

## Semifluidization: a Review

J. S. N. MURTHY AND G. K. ROY

Department of Chemical Engineering, Regional Engineering College, Rourkela 769 008

*Semifluidization is a novel solid-fluid contact technique conceived in the late fifties. Over the last two-and-a-half decades considerable investigation has been made into the various aspects of the phenomenon which have been reported in the literature. In view of its unique usefulness, semifluidization will become increasingly important, specifically in the design of several physical and chemical systems. This review presents a concise picture of the various aspects of semi-fluidization (momentum, heat and mass transfer, design of reactors and industrial applications) for a better understanding of the phenomenon and to provide more rational approaches to the design of semi-fluidization systems.*

THE phenomenon of semifluidization, a unique and novel technique of fluid-solid contact proposed by Fan, *et al.*<sup>1</sup> in the late fifties, can be viewed as a combination of a batch fluidized bed at the bottom and a fixed bed at the top. This is obtained by providing enough space available for free expansion of the bed, and increasing the flow rate of fluid through the bed of particles sufficiently high to buoy the solid particles. A restraint provided at the top of the bed prevents the escape of the particles from the system and helps to form a packed-bed section just below the restraint. A semifluidized bed overcomes certain inherent disadvantages of fluidized and fixed beds. Semifluidized beds find wide applications as exothermic chemical reactors,<sup>2</sup> ion-exchangers,<sup>3</sup> filters,<sup>3,4</sup> and for the removal of fine particles suspended in gases or liquids and bioreactors.<sup>3,5</sup>

The present article is designed to review the available literature on semifluidization under the following headings: (a) momentum transfer, (b) heat and mass transfer, (c) application to the design of reactors, and (d) miscellaneous industrial applications.

### Momentum Transfer

This part deals with the determination of (a) pressure drop, (b) minimum semifluidization velocity (onset of semifluidization velocity), (c) maximum semifluidization velocity (d) height of the packed bed and lean bed, and (e) residence-time distribution.

#### Pressure Drop

In semifluidization the total pressure drop is ideally the algebraic sum of the pressure drop across the fluidized and the packed sections. Hence,

$$\Delta P_T = \left(\frac{\Delta P}{L}\right)_f (h - h_{pa}) + \left(\frac{\Delta P}{L}\right)_{pa} h_{pa} \quad (1)$$

Through an overall mass balance Fan and co-workers<sup>1,6</sup> established that the following relationship exists between the magnitude of the space occupied by the packed bed and that by the fluidized bed in a semifluidized bed

$$h_{pa} = (h_f - h) \frac{1 - \epsilon_f}{\epsilon_f - \epsilon_{pa}} \quad (2)$$

While there is essentially only one generalised equation<sup>1,7</sup> for the prediction of the pressure drop across a fluidized bed,

$$\left(\frac{\Delta P}{L}\right)_f = (\rho_s - \rho_f) (1 - \epsilon_f) \quad (3)$$

there are various correlations for the determination of the pressure drop across a packed bed. These include the Kozeny-Carman equation,<sup>7</sup> Leva's equation,<sup>8</sup> and Ergun's equation.<sup>8</sup> The equation for pressure drop across a semifluidized bed using Ergun's equation is given by

$$(\Delta P)_T = \frac{1}{g_c} \left[ 150 \frac{(1-\epsilon_{pa})^3}{\epsilon_{pa}^3} \frac{\mu_f v_o}{d_p^2} + 1.75 \frac{(1-\epsilon_{pa})}{\epsilon_{pa}^3} \frac{G v_o}{d_p} \right] \left[ (h_t - h_s) \frac{1-\epsilon_t}{\epsilon_t - \epsilon_{pa}} \right] + \left[ h_t - \frac{(1-\epsilon_{pa})(h_t - h)}{\epsilon_t - \epsilon_{pa}} \right] (1-\epsilon_t)(\rho_s - \rho_t) \quad (4)$$

Fan and Wen<sup>9</sup> measured the pressure drop in fixed and fluidized beds separately and obtained the total pressure drop using equation (1). This was compared with that calculated using equation (4). The experimental values were nearer to those calculated using equation (1) whereas equation (4) gave lower values. For small and irregular-shaped particles, Kurian and Raja Rao<sup>10</sup> observed that the pressure drop was greater than that given by equation (4), and the following equation was proposed to take into account this additional pressure drop

$$\Delta P_a = 2.10 \times 10^{-3} G_{st}^{1.56} d_p^{-0.94} h_{pa}^{0.59} \quad (5)$$

To overcome wide discrepancies between the experimental and the calculated values, Roy and Sen Gupta<sup>11</sup> and Roy and Sarma<sup>12</sup> have suggested a correction factor,  $G$ , in terms of system parameters. The correction factor is defined as

$$G = \frac{\Delta(P_T)_{exp}}{(\Delta P_T)_{cal}} \quad (6)$$

The calculated values of  $(\Delta P_T)$  were obtained using equation (4). The proposed correlations for the correction factor are presented in Table I.

Roy and Sarma<sup>13,14</sup> have also suggested another dimensionless correlation for pressure drop in liquid-solid and gas-solid systems (Table II).

**TABLE I**  
CORRELATIONS FOR THE CORRECTION FACTOR

Sl. No.	Correlation	System
1. (a)	$G = 7.3 \times 10^{-3} \left( \frac{D_c}{d_p} \right)^{-0.53} \left( \frac{\rho_s}{\rho_t} \right)^{1.18} \left( \frac{h_s}{D_c} \right)^{-2.05} (R)^{1.56} \left( \frac{h_{pa}}{h_s} \right)^{0.64}$	Gas-spherical solids <sup>11</sup> (Roy and Sen Gupta, 1973)
(b)	$G = 1.95 \times 10^{-1} \left( \frac{D_c}{d_p} \right)^{-0.59} \left( \frac{\rho_s}{\rho_t} \right)^{0.55} \left( \frac{h_s}{D_c} \right)^{-0.94} \left( \frac{h_{pa}}{h_s} \right)^{0.29} (R)^{0.72}$	Gas-non spherical solids <sup>11</sup> (Roy and Sen Gupta, 1973)
2.	$G = 1.67 \left( \frac{D_c}{d_p} \right)^{-0.59} \left( \frac{\rho_s}{\rho_t} \right)^{0.67} \left( \frac{h_s}{D_c} \right)^{-0.43} \left( \frac{h_{pa}}{h_s} \right)^{0.08} (R)^{0.08}$	Liquid-solid <sup>12</sup> (Roy and Sarma, 1974)

