0.00690	10.38
0.2373	0.4034	2.0702	0.1152	0.00690	5.60
0.1167	0.3683	2.6078	0.8182	0.00689	5.59
0.0417	0.1316	2.6078	0.8182	0.00690	5.60
			0.7002	0.00691	5.63
0.2667	0.8420	1.5815		0.00689	5.57
0.2458	0.7762	1.5815	0.7002	0.00689	5.58
0.2250	0.7104 0.6182	1.5815	0.7002 0.7002	0.00689	5.58
0.1958		1.5815		0.00689	5 - 59
0.1292	0.4080	2.1078	0.7751	0.00688	5 • 59 5 • 57
0.2842	0.8972	1.3732	0.6548	0.00689	5.52
0.2925	0.9233	1.3732	0.6548		
0.3100	0.9785	1.3732	0.6548	0.00685	5.53
0.1375	0.1311	1.5809	0.7002	0.00690	5.60 5.58
0.3000	0.9471	1.5809	0.7002		
0.1984	0.6265	2.0888	0.7731	0.00688	5.56
0.2651	0.8371	2.0888	0.7731	0.00686	5-53
0.1333	0.4207	2.6078	0.8182	0.00690	5 - 59
0.2416	0.7625	2.6078	0.8182	0.00688	5.57
0.4500	0.7725	1.3890	0.6587	0.00688	10.42
0.1666	0.5555	1.6700	0.7129	0.00453	1 7.62
0.2416	0.8054	1.6700	0.7129	0.00453	
0.0791	0.2637	2.2118	0.7832	0.00453	
0.1291	0.4303	2.2118	0.7832	0.00452	
0.1625	0.5416	2.2118	0.7832	0.004520	
0.0358	0.1148	4.7218	0.8985	0.00452	
0.0330	0.5454	1.9987	1.7601	0.00453	
0.2000	0.6383	1.9987	0.7601	0.00452	
0.2250	0.7180	1.9987	0.7601	0.00451	
0.2667	0.8510	1.9987	0.7601	0.00451	
0.2542	0.8114	2.1185	0.7737	0.004509	
0.2884	0.9206	1.6688	0.7127	0.00450	
0.2358	0.7207	1.6688	0.7127	0.00450	
0.1633	0.5532	1.6965	0.7174	0.00452	7.53
0.1633	0.8203	1.7005	0.7174	0.00451	7.50
0.2333	0.8071	1.7328	0.7233	0.001178	7.4L
0.2)))	0.0017	T. 1250	0.12)	о воотщо	1044

0.2125	0.6873	1.5114	0.6787	0.002610	12.50
0.2375	0.7682	1.5114	0.6787	0.002606	12.46
0.2500	0.8083	1.5114	0.6787	0.002603	12.44
0.3000	0.9701	1.5114	0.6787	0.002598	12.39
0.1000	0.3232	2.0504	0.7632	0.002609	12.47
0.1250	0.4041	2.0504	0.7632	0.002604	12.46
0.1667	0.5391	2.0504	0.7632	0.002600	12.42
0.2000	0.6469	2.0504	0.7632	0.002599	12.40
0.2042	0.6758	1.5470	0.6861	0.002602	12.38
0.2292	0.7624	1.5547	0.6876	0.002591	12.30
0.2417	0.8059	1.5583	0.6884	0.002585	12.25
0.2875	0.9636	1.5672	0.6901	0.002572	12.17
0.0917	0.3044	2.1062	0.7694	0.002592	12.32
0.1167	0.3883	2.1111	0.7700	0.002592	12.27
0.1584	0.5295	2.1203	0.7710	0.002580	12.20
0.1917	0.6416	2.1224	0.7712	0.002578	12.18