**TABLE II**  
CORRELATIONS FOR  $\Delta P_T / \Delta P_{ost}$

Sl. No.	Correlation	System
1.	$\frac{\Delta P_T}{\Delta P_{ost}} = 0.22 \left( \frac{D_c}{d_p} \right)^{-0.21} \left( \frac{\rho_s}{\rho_t} \right)^{0.48} (R)^{1.71} \left( \frac{h_{pa}}{h_s} \right)^{1.24}$	Gas-solids <sup>13</sup> (Roy and Sarma, 1977)
2.	$\frac{\Delta P_T}{\Delta P_{ost}} = 19.5 \left( \frac{D_c}{d_p} \right)^{-0.17} \left( \frac{\rho_s}{\rho_t} \right)^{0.48} (R)^{0.28} \left( \frac{h_{pa}}{h_s} \right)^{0.89}$	Liquid-solids <sup>14</sup> (Roy and Sarma, 1978)

Saha and Dutt<sup>15</sup> have put forward a generalized correlation for the prediction of pressure drop in a solid-liquid semifluidized bed. The correlation includes two friction-factor terms, one for the fluidized section and the other for the packed section of the bed. The correlation obtained for the prediction of pressure drop in a semifluidized bed is

$$\Delta P_T = \left( \frac{130}{Re_f} + 1.25 \right) \left( \frac{\rho_s}{\rho_f} \right)^{0.24} R^{-0.15} \frac{G_{sf} V_o (1 - \epsilon_f) (h_s - h_{pa})}{\epsilon_f^3 g_c d_p} + \left( \frac{150}{Re_{pa}} + 1.75 \right) R^{0.14} \phi_s^{-1.8} \left[ 1 + 0.02 F' \frac{h_{pa} G_{sf} V_o (1 - \epsilon_{pa})}{g_c d_p \epsilon_{pa}^3} \right] \quad (7)$$

where  $F'$  is the fraction of solids transferred to the packed bed section.

#### Minimum Semifluidization Velocity, $G_{osf}$

The minimum semifluidization velocity, or onset velocity of semifluidization, is defined as the fluid velocity at which the top of the fluidized bed just touches the top restraint of the semifluidizer. A number of correlations have been proposed to predict this velocity which are summarized below.

Poddar and Dutt<sup>15</sup> have reported data on semifluidization in liquid-solid systems. They employed the correlation of Wen and Yu<sup>17</sup> for the voidage function and the relationship between  $f$  and the voidage function (Wilhelm and Kwauk,<sup>18</sup> Lewis, *et. al.*<sup>19</sup> Richardson and Zaki<sup>20</sup>) to obtain the condition for minimum semifluidization. Roy and Sarma<sup>21</sup> obtained the experimental data for a few water-solid systems, and developed the following correlation for the prediction of minimum semifluidization velocity

$$G_{osf} = 0.2042 G_a^{0.675} [1 - (1/R) (1 - \epsilon_{pa})]^{2.87} \left( \frac{\mu_f}{d_p} \right) \quad (8)$$

Empirical correlations are also available in the form of equation (9)<sup>21-26</sup> and presented in Table III

$$f \left[ \frac{G_{osf}}{G_{mf}} \text{ or } \frac{G_{osf}}{G_{mf}}, \frac{D_o}{d_p}, \frac{\rho_s}{\rho_f}, R \right] = 0 \quad (9)$$

A few monographs<sup>27,28</sup> are also available for the rapid estimation of minimum semifluidization velocity (Figures 1 and 2).

**TABLE III**  
**CORRELATIONS FOR PREDICTION OF MINIMUM SEMIFLUIDIZATION VELOCITY**

Sl. No.	Correlation	System
1.	$\frac{G_{osf}}{G_{mf}} = 0.015 (R) + (\log Ar + 2.456)/52$	Water-solids <sup>21</sup> (Roy and Sarma, 1971)
2.	$\frac{G_{osf}}{G_{mf}} = 0.30 \left( \frac{D_o}{d_p} \right)^{-0.21} \left( \frac{\rho_s}{\rho_f} \right)^{0.41} (R)^{0.66}$	Water-solids <sup>21</sup> (Roy and Sarma, 1972)
3. (a)	$\frac{G_{osf}}{G_{mf}} = 1.625 \left( \frac{D_o}{d_p} \right)^{0.266} \left( \frac{\rho_s}{\rho_f} \right)^{-0.228} (R)^{0.585}$	Liquid-non-spherical particles <sup>25</sup> (Roy and Sarma, 1972)
(b)	$\frac{G_{osf}}{G_{mf}} = 1.875 \left( \frac{D_o}{d_p} \right)^{0.266} \left( \frac{\rho_s}{\rho_f} \right)^{-0.228} (R)^{0.585}$	Liquid-spherical solids <sup>25</sup> (Roy and Sarma, 1972)
4. (a)	$\frac{G_{osf}}{G_{mf}} = 2.66 \times 10^3 \left( \frac{D_o}{d_p} \right)^{0.62} \left( \frac{\rho_s}{\rho_f} \right)^{-1.00} (R)^{0.50}$	Gas-non-spherical particles <sup>24</sup> (Roy and Sen Gupta, 1972)
(b)	$\frac{G_{osf}}{G_{mf}} = 3.4 \times 10^3 \left( \frac{D_o}{d_p} \right)^{1.11} \left( \frac{\rho_s}{\rho_f} \right)^{-1.78} (R)^{0.89}$	Gas-spherical particles <sup>24</sup> (Roy and Sen Gupta, 1972)
5. (a)	$\frac{G_{osf}}{G_{mf}} = 0.475 \left( \frac{D_o}{d_p} \right)^{-0.20} \left( \frac{\rho_s}{\rho_f} \right)^{0.17} (R)^{0.38}$	Liquid-solids <sup>23</sup> (Roy, 1975)
(b)	$\frac{G_{osf}}{G_{mf}} = 0.071 \left( \frac{D_o}{d_p} \right)^{-0.20} \left( \frac{\rho_s}{\rho_f} \right)^{0.17} (R)^{0.38}$	Gas-solids <sup>23</sup> (Roy, 1975)
6.	$\frac{G_{osf}}{G_{mf}} = 48.0 \left( \frac{D_o}{d_p} \right)^{0.38} \left( \frac{\rho_s}{\rho_f} \right)^{-1.05} (R)^{0.64}$	Gas-solids <sup>26</sup> (Roy and Sen Gupta, 1975)

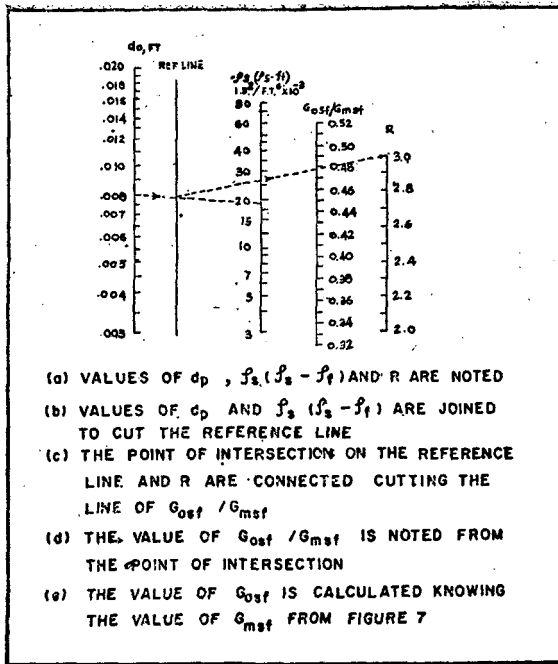


Fig. 1 Prediction of minimum semifluidization velocity for liquid-solid systems

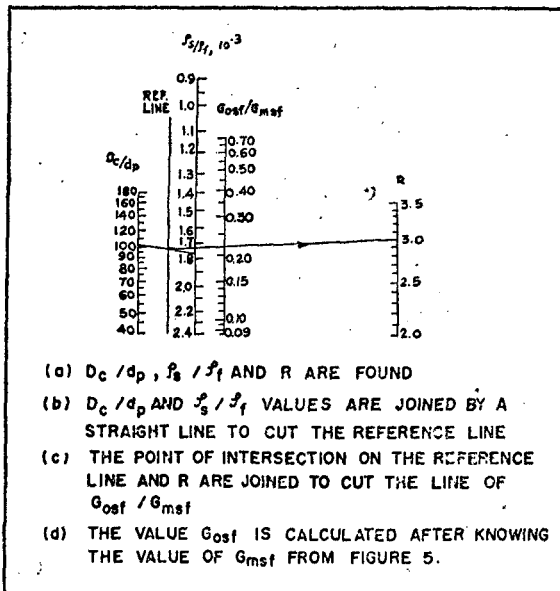


Fig. 2 Prediction of minimum semifluidization velocity for air-solid systems

Roy and Sharatchandra<sup>29</sup> obtained the data on liquid-solid semifluidization characteristics for systems of a heterogeneous mixture, viz. dolomite-chromite, dolomite-baryte, dolomite-iron ore, iron ore-chromite and iron ore-baryte. They have substituted an average density ( $\rho_{sav}$ ) for the term particle density by

$$(\rho_{sav}) = \frac{\sum W_i}{\sum \frac{W_i}{\rho_i}} \quad (10)$$

in the equation for minimum semifluidization proposed by Roy<sup>23</sup>

$$\frac{G_{ost}}{G_{msf}} = 0.473 \left( \frac{D_c}{d_p} \right)^{0.20} \left( \frac{\rho_s}{\rho_f} \right)^{0.17} (R)^{0.38} \quad (11)$$

The values of  $G_{msf}$  were calculated from equation (12) replacing the particle density term with the average particle density as defined in equation (10), and compared with the experimental values

$$G_{msf} = \frac{1.85 \times 10^4 d_p^{0.65} [\rho_f (\rho_s - \rho_f)]^{0.55}}{\mu_f^{0.1}} \quad (12)$$

For homogeneous mixtures, Roy and Dash<sup>30</sup> developed a correlation relating the ratio of maximum semifluidization velocity to minimum semifluidization velocity with the system parameters, substituting average particle size  $(d_p)_{av}$  instead of  $d_p$  in equations (11) and (12), where  $(d_p)_{av}$  was given by

$$\frac{1}{(d_p)_{av}} = \sum \frac{x}{d_p} \quad (13)$$

$x$  being the mass fraction of the components of the mixtures.

#### Maximum Semifluidization Velocity, $G_{msf}$

This is defined as the fluid velocity at which all solid particles are supported by the fluid in the packed portion of the bed. There are three methods<sup>10</sup> for the determination of  $G_{msf}$  experimentally (Figures 3 and 4 refer to two of them).

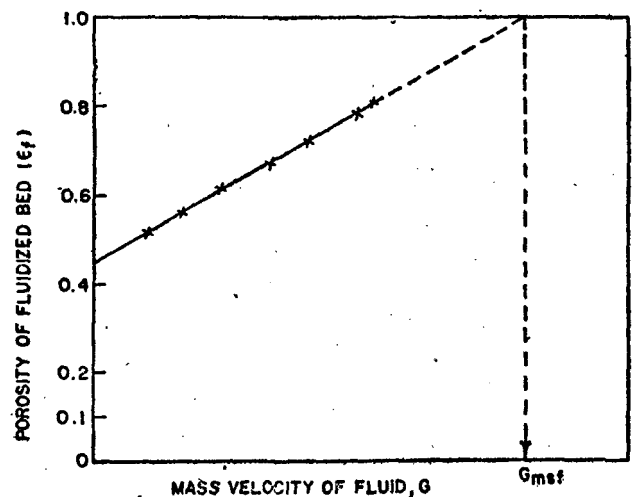


Fig. 3 Determination of  $G_{msf}$  from  $\epsilon_f$  vs.  $G$  plot

A number of correlations are also proposed in the literature<sup>16,21,28,30,31</sup> for the prediction of maximum

semifluidization velocity from system properties. Nomographs are also available for its rapid prediction as shown in Figures 5 and 6 for liquid-solid and gas-solid systems respectively.<sup>27,30</sup> Equations have also been

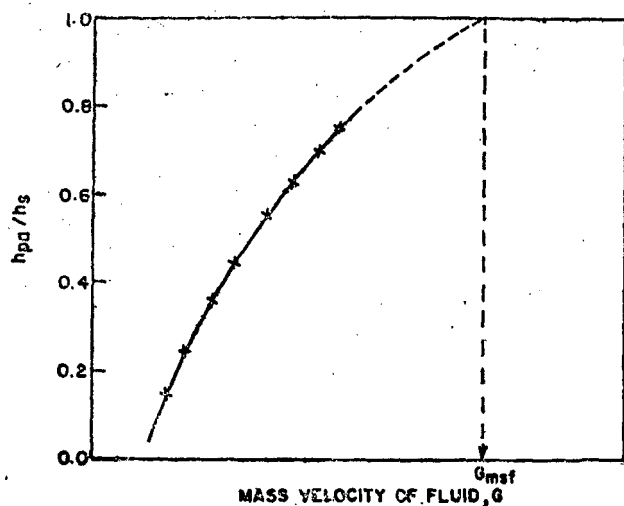


Fig. 4 Determination of  $G_{msf}$  from  $h_p/h_s$  vs.  $G$  plot

developed for its prediction from the minimum fluidization velocity<sup>32,33</sup>. These correlations are presented in Table IV.