G	G1	Ca	C <sub>2</sub>	(kr)	NSc
Superficial	Mass	Saturated	Outlet	(k <sub>L</sub> ) <sub>l.m.</sub> Mass Transfer	Schmidt
				Mass Hallster	
mass velocity		Concentration	Concentration	Coefficient	Number
$lb/(hr)(ft^2)$	lb/hr	<b>l</b> b/lb	1b/1b	$1b/(hr)(ft^2)(\triangle t)$	3)
16067	350.52	0.003140	0.001640	24.90	1120
13338	290.98	0.002902	0.001618	22.77	1290
18795	410.03	0.002902	0.001483	28.32	1290
16067	350.52	0.002902	0.001587	26.62	1290
13338	290.98	0.003025	0.001436	17.58	1200
21524	496.57	0.003057	0.001158	21.48	1180
24150					
	529.04	0.003057	0.001189	24.89	1180
30400	663.21	0.002907	0.000446	19.71	1290
33000	719.93	0.002803	0.000496	25.10	1360
28000	610.85	0.002619	0.000131	19.88	1530
33000	719.93	0.002600	0.000628	34.72	1540
28000					
	610.85	0.002619	0.0006717	30.92	1530
24150	496.57	0.002842	0.000700	23.77	1335
21504	528.04	0.002878	0.000758	28.87	1300
28000	610.85	0.003121	0.000676	26.64	11110
24150	469.57	0.003336	0.000952	27.29	1020
31000	676.30	0.003336	0.000949	41.03	1020
39600	863.92	0.003336	0.000882	47.98	1020
18795	410.03	0.002907	0.000675	19.31	1250
39600	863.91	0.002907	0.000550	32.46	1250
31000	676.30	0.003336	0.000799	33.31	1020
39600	863.91				
		0.003336	0.000835	44.71	1020
31000	676.30	0.002902	0.000488	22.29	1290
39600	863.91	0.002902	0.000570	33.84	1290
13338	290.98	0.002902	0.001551	21.51	1290
-,,,,,	2,00,0	00002/02	240040		12/0
10610	027 15	0.000007	0.0027/0		
	231.47	0.003005	0.001563	22.30	1220
13338	290.98	0.003024	0.001437	28.42	1200
16067	350.52	0.003121	0.001137	20.89	1140
18795	410.03	0.003154	0.001127	24.04	1120
21524	469.57	0.003223	0.001197	28.81	1080
30400	663.20				
		0.002709	0.000635	23.59	1440
18795	110.03	0.003223	0.001399	29.63	1080
21524	469.57	0.003223	0.001380	33.25	1080
2117.20	529.04	0.003223	0.001380	37.55	1080
28000	610.85	0.003036	0.001286	42.69	1200
28000	610.85	0.003100	0.001260	40.35	1160
28000	610.85	0.003100	0.001395	46.56	11.60
21524	469.57	0.002953	0.001209	31.17	1250
10610	231.47	0.003005	0.001508	21.37	1220
13338	290.98	0.003024	0.001437	28.79	1200
13338	290.98	0.003024	0.001408		1200
∪ررر⊥	£70.70	0.000024	O.OOTTOO	24.93	1200

4500	98.17	0.003177	0.003051	25.38	1110
6450	140.71	0.003177	0.002968	29.98	1110
7700	167.98	0.003177	0.002812	27.58	1110
10000	218.16	0.003177	0.002719	31.38	1110
73 <i>5</i> 0	160.35	0.002882	0.002387	20.2և	1300
8750	190.89	0.002882	0.002267	20.00	1300
10200	222.52	0.002882	0.002402	28.8և	1300
11600	253.06	0.002882	0.002226	25.08	1300
4500	98.17	0.003177	0.002968	21.05	1110
6450	140.71	0.003177	0.002861	24.48	1110
7700	167.98	0.003177	0.002802	27.58	1110
10000	218.16	0.003177	0.002657	29.26	1110
7350	160.35	0.002882	0.002350	19.35	1300
8750	190.89	0.002882	0.002257	19.97	1300
10200	222.52	0.002882	0.002361	27.57	1300
11600	253.06	0.002882	0.002200	24.30	1300