Roy and Dash<sup>34</sup> developed a correlation for maximum semifluidization velocity in homogeneous mixtures. A similar correlation for heterogeneous mixtures has also been proposed by Roy and Sharat Chandra.<sup>35</sup>

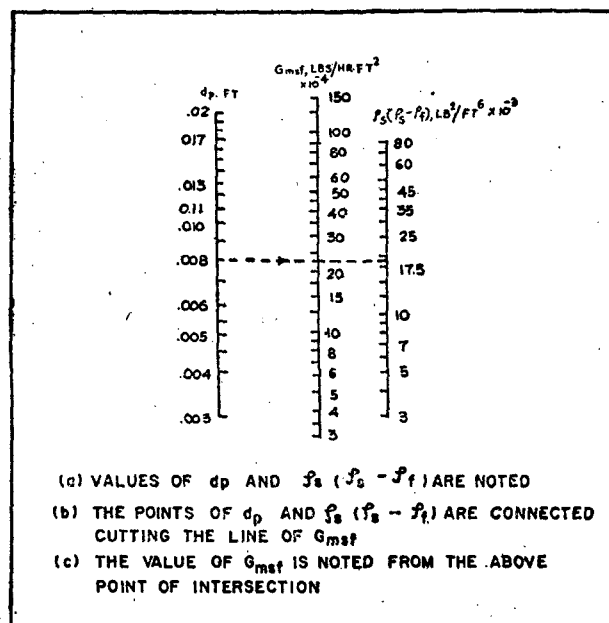


Fig. 5 Prediction of maximum semifluidization velocity for liquid-solid systems

TABLE IV

# CORRELATIONS FOR PREDICTION OF MAXIMUM SEMIFLUIDIZATION VELOCITY

Sl. No.	Correlation	System
1.	$18 Re_{msf} + 2.7 Re_{msf}^{1.687} = Ga$	Liquid-solids (Poddar and Dutta, 1969) <sup>18</sup>
2.	$G_{msf} = 0.3 (Ar)^{0.56} (\mu_t/d_p)$	Liquid-solids (Roy and Sarma, 1971) <sup>21</sup>
3.	$G_{msf}/G_{mf} = 5.71 (D_o/d_p)^{0.42} (\rho_s/\rho_t)^{0.67}$	Liquid-solids (Roy and Sarma, 1974) <sup>22</sup>
4. (a)	$Re_{msf} = 1.15 \times 10^{-3} (G_s)^{0.676}$	Gas-solids (Roy and Sen Gupta, 1975) <sup>28</sup>
(b)	$Re_{msf} = 1.15 \times 10^{-3} (Ar)^{0.676}$	Gas-solids (Roy and Sen Gupta, 1975) <sup>29</sup>
5. (a)	$G_{msf} = \frac{1.85 \times 10^4 (d_p)^{0.68} [\rho_t (\rho_s - \rho_t)]^{0.55}}{\mu_t 0.1}$	Liquid-solids (Roy, 1975) <sup>23</sup>
(b)	$G_{msf} = \frac{1.37 \times 10^4 (d_p)^{0.65} [\rho_t (\rho_s - \rho_t)]^{0.55}}{\mu_t 0.10}$	Gas-solids (Roy, 1975) <sup>23</sup>
6.	$G_{msf}/G_{mf} = 130 (D_o/d_p)^{0.5} (\rho_s/\rho_t)^{-0.72}$	Gas-solids (Roy and Saratchandra, 1976) <sup>32</sup>

Note: Equation (5):  $d_p$  in metres;  $\rho_s$  and  $\rho_t$  in  $kg/m^3$ ;  $\mu_t$  in  $kg/m.hr$ ;  $G_{msf}$  in  $kg/m^2.hr$ .

### Height of Packed Bed

For purposes of design it is imperative to know the variation in the height of the packed bed-formed below the restraint-with the change in fluid mass velocity,

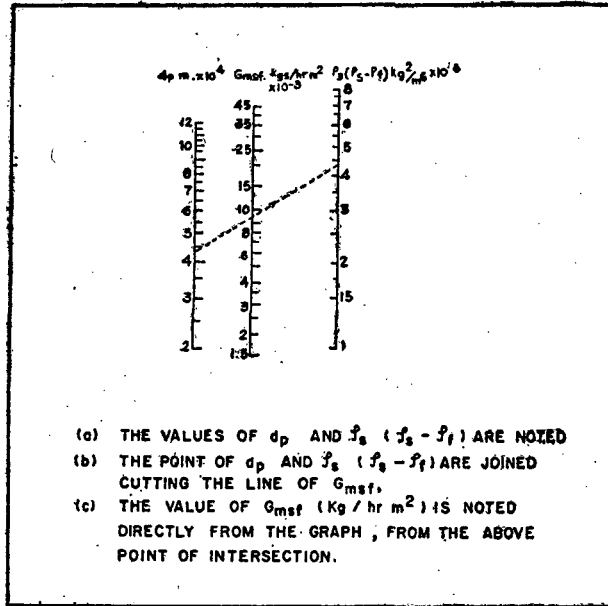


Fig. 6 Prediction of maximum semifluidization velocity for air-solid systems

the limits for the same being values corresponding to the onset and the maximum semifluidization conditions.

The methods available for the prediction of the height of the packed section in semifluidization are: (a) experimental determination with the parameters such as the position of screen and the fluid mass velocity, and (b) the method of Fan, *et al.*<sup>1</sup> based on a material balance approach. Fan and Wen<sup>9</sup> and Wen, *et al.*<sup>36</sup> have also suggested a dimensionless correlation for the prediction of the packed-bed height. Poddar and Dutt<sup>37</sup> have given a mathematical explanation for the packed bed formation under semifluidized condition, and have proposed a correlation for the estimation of the same for a solid-liquid system, based on the properties of solid and liquid and the flow conditions.

Since the formation of the packed bed begins at the onset of semifluidization and ends with the maximum semifluidization velocity, the introduction of the onset velocity of semifluidization,  $G_{osf}$ , instead of minimum fluidization velocity was considered to be more relevant by Roy and Sarma<sup>21</sup> in the derivation of Fan and Wen\* and Wen, *et al.*<sup>36</sup>.

All the correlations proposed for predicting the height of the packed-bed are presented in Table V.

**TABLE V**  
CORRELATIONS FOR PREDICTION OF PACKED BED HEIGHT

Sl. No.	Correlation	System
1.	$h_{pa} = \frac{h_{sf}(1-\epsilon_{pa})}{(\epsilon_f - \epsilon_{pa})} - \frac{h_s(1-\epsilon_{pa})}{(\epsilon_f - \epsilon_{pa})}$ $\epsilon_f = \frac{(18 Re + 2.7 Re^{1.687})^{0.2125}}{Ga}$	Liquid-solids <sup>37</sup> (Poddar and Dutta, 1970)
2.	$h_{pa} = (h_f - h_{sf}) \frac{(1 - \epsilon_f)}{(\epsilon_f - \epsilon_{pa})}$	Liquid-solids <sup>10,38</sup> (Fan and Wen, 1961; Wen, <i>et al.</i> , 1963)
3.	$\frac{(h_{sf} - h_s)}{(h_{sf} - h_{pa})} = \left( \frac{G_{sf} - G_{osf}}{G_{msf} - G_{osf}} \right)^{0.20}$	Liquid-solids <sup>21</sup> (Roy and Sarma, 1971)
4. (a)	$\frac{h_{pa}}{h_s} = 1.09 (G_{sf}/G_{msf})^{1.51} (D_c/d_p)^{0.23} (\rho_s/\rho_f)^{0.17} (R)^{-0.88} (h_s/D_c)^{-0.15}$	Liquid-non-spherical solids <sup>45</sup> (Roy and Sarma, 1973)
(b)	$h_{pa}/h_s = 2.21 (G_{sf}/G_{msf})^{2.08} (D_c/d_p)^{0.23} (\rho_s/\rho_f)^{0.17} (R)^{-0.88} (h_s/D_c)^{-0.15}$	Liquid-spherical solids <sup>45</sup> (Roy and Sarma, 1973)
5. (a)	$h_{pa}/h_s = 1.07 \times 10^{-5} (G_{sf}/G_{msf})^{3.22} (D_c/d_p)^{-0.17} (\rho_s/\rho_f)^{2.18} (h_s/D_c)^{2.78} (R)^{-1.66}$	Gas-non-spherical solids <sup>48</sup> (Roy and Sen Gupta, 1974)
(b)	$h_{pa}/h_s = 1.25 \times 10^{-5} (G_{sf}/G_{msf})^{3.13} (D_c/d_p)^{-0.72} (\rho_s/\rho_f)^{2.19} (h_s/D_c)^{2.19} (R)^{-1.66}$	Gas-spherical solids <sup>48</sup> (Roy and Sen Gupta, 1974)

### Lean Bed Zone

It has been observed by Wen, *et al.*<sup>36</sup> and Rao and Ramalingam that there exists a zone more or less devoid of solids in between the fluidized and packed-bed sections of a fluid-solid semifluidized bed and that this was more prominent in the case of gas-solid systems. Murthy, *et al.*<sup>40</sup> obtained data on the heights of the lean-bed zone in semifluidized beds for air-solid systems. Based on the lines of statistical design and dimensional analysis, an equation was presented for the prediction of the height of the lean-bed zone in terms of solid, fluid and bed properties as given below

$$Y = 138.7 + 2.5 X_1^2 + 5.1 X_2^2 + 20.6 X_3^2 + 6.0 X_4^2 + 2.5 X_5^2 - 16.1 X_1 + 26.3 X_4 + 19.5 X_5 + 10.3 X_4 X_5 + X_3(16.8 X_1 - 17.4 X_4 - 16.9 X_5 - 10.2 X_2) \quad (14)$$

The equation has been found to hold good within  $\pm 15$  per cent.

### Total Bed Height

Sunkoori, *et al.*<sup>41</sup> have observed an exponential relationship between the ratio of free surface fluid-bed height to fluid-bed height in semifluidization; and the ratio of total semifluidization height to static-bed height is given by the following equations

$$\left(\frac{h_{ff}}{h_f}\right) = A e^{0.1G} \quad (15)$$

$$A = 0.007 \left(\frac{h_{st}}{h_s}\right)^{2.5} \quad (16)$$

These authors have expressed the variation of the bed height with mass velocity in a phase diagram for the water-quartz system as shown in Figure 7.

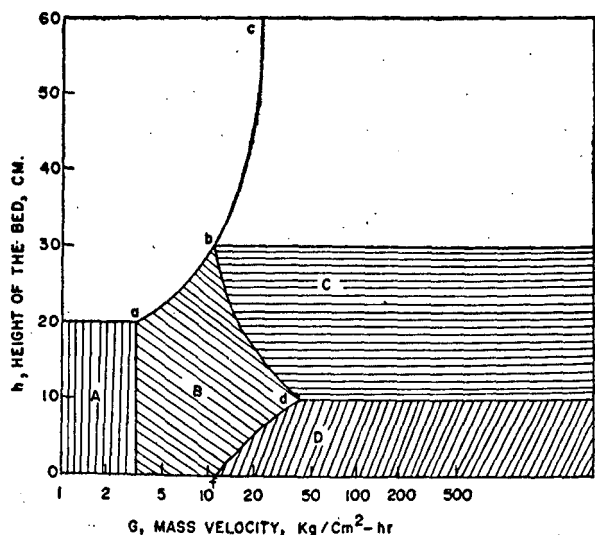


Fig. 7 Phase diagram: variation of bed height with mass velocity ( $h_s = 20$  cm)

### Residence Time Distribution (RTD)

This is an essential aspect of the design and proper working of semifluidized bed reactors. The only reported study on this aspect is by Saxena and Pandey.<sup>42</sup> They have measured RTD in semifluidized beds for glass beads and crushed calcite with water as the fluidizing medium. The effect of different variables has been studied and interpreted in the light of work done for fixed and fluidized systems.

It has been found that particle size has no effect on RTD for a fluidized bed but has an effect on a fixed bed. In the case of a semifluidized bed, the effect of particle diameter on residence time depends upon the extent to which the fixed-bed portion is formed. There was an increase in residence time with decrease in column diameter which has been attributed to the greater wall effect. The relationship between particle Reynolds number and residence time was given by the equation

$$t_{st} = 13.78 \times 10^3 (N_{Re})^{-0.78} \left(\frac{d_p}{D_c}\right)^{0.72} \quad (17)$$

with the maximum deviation of  $\pm 20$  per cent.

By considering the semifluidized bed as a combination of fluidized and packed beds, they proposed another correlation relating  $t_{st}$  to  $t_{fo}$  (mean residence time of free fluidized bed, i.e. without top restraint under similar conditions)

$$\frac{t_{st}}{(t_{fo})_{calc}} = 8.3 \times 10^{-1} \left[ \frac{(1-\epsilon_e)}{(1-\epsilon_{fo, calc})} \times \frac{\epsilon_p}{\epsilon_{fo, calc}} \right]^{0.42} \quad (18)$$

Pandey and Saxena<sup>43</sup> studied the axial dispersion in liquid semifluidized beds in glass columns of varying diameters with glass beads of different sizes as the bed material and water as the fluidizing medium. The effect of different variables has been investigated and the following correlation has been developed for the determination of the axial dispersion coefficient in a semifluidized bed

$$\left(\frac{D}{V_o L}\right)_{st} = 1.3 \times 10^{-4} (N_{Rem})^{1.85} \frac{SF}{SF} (dp/D_c)^{-0.80} \quad (19)$$

### State of Semifluidization

Empirical equations based on the dimensional analysis approach have been proposed to relate the various parameters when the bed is in a semifluidized condition. Thus, based on the dimensional analysis of the continuity and momentum equations for particle fluidization proposed by Beranek<sup>38</sup> and Beranek and Klumper,<sup>43</sup> Fan and Wen<sup>9</sup> have obtained the following relationship for a semifluidized bed

$$f \left[ \frac{(h-h_s)}{(h-h_{pa})}, \frac{G-G_{mf}}{G_t-G_{mf}} \right] = 0 \quad (20)$$

A two-parameter fit by Kurian and Rajarao<sup>10</sup> yielded an explicit form of the relationship as

$$\frac{G-G_{mf}}{G_t-G_{mf}} = 0.61 \left[ \frac{h-h_{pa}}{h-h_s} \right]^{-1.3} \quad (21)$$

On the other hand, Roy and Sarma<sup>45</sup> introduced the minimum semifluidization velocity in place of fluidization velocity in equation (21) and developed the following expression

$$\frac{h-h_s}{h-h_{pa}} = \left[ \frac{G-G_{ost}}{G_t-G_{ost}} \right]^{0.2} \quad (22)$$