NSh	N Re	N¹Re/(1.€) Dimensionless	J <sub>d</sub> Mass Transfer	$(N_{Sh} \in -2)/N_{Sc}^{1/2}$	J'd Modified Mass
Sherwood	Modified			Dimensionless	Modified Mass
Number	Rsynolds Number	Group	Factor	Group	Transfer Factor
		app may stall	400	400	40 40 40
() 10	10.00	710.0	0.1670	3.87	0.1245
64.48	48.33	140.0	0.1670	3.66	0.1500
63.84	37.89	109.7	0.2023 0.1785	4.60	0.1333
79.42	53.36	154.5	0.1964	4.31	0.1454
74.64	45.93	133.0	0.1525	2.73	0.1084
46.46	38.14	115.7	0.1115	3.66	0.0753
57.34 66.41	63.48	280.0 315.4	0.1115	4.67	0.0753
	71.52	546.7	0.0768	4.08	0.0574
54.36	86.33	502.4	0.0933	5.18	0.0698
72.59 61.80	91.30	470.3	0.0913	h.22	0.0607
	85.47	288. h	0.1403	6.43	0.1070
108.88	86.57		0.1466	5.66	0.0955
96.00	85.35	284.8		ь.32	0.0955
67.85	67.41	224.8	0.1193	4•32 5•22	0.1295
81.36	60.67	202.3	0.1597	5.22 5.97	0.1295
69.55	83.21	370.2	0.1003		0.0778
66.55	75.11	217.6	0.1145	4.13	
100.20	96.56	279 • 7	0.1342	6.32	0.1030
116.50	122.63	355.2	0.1228	7.38	0.0935
53.27	53.97	180.0	0.1192	3.28	0.0884
89.29	113.38	378.1	0.0965	5.62	0.0721
81.24	96.42	424.9	0.1089	6.04	0.0818
108.72	122.81	541.2	0.1144	8.16	0.0864
62.51	87.81	483.0	0.0852	4.52	0.0686
94.63	111.85	615.2	0.1012	6.98	0.0765
60.14	37•73	110.50	0.1911	3.64	0.1416
39.74	20.25	70.53	0.2400	2.46	0.1745
50.29	25.58	89.09	0.2405	3.18	0.1778
35.88	31.42	144.92	0.1423	2.49	0.1030
40.77	37.06	170.94	0.1380	2.88	0.1002
	12.96	198.15	0.1409	3.45	0.1038
47.75 46.47	53.92	531.23	0.1409	3.52	0.0730
49.24	37.53	156.44	0.1660	3.16	0.1230
55.05	12.92	178.90	0.1620	h.08	0.1202
		200.62	0.1637	4.00	0.1219
62.14 74.89	48.13 53.56	223.25	0.1037	5.17	0.1219
69.43	54.32	240.03	0.1590	4.93	0.1357
80.01	54.32 54.28	188.93	0.1835	5.24	0.1370
56.36	40.51	141.00	0.1680	3.54	0.1246
37.94	20.16	71.33	0.2300	2.55	0.1669
50.60	25.10	90.07	0.2437	3.23	0.1800
43.60	25.40	91.36	0.2110	2.78	0.1547
45.00	27.20	71070	0.5770	2010	0.174

24.63 29.05	5.138 7.352	15.99 22.88	0.6045	1.61	0.4255 0.3554
26.69	8.768	27.09	0.3839	1.56	0.2721
30.31	11.365	35.37	0.3363	1.79	0.2407
21.60	7.845	33.13	0.3279	1.33	0.2290
21.30	9.322	39.37	0.2790	1.31	0.1897
30.67	10.850	45.82	0.3367	1.96	0.2421
26.66	12.334	52.09	0.2574	1.68	0.1832
20.36	5.122	16.32	0.5014	1.16	0.3463
24.96	7.310	23.40	0.4068	1.46	0.3045
26.51	8.707	27.94	0.3839	1.57	0.2720
27.98	11.251	36.31	0.3136	1.67	0.2230
20.52	7.794	33.80	0.3134	1.26	0.2178
21.17	9.279	40.34	0.2873	1.31	0.1894
29.09	10.766	47.01	0.3219	1.87	0.2306
25.62	12.235	53.47	0.2494	1.63	0.1769

Table 3. Calculation of second series of semi-fluidized bed data.

Run No	:	W Weight of bed lb.	:	ho Height of bed ft.	: : : : :	h Overall depth of bed ft.	:	hf Height of fluidized bed section ft.
D-1		0.3307		0.3613		0.5717		0.3484
D-2		0.3307		0.3613		0.5717		0.3151
<b>D-</b> 3		0.3307		0.3613		0.5717		0.2913
D-4		0.3307		0.3613		0.5717		0.2676
E-1		0.2899		0.3131		0.6667		0.5650
E-2		0.2899		0.3131		0.6667		0.5158
E-3		0.2899		0.3131		0.6667		0.4833
E-4		0.2899		0.3131		0.6667		0.4125
F-l		0.2645		0.2851		0.5833		0.4921
F-2		0.2645		0.2851		0.5833		0.4693
F-3		0.2645		0.2851		0.5833		0.4313
F→4		0.2645		0.2851		0.5833		0.4009

pa ght of ked bed tion ft.	: X : Weight fraction : in packed bed : section :	 D <sub>p</sub> Particle diameter ft.	 Total : particle : surface : sq.ft. :	€ Porosity	: S : Cross-section : Area of : Column : sq.ft.
.2233	0.6182	0.006890	6.97	0.7004	0.023916
.2566	0.7104	0.006890	6.97	0.7004	11
. 2804	0.7762	0.006890	6.97	0.7004	II
-3041	0.8420	0.006890	6.97	0.7004	Ħ
.1017	0.3240	0.004530	8.68	0.7740	11
.1509	0.4800	0.004523	8.65	0.7740	Ħ
.1834	0.5830	0.004520	8.64	0.7740	н
-2542	0.8110	0.004510	8.60	0.7740	н
.0912	0.3232	0.002609	12.47	0.7617	Ħ
.1140	0.4041	0.002604	12.46	0.7617	11
.1520	0.5391	0.002600	12.40	0.7617	Ħ
.1824	0.6469	0.002599	12.40	0.7617	11

Heigh	t of samp	ling outlet	: Concent:	c ration, lb/	lb. at	: : :	C <sub>1</sub> Conc. at interface lb/lb		Conc. at outlet lb/lb	
a	: b	: c	: a	b		:	10/ 10			
0.2033	0.3700	0.5717	0.000215	0.000359	0.000913		0.00030		0.000943	
0.2033	0.3700	0.5717	0.000161	0.000287	0.000862		0.00021		0.000862	
0.2033	0.3700	0.5717	0.000107	0.000251	0.000799		0.00012		0.000799	
0.2033	0.3700	0.5717	0.000099	0.000242	0.000745		0.00007		0.000745	
0.3700	0.5367	0.6667	0.000585	0.000690	0.0010ग्रि		0.00071		0.001014	
0.3700	0.5367	0.6667	0.000456	0.000591	0.001027		0.00053		0.001027	
0.3700	0.5367	0.6667	0.000363	0.000580	0.001017		0.00039		0.001017	
0.3700	0.5367	0.6667	0.0002111	0.000623	0.000992		0.00025		0.000992	
0.2033	0.3700	0.5833	0.001428	0.001921	0.002460		0.00198		0.002460	
0.2033	0.3700	0.5833	0.001383	0.001886	0.002280		0.00184		0.002280	
0.2033	0.3700	0.5833	0.001033	0.001364	0.002263		0.00142		0.002263	
0.2033	0.3700	0.5833	0.000970	0.001176	0.002245		0.00118		0.002245	

Saturated concentration		(AC) <sub>1.m.</sub>					( \( C \) t.m.				
1b/1b		ed bed :	Fluidized bed section	:	Overall	:	Packed bed section	:	Fluidized bed section	: Overall	
0.002902	0.00	2271	0.002752		0.002405		0.002380		0.002752	0.002460	
0.002902	0.00	2385	0.002763		0.002480		0.002397		0.002792	0.002511	
0.002902	0.00	2470	0.002842		0.002505		0.002474		0.002842	0.002556	
0.002902	0.00	2466	0.002862		0.002515		0.002499		0.002862	0.002651	
0.002953	0.00	2640	0.002590		0.002367		0.002076		0.002507	0.002556	
0.002953	0.00	2240	0.002780		0.002414		0.002272		0.002703	0.002567	
0.002953	0.00	2240	0.002760		0.002428		0.002224		0.002743	0.002722	
0.002953	0.00	2270	0.002790		0.002439		0.002260		0.002731	0.002725	
0.003223	0.00	09.83	0.002077		0.001709		0.001003		0.001943	0.001639	
0.003223	0.00	1160	0.002175		0.001846		0.001171		0.002010	0.001671	
0.003223	0.00	1329	0.002435		0.001854		0.001371		0.002347	0.001821	
0.003223	0.00	1445	0.002582		0.001880		0.001510		0.002453	0.001843	