Other investigators 23,28,45-48 empirically correlated the state of semifluidization in various forms of the function

$$f \left[ \frac{G}{G_t}, \frac{h_{pa}}{h_s}, \frac{D_c}{d_p}, \frac{\rho_s}{\rho_t}, R, \frac{h_s}{D_c} \right] = 0 \quad (23)$$

All these correlations are given in Table VI

**TABLE VI**  
**EQUATIONS WHICH DESCRIBE STATE OF SEMIFLUIDIZATION**

Sl. No.	Equation	Systems
1.	$\frac{G_{sf}}{G_t} = \frac{4.62}{D_c^{-0.372}} (Ga)^{-0.15} (Sf)^{-0.186}$	Gas-solids <sup>47</sup> (Baburao and Doraiswamy, 1967)
2.	(a) $\frac{G_{sf}}{G_{msf}} = 0.684 \left( \frac{D_c}{d_p} \right)^{-0.11} \left( \frac{h_s}{D_c} \right)^{-0.07} \left( \frac{\rho_s}{\rho_t} \right)^{-0.08} \left( \frac{h_{pa}}{h_s} \right)^{1.00} (R)^{0.43}$ (b) $\frac{G_{sf}}{G_{msf}} = 0.945 \left( \frac{D_c}{d_p} \right)^{-0.18} \left( \frac{h_s}{D_c} \right)^{-0.10} \left( \frac{\rho_s}{\rho_t} \right)^{-0.11} \left( \frac{h_{pa}}{h_s} \right)^{0.66} (R)^{0.57}$	Liquid-spherical solids <sup>48</sup> (Roy and Sarma, 1973)  Liquid-non-spherical solids <sup>45</sup> (Roy and Sarma, 1973)
3.	(a) $\frac{G_{sf}}{G_{msf}} = 37.07 \left( \frac{D_c}{d_p} \right)^{0.23} \left( \frac{h_s}{D_p} \right)^{-0.89} \left( \frac{\rho_s}{\rho_t} \right)^{-0.71} \left( \frac{h_{pa}}{h_s} \right)^{0.32} (R)^{0.53}$ (b) $\frac{G_{sf}}{G_{msf}} = 35.16 \left( \frac{D_c}{d_p} \right)^{0.23} \left( \frac{h_s}{D_c} \right)^{-0.86} \left( \frac{\rho_s}{\rho_t} \right)^{-0.68} \left( \frac{h_{pa}}{h_s} \right)^{0.31} (R)^{0.51}$	Gas-spherical solids <sup>26</sup> (Roy and Sen Gupta, 1975)  Gas-non-spherical solids <sup>46</sup> (Roy and Sen Gupta, 1975)
4.	(a) $\frac{G_{sf}}{G_{msf}} = 4.8 \left( \frac{D_c}{d_p} \right)^{-0.18} \left( \frac{\rho_s}{\rho_t} \right)^{-0.32} \left( \frac{h_{pa}}{h_s} \right)^{0.59} (R)^{0.81}$ (b) $\frac{G_{sf}}{G_{msf}} = 0.925 \left( \frac{D_c}{d_p} \right)^{-0.15} \left( \frac{\rho_s}{\rho_t} \right)^{-0.12} \left( \frac{h_{pa}}{h_s} \right)^{0.32} (R)^{0.43}$	Gas-solids <sup>23</sup> (Roy, 1975)  Liquid-solids <sup>23</sup> (Roy, 1975)
5.	$\frac{G_{sf}}{G_{msf}} = 0.925 \left( \frac{D_c}{d_p} \right)^{-0.15} \left( \frac{\rho_s}{\rho_t} \right)^{-0.12} (R)^{0.43} \left( \frac{h_{pa}}{h_s} \right)^{0.32}$	Liquid-solids <sup>48</sup> (Irregular particles) (Roy and Biswal, 1977)

The hydrodynamic behaviour of semifluidization has been investigated by S.H. Chem, *et al.*<sup>49</sup> Various parameters under extensive analysis include pressure drop, gas hold-up, onset liquid velocity for semifluidization and the height of the packed bed and the fluidized bed sections. The pressure drop of the semifluidized bed obtained experimentally was found to compare favourably with that predicted by the model equations.

### Heat and Mass Transfer Operations

A brief review dealing with heat and mass transfer studies was made by Murthy, *et al.*<sup>50</sup> A summary of the investigations conducted so far on heat and mass transfer in semifluidized beds is given below.

#### Heat Transfer

There are three important aspects that may be investigated in connection with semifluidized bed heat transfer: (i) fluid-to-particle heat transfer; (ii) wall-to-fluid heat transfer; and (iii) particle-to-fluid heat transfer. The published literature refers to wall-to-fluid heat transfer.

Purnachandra Rao and Kaparthi<sup>39</sup> reported the studies on the wall-heat transfer coefficients of semifluidized beds in a 25 mm copper column, employing different sizes of glass beads, quartz and aluminium particles fluidized with air. The overall heat transfer

coefficient of the bed was predicted using Leva and Grummers<sup>51</sup> equation for packed bed, the equation of Urie<sup>52</sup> for the fluidized bed and the Sieder Tate<sup>53</sup> equation for the empty-section heat transfer coefficient from a knowledge of the heights of the various sections of a semifluidized bed. The equation was given as

$$H_{sf} = \frac{H_{pa} h_{pa} + H_f h_f + H_e h_e}{h_{pa} + h_f + h_e} = \sum \frac{hH}{h} \quad (24)$$

Based on experimental observations, the authors concluded that the wall-to-bed heat transfer coefficients decreased with porosity and particle density and increase with particle diameter.

Verma, *et al.*<sup>54</sup> studied the heat transfer characteristic of semifluidized beds using liquid-solid systems and reported similar observations as those by Rao and Kaparthi<sup>39</sup> for gas-solid systems.

Murthy, *et al.*<sup>55</sup> obtained data for heat transfer coefficients for air-solid systems. With the help of dimensional analysis and based on statistical experimental design, an equation was formulated to predict the Nusselt number in terms of different system variables.