G : Mass : velocity :	G: Mass velocity		(k <sub>L</sub> )1.m.		:	(kL)t.m.	
1b/(hr)(sq.ft.)		Packed bed section	:Fluidized : bed : section	: Overall	Packed bed section	:Fluidized : bed : section	Overal
21524	514.77	30.83	21.08	28.97	28.68	21.08	28.32
24150	577•57	30.87	21.75	28.81	30.83	21.52	28.45
28000	669.65	33.83	18.14	30.66	33.70	18.14	30.04
33000	789.22	36.32	17.53	33.56	35.85	17.53	31.83
16067	384.26	30.10	13.28	19.57	20.70	13.71	18.08
18795	449.59	22.25	17.85	22.10	21.90	18.40	20.79
21524	514.77	26.30	22.70	24.93	26.50	22.40	22.24
28000	669.65	28.20	33.00	31.67	28.30	24.30	25.35
7350	175.78	21.31	19.85	20.29	21.88	20.88	21.15
8750	209.26	18.15	23.82	20.74	23.77	20.71	22.91
10200	243.94	23.49	24.74	24.98	24.74	22.77	24.41
11600	277.42	25.33	28.82	26.24	28.52	24.25	26.77

N'Re Reynold	: s:D:	N'Re/(1-€ imensionles	8 Sherwood	Sh number	Mass-transfe	d er coefficient
	*	group			ean: Log. mean	:Integrated mean driving force
60.88		203.30	81.00	79 • 3	0.1588	0.1552
68.31		228.00	80.6	79.6	0.1407	0.1390
79.20		264.35	85.8	84.1	0.1292	0.1265
93.34		311.54	94.0	89.0	0.1199	0.1137
31.40		139.00	35.4	32.7	0.1413	0.1305
35.50		159.00	40.0	37.6	0.1363	0.1283
40.50		179.10	45.1	40.3	0.1343	0.1198
52.60		233.00	57.3	51.3	0.1312	0.1174
8.47		35.54	19.4	20.2	0.2907	0.3030
10.06		42.21	22.9	21.9	0.2495	0.2756
11.71		49.14	19.8	23.3	0.2475	0.2519
13.31		55.85	25.1	25.6	0.2381	0.2430

N <sub>Sc</sub> Schmidt number		Dimens	-2)/NSc1/3 sionless	: J'd :Modified mass-transfer coefficien :					
	: Log.	mean g force	:Integrated mean :driving force	: Log. m		driving force			
1290	5.0	03	4.91	0.1	192	0.1167			
1290	5.0	00	4.94	0.1	1057	0.1014			
1290	5.2	34	5.23	0.0	0972	0.0953			
1290	5.8	36	5.54	0.0	906	0.0857			
1250	2.	36	3.16	0.0	987	0.0908			
1250	2.0	69	2.52	0.0	994	0.0931			
1250	3.0	05	2.71	0.0	988	0.0878			
1250	3.5	94	3.50	0.0	976	0.0870			
1080	1.	25	1.30	0.2	2004	0.2097			
1080	1.2	27	1.43	0.1	1726	0.1930			
1080	1.	51	1.54	0.1	1742	0.1775			
1080	1.0	57	1.71	0.1	1693	0.1693			

Table 4. Results of recalculation of Evan and Gerald's fluidized bed data (8).

Run No.	: Temperature	: Weight of : Particle Bed : lb.	: hf : Height of :Fluidized Bed : ft.	: D <sub>p</sub> :Particle :Diameter : ft.	Porosity
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24	21.00 24.05 23.41 21.70 23.40 23.40 23.05 23.05 23.05 23.35 23.35 23.35 23.37 23.65 23.75 23.65 23.75 23.65 23.75 23.81 23.70 22.07 23.61 23.29 24.73 24.20 24.07 23.40	0.1012 0.1009 0.1050 0.1051 0.1021 0.1039 0.1061 0.0601 0.0616 0.0640 0.0376 0.0569 0.0572 0.0583 0.0583 0.0589 0.0565 0.0492 0.0961 0.0966 0.0875 0.0965	0.1 lpl 0.1 sl 0.1 sl 0.1 sl 0.1 sl 0.200 0.272 0.295 0.121 0.131 0.108 0.1 lpl 0.095 0.193 0.262 0.082 0.082 0.082 0.082 0.182 0.226 0.226 0.226 0.226 0.226 0.226	0.00660 0.00660 0.00678 0.00678 0.00678 0.00678 0.00134 0.00134 0.001392 0.00126 0.00219 0.00219 0.00219 0.00216 0.00212 0.00212 0.00213	0.56 0.59 0.67 0.69 0.77 0.78 0.70 0.71 0.69 0.75 0.82 0.87 0.56 0.56 0.56 0.56 0.59

G	_	Cs	Co	-	()-1)-	, No	=	N
Superficial	:	Saturated	: C <sub>2</sub>		(kL) <sub>1.m</sub> . Mass Transfer	: NSc :Schmidt	:	N <sub>Sh</sub> Sherwood
Mass Velocity		Concentration	: Concentration		Coefficient	Number	i	Number
1b/(hr)(ft2)		1b/1b	: lb/lb	:	1b/(hr)(ft2)(A			
3300	Ť	0.00328	0.001120	÷	12.6	1020	÷	29.42
8650		0.00328	0.000594		15.9	1020		37.21
11420		0.00321	0.000462		15.9	1060		38.98
15200		0.00373	0.000405		15.7	385		36.20
21870		0.00373	0.000316		16.8	385		38.84
23 750		0.00321	0.000274		18.9	1060		46.35
8320		0.00321	0.000552		16.2	1060		25.42
11150		0.00313	0.000432		17.6	1100		28.60
11420		0.00317	0.000467		18.9	1080		30.63
13590		0.00323	0.000380		18.6	1050		29.02
13600		0.00320	0.000290		21.8	1060		30.98
19600		0.00320	0.000268		20.2	1060		31.19
23000		0.00321	0.000269		23.4	1060		36.01
1381		0.00324	0.002200		11.4	1040		10.17
1386		0.00325	0.002068		10.0	1040		8.86
3020		0.00326	0.001460		13.3	1030		11.61
3890		0.00324	0.001290		15.6	1040		11.66
5840		0.00315	0.001531		17.2	1100		15.77
8380		0.00324	0.001303		20.2	1040		17.85
11510		0.00320	0.001040		20.6	1070		18.08
14950		0.00337	0.000930		20.2	390		16.58
17490		0.00330	0.000796		23.5	1015		19.99
3400		0.00318	0.001739		14.2	1085		9.30
6620		0.00321	0.000704		17.7	1060		11.65
8320		0.00321	0.000589		17.7	1060		11.52

N'Re Modified Reynolds Number	: N¹Re : 1 -€ : Dimensionless : Group :	Jd : Mass Transfer : Factor :	N <sub>sh</sub> ∈ -2 N <sub>sc</sub> 1/3 Dimensionless Group	J'd : Modified Mass : Transfer Factor :
9.80 25.80 34.60 46.50 67.20 71.6 16.0 21.3 22.2 26.2 23.6 37.0 43.4 1.53 1.53 3.32 3.62 6.40	22.27 62.92 101.81 150.00 292.17 325.45 53.33 73.44 71.61 101.80 98.33 205.50 333.84 3.59 3.47 8.09 9.78	0.390 0.188 0.146 0.102 0.076 0.084 0.204 0.170 0.175 0.142 0.168 0.106 0.855 0.745 0.453 0.414 0.314	1.43 1.98 2.36 2.30 2.80 3.34 1.54 1.77 1.86 1.94 2.10 2.30 2.87 0.36 0.29 0.48 0.53 0.81	0.276 0.135 0.101 0.071 0.055 0.061 0.113 0.120 0.125 0.105 0.120 0.077 0.077 0.505 0.137 0.286 0.262 0.208
9.20 12.3 16.3 19.1 2.69 5.34 6.66	36.80 68.33 135.8 212.2 8.15 20.54 74.00	0.250 0.188 0.134 0.136 0.142 0.280 0.222	1.12 1.25 1.26 1.61 0.41 0.65 0.83	0.169 0.127 0.090 0.093 0.264 0.176 0.139

Table 5. Results of recalculation of Evan and Gerald's packed bed data (8).

Run No. :	T Temperature	W Weight of Particle Bed lb.	h <sub>pa</sub> Height of Fluidized Bed ft.	D <sub>p</sub> Particle Diameter ft.	Porosity
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18	23.49 23.52 23.60 23.40 23.60 23.70 23.30 23.10 23.91 23.60 23.40 23.50 23.40 23.40 23.61 23.61 23.61	0.1039 0.1038 0.1060 0.1033 0.1010 0.0619 0.0629 0.0596 0.0512 0.0651 0.0610 0.1058 0.0596 0.0590 0.0479 0.0621 0.0853	0.138 0.130 0.132 0.131 0.122 0.079 0.081 0.083 0.085 0.085 0.082 0.079 0.076 0.129 0.071 0.076 0.062	0.00676 0.00667 0.00673 0.00673 0.00664 0.00433 0.00440 0.00435 0.00431 0.00436 0.00249 0.00256 0.00249 0.00230 0.00183	0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.0