Empirical equations reported in the literature for the prediction of heat transfer coefficient in semifluidized beds are given in Table VII

**TABLE VII**  
**EQUATIONS FOR PREDICTION OF HEAT TRANSFER COEFFICIENTS**

Sl. No.	Equation	System
1.	$Nu_p = 0.014 (Re_p)^{1.1} \left( \frac{1-\epsilon}{\epsilon} \right)^{0.4}$	Gas-solids <sup>39</sup> (Purnachandra Rao and Ramalingam, 1969)
2. (a)	$Nu_p = 0.0285 (Re_p)^{1.21} (Pr)^{0.33} (R)^{-1.1} \left( \frac{d_p}{D_c} \right)^{-0.58}$	Water-glass beads <sup>54</sup> (Verma, <i>et al.</i> 1972)
(b)	$Nu_p = 0.0032 (Re_p)^{1.21} (Pr)^{0.33} (R)^{-1.1} \left( \frac{d_p}{D_c} \right)^{-0.58}$	Water-aluminium particles <sup>54</sup> (Verma, <i>et al.</i> 1972)
3.	$Z = 0.975 - 0.067 Y_1 - 0.052 Y_2 + 0.093 Y_3 - 0.043 Y_4 + 0.07 Y_1 Y_4$	Air-solids <sup>55</sup> (Murthy, <i>et al.</i> 1983)

### Mass Transfer

Fan, *et al.*<sup>1</sup> initiated the studies in the field of mass transfer with a benzoic acid-water system, and correlated the mass transfer data in terms of the  $j_d$  factor and the modified Reynolds number. Fan, *et al.*<sup>\*</sup> also suggested the correlations in terms of the Sherwood and Reynolds numbers for a packed bed and fluidized bed and for a combination of the fluidized and packed beds.

Rai, *et al.*<sup>56</sup> undertook the work to study the effect of variables, viz. the properties of the solid particles (density, shape and size) and the fluid (density and viscosity) and operating conditions (fluid velocity and diffusivity). Benzoic acid, cinnamic acid and 2-naphthol were used as the solid particles with water as the fluidizing medium. Govindarajan and Sen Gupta<sup>57</sup> obtained mass transfer data for the benzoic acid-water system for various particle sizes and initial static bed heights. The correlations proposed by the various investigators relating to mass transfer studies are presented in Table VIII.

### Design of Semifluidized Beds

The application of a semifluidized bed reactor was illustrated by Baburao<sup>58</sup> in a study involving the

Oxidation of benzene. The optimal conversion condition was obtained by modelling the semifluidized bed reactor as a CSTR and a tubular reactor in series. After the relative sizes of the fluidized and the packed sections were determined, the equation for the state of semifluidization described earlier<sup>48</sup> was used to determine the operating parameters. Experimental conversion values of the semifluidized bed, the mixed and the tubular reactors were compared and an improved performance was obtained in the case of the semifluidized one.

Murthy, *et al.*<sup>59</sup> suggested a detailed procedure for the design of a semifluidized bed reactor. They proposed a modified form of the reactor suggested by Baburao, *et al.*<sup>2</sup> Equations proposed for the study of momentum transfer operations and the optimal criteria suggested by Baburao<sup>58</sup> were made use of in the suggested design procedure. Upadhrashta, *et al.*<sup>60</sup> have presented the salient design features of a semifluidized bed reactor for the vapour-phase oxidation of naphthalene to phthalic anhydride (an exothermic reaction). They have observed that the lowest catalyst-to-reactant-flow rate ratio is achieved for such a reactor.

**TABLE VIII**  
**CORRELATIONS FOR MASS TRANSFER IN SEMIFLUIDIZED BEDS**

Sl. No.	Correlation	System
1.	$J_D = 1.865 (Re_m)^{-0.48}; Re_m = Re_p / (1 - \epsilon); 5 < Re_m < 30$	Water-benzoic acid <sup>1</sup> (Fan, <i>et al.</i> , 1959)
2.	$N_{sh} = 2.0 + 0.312 (Re_p)^{0.408} (N_{so})^{0.333}$	Water-solids <sup>6</sup> (Fan, <i>et al.</i> , 1960)
3. (a)	$N_{sh} = 0.0637 \left( \frac{\rho_s - \rho_f}{\rho_f} \right) \left( \frac{d_p}{D_o} \right)^{0.63} (N_{Rep})^{0.729} (N_{so})^{0.333}$	Water-solids <sup>56</sup> (Rai, <i>et al.</i> , 1970)
(b)	$N_{sh} = 0.226 \left( \frac{\rho_s - \rho_f}{\rho_f} \right) \left( \frac{d_p}{D_o} \right)^{-0.588} (N_{Rep})^{0.729} (N_{so})^{0.5}$	Water-benzoic acid <sup>56</sup> (Rai, <i>et al.</i> , 1970)
4. (a)	$J_D = 2.745 (N_{Rep})^{-0.484}$	Water-benzoic acid <sup>56</sup> (Rai, <i>et al.</i> , 1970)
(b)	$J_D = 2.35 (N_{Rep})^{-0.484}$	Water-cinnamic acid <sup>56</sup> (Rai, <i>et al.</i> , 1970)
(c)	$J_D = 1.865 (N_{Rep})^{-0.484}$	Water-2-naphthol <sup>56</sup> (Rai, <i>et al.</i> , 1970)
5.	$k_L a = 2.512 \times 10^{-5} \left( \frac{d_p G_{sf}}{\mu_f} \right)^{0.51} (R)^{-1.03}$	Water-benzoic acid <sup>57</sup> (Govindarajan, 1976)

Note:  $k_L$  in cm/sec

The design of a semifluidizer for a liquid-solid system was given by Roy<sup>61</sup> and for a gas-solid system by Roy and Sen Gupta.<sup>62</sup> The method of calculating the various quantities including (a) the range of semifluidization operation, (b) the height of the packed and fluidized sections, (c) semifluidization velocity, and (d) the power requirement, were described for given operating conditions in the case of a dolomite-water system.

### Miscellaneous Industrial Applications

The application of semifluidized beds has been broadly stressed by Fan and Hsu<sup>3</sup> and Baburao, *et al.*<sup>2</sup> Semifluidized beds find wide application as reactors (for exothermic and bioreactions) in ion exchange and in filtration operations for the removal of suspended particles from gases or liquids. A brief description of the utility of semifluidized beds in each of the above operations is given below.

#### *Ion Exchange*

Mobay Chemical Company, Asahi Chemical Company of Japan, Bayer A G (Germany) and its U.S. counterpart, Lewatit Company, are advocating the use of semifluidized beds with ion-exchange resins.<sup>3</sup> This action arises from the discovery that a fluidized bed followed by a fixed bed increases the efficiency of resin utilization by improving liquid-resin contact. The fixed bed acts as a polishing section, handles the ion leakage from the fluidized bed and prevents elutriation of resin particles.<sup>63-65</sup> In addition to a higher resin utilization efficiency, a semifluidized bed also minimizes the volume of regenerant and wash water needed, reduces pressure drop and operates more consistently than the conventional fixed bed process.

#### *Filtration*

For filtration purposes the fixed bed section functions as a deep bed filter, and turbulence generated in the fluidized section scours the deposited solids, delaying the build-up of pressure drop. When the fixed portion of a filtration bed is used in conjunction with a fluidized portion of the bed, as is possible in a semifluidized bed, unique advantages are obtained. During the filtration cycle most of the filtered particles are retained in the fluidized bed portion. As the cake of the filtered particles collects at the bottom of the fixed-bed portion, the filtered particles then circulate with the fluidized media in the bed. This action eliminates the blinding of the filter bed and thus prolongs the filtration cycle. Moreover, the pressure drop across the fixed bed can be maintained uniformly.