Superficial Mass Velocity lb/(hr)(ft <sup>2</sup> )	Cs Saturated Concentration b/lb	C2 Outlet Concentration lb/lb	(kL) <sub>1.m</sub> . Mass Transfer Coefficient lb/(hr)(ft <sup>2</sup> )(\(\triangle C)	NSc Schmidt Number	NSh Sherwood Number
2345 2870 8660 15220 19410 1386 1843 4205 4625 17780 18620 22400 1669 14000 16680 17780 1148	0.00322 0.00323 0.00323 0.00321 0.00323 0.00324 0.00320 0.00318 0.00327 0.00324 0.00321 0.00322 0.00315 0.00322 0.00315	0.001475 0.001280 0.000805 0.000588 0.000562 0.001820 0.001665 0.00119 0.000950 0.000585 0.002945 0.00131 0.002990 0.002830	12.8 13.1 21.6 27.2 35.2 12.5 15.8 19.0 18.0 18.1 148.0 18.1 52.5 55.5 54.6 17.4 24.3	1050 1050 1045 1060 1045 1040 1055 1080 1030 1045 1030 1050 1050 1050 1050	31.15 31.41 52.21 66.11 83.97 19.35 22.00 30.20 28.53 65.16 66.44 75.00 17.03 47.28 49.78 45.11

N'Re :	N°Re.	. I.	: N <sub>sh</sub> ∈ -2	
Modified :		: J <sub>d</sub> : Mass Transfer		: J'd
Reynolds:	$1-\epsilon$ Dimensionless		Nsc1/3	Modified Mas
		: Factor	: Dimensionless	: Transfer Facto
Number :	Group	:	: Group	\$
:			:	:
7.09	15.41	0.545	1.45	0.401
8.53	17.40	0.476	1.37	0.438
26.0	53.06	0,259	2.41	0.189
45.5	94.79	0.187	3.17	0.138
57.5	117.34	0.190	3.99	0.139
2.68	5.70	0.942	0.81	0.638
3.58	7.30	0.785	0.90	0.545
8.09	17.58	0.478	1.39	0.339
8.74	20.80	0.415	1.41	0.297
33.0	86.84	0.260	3.76	0.187
37.0	72.55	0.230	2.99	0.171
43.2	86.40	0.23	3.48	9.165
1.87	3.74	1.16	0.63	
15.5				0.772
18.5	30.39	0.392	2.07	0.288
	38.54	0.348	2.33	0.252
18.1	36.20	0.318	2.02	0.234
0.92	1.91	1.60	0.39	1.020
2.48	4.67	0.820	0.53	0.505

## MASS TRANSFER IN SEMI\_FLUIDIZED BEDS

by

Yung Chia Yang

B. S., National Taiwan University, China, 1955

AN ABSTRACT OF A THESIS

submitted in partial fulfillment of the

requirements for the degree

MASTER OF SCIENCE

Department of Chemical Engineering

KANSAS STATE UNIVERSITY
OF AGRICULTURE AND APPLIED SCIENCE

Semi-fluidization is a new fluid-solid contact process in which two distinct beds of the solid particles are formed simultaneously within a single vessel. This can be accomplished by partial restriction or prevention of the bed expansion of the fluidized bed. Two types of the solid particles beds formed in the semi-fluidized operation are a packed bed and a fluidized bed. Therefore, the semi-fluidized bed should possess the characteristics of both the packed bed and the fluidized bed. In this investigation, the mass-transfer aspect of semi-fluidization was investigated by use of the benzoic acid-water system.

Several dimensionless correlatione were obtained based on the overall mean driving force. It was found that the correlations obtained are very similar to those for the ordinary packed and fluidized beds.

As a matter of fact, these correlations can be used in calculation of the overall mass-transfer coefficients in the semi-fluidized bed for purpose of approximation. However, the porosity of the semi-fluidized bed is an independent variable of the operation unlike that of ordinary fluidized bed. In other words, the rate of mass-transfer can be controlled by means of the bed expansion only.

Some additional experiments were conducted in order to study the effect of axial mixing on the overall mass transfer correlation and also the effect of sudden change in the concentration gradient. It was found that such effecte were small.

It was also shown that the overall mean  $\mathbf{k}_{L}$  is expressed as a linear combination of effective  $\mathbf{k}_{L}$  of each bed.