This method of filtration has application in the removal of insoluble ash and mineral matter from coal

oil obtained from the hydrogenation and dissolution of coal; and in the removal of fine particles in a hot flue gas from power plants or from the product gases of coal-conversion reactors.

Hsu and Fan<sup>4</sup> carried out an experimental study of filtration of a slurry composed of 50-mesh coal particles dispersed in water. The filtering medium consisted of —20 to +50-mesh silica sand. Filter performance was determined with 25, 50, 75 and 100 per cent of the filtering medium initially in the packed section of the semifluidized bed. Filtration with 100 per cent of the bed particles corresponds to the conventional up-flow deep bed filtration. Samples of the filtrate were collected intermittently from a sample point located between the upper porous plate and the outlet. These samples were then filtered to determine the solid content.

The results of the study showed that the performance of the semifluidized bed filter was far superior to that of the deep bed filter. It was also established that filtration runs lasting as long as 6 times that of the deep bed filter could be attained without apparent deterioration in the quality of the filtrate.

#### *Bio-Reactors*

The unique advantage of a semifluidized bed as a bioreactor has been mentioned in detail by Fan and Hsu.<sup>3,5</sup> Studies have shown that when micro-organisms are attached to inert supports their effectiveness is immensely improved in the degradation of hazardous organic contaminants and nitrogen compounds, in the reduction of total organic carbon (TOC), BOD and total suspended solids (TSS), and in the conversion of volatile solids to methane gas. A packed-bed operation using immobilized micro-organisms in anaerobic wastewater treatment achieved removal of more than 70 TSS, 50 BOD and 60 TOC with a residence time of less than 5 hours.<sup>5</sup>

In fluidized bed operations the use of immobilized micro-organisms can achieve near complete removal of organic toxicants, nitrogen and BOD with a residence time of a few minutes compared to 1 to 3 hours for conventional processes.

The use of a semifluidized bed would eliminate the disadvantages of a fluidized bed, for example elutriation of the particles coated with micro-organisms, and unstable bed expansion. The semifluidized bed would also reduce or eliminate plugging of the bed by solids (waste or microbial cells) experienced in fixed-bed operations.

Initially, the semifluidized bed may be seeded by introducing micro-organisms and nutrients to the bed under flow rates slightly higher than the minimum fluidization velocity. The micro-organisms will first grow in suspended phase and then attach themselves to the

available surface area, i.e., the column wall and the fluidizing particles. The particles may consist of sand, coal or packings with large surface area. Once the particles are covered with microbes the semifluidized bed reactor is in operation.

The fluidized bed portion of the bed will carry the main load of digestion with the packed bed acting as a polishing section. The fluid reaching the latter section will be lean in nutrients (or waste constituents) and thus will not contribute excessively to an increase in cell masses that can clog the bed. To control the microbial population the entire bed should periodically be fluidized. A saving of about 95 per cent in space can be realized by installing a semifluidized bed bioreactor instead of a conventional suspended-growth waste-treatment unit.

#### *Chemical Reactors*

Cholette, Blanche and Cloutier<sup>66,67</sup> have shown that a combination of mixed and tubular reactors is often theoretically more efficient than either of these reactors operated independently. For endothermic reactions, a tubular reactor is always superior, while for exothermic reactions a CSTR is superior to a tubular reactor up to a certain conversion after which a tubular reactor is more efficient.

In carrying out an exothermal reaction under adiabatic conditions in a tubular reactor two opposite influences may be noted: (i) an increase in reaction rate due to an adiabatic rise in temperature, and (ii) a decrease in reaction rate due to a decrease in reactants' concentration. In the earlier action of the reactor the influence of the temperature is normally more marked leading to an increase in reaction rate. After a certain stage the effect of concentration exerts a greater influence leading to a decrease in reaction rate. That portion of the reaction which occurs in the increasing-rate zone can be advantageously carried out in a fully mixed reactor. The subsequent part, occurring in the decreasing rate zone, should be carried out in a tubular reactor.

It is clear from the above argument that for an exothermal adiabatic reaction the first reactor should be a CSTR which will give an outlet conversion corresponding to the maximum reaction rate. The products can then be led into a tubular reactor for achieving the final degree of conversion required.

The theoretical advantage of the mixed reactor-tubular reactor (MT) combination has been practically realized in a simple reactor system utilizing the principle of semifluidization.<sup>2</sup> For this purpose, a bundle of rigid plastic tubes were fixed to a perforated plate. This bundle was inserted into a fluidized bed reactor. It was possible to form fixed beds in the tubes

while at the same time retaining a portion of the solids in a fluidized state at the bottom of the reactor. The proportion of solids in the tubular fixed and fluidized beds was varied both by varying the velocity of the fluid and the portion of the tubular bundle. Thus, this type of reactor can be operated as an MT combination, the lower fluidized bed providing the mixed reactor and the top packed portion as the tubular reactor. The fraction of the bed that should act as a CSTR can be calculated from the equation developed by Aris.<sup>68</sup> With this conversion as the starting point the length of the tubular reactor for attaining a given conversion can be calculated. With the help of available methods.

The reactor of the above type has specific advantages for fast exothermic reactions such as the vapour-phase oxidation and chlorination of hydrocarbons.

#### *Scope for Future Work*

The review of the literature reveals that the information on semifluidization is highly incomplete especially with respect to heat and mass transfer studies. In addition, information relating to the design of semifluidized systems is too meagre.

Available work in the field of heat transfer is too limited to assess the effect of the system variables, viz. fluid and solid properties and the operating conditions. Hence, systematic and thorough investigations are warranted in this field to study the effect of the above variables on heat transfer rate.

Further, for the prediction of heat transfer rates in a semifluidized bed the formulation of equations based on a theoretical standpoint will be more dependable and of wider applicability than the empirical relations developed so far.

Mass transfer in a semifluidized bed is considered to be a gray area, most of which still remains unexplored. Available information is highly empirical in nature and of limited use. Hence it is desirable to make a detailed study of the effect of all the variables on mass transfer rate in a semifluidized bed, and to propose correlations based on a theoretical background which will be of a general nature and valid beyond experimental conditions.

Prediction of pressure drop is still not very accurate due to non-availability of a suitable method for the correct determination of porosity in the packed section of the semifluidized bed. Hence, independent determination of this important parameter is required.

The literature on the determination of the heights of packed, fluidized and lean beds in semifluidization is very limited. Much additional work is warranted for the accurate determination of these heights.

The design of semifluidized beds for various applications is a field virtually untouched. Hence,

there is wide scope to deal with the design aspects of a number of important systems of both a chemical and physical nature where the semifluidized bed can find a potential use in view of its proven advantages over the conventional fixed and fluidized beds.

Dynamic and microscopic studies, e.g. analysis of pressure fluctuations and particle motion, need to be intensified.

### Conclusion

In recent years, the application of semifluidization, to different, physical and chemical processes is gaining momentum. Recent developments of semifluidized; bed ion-exchangers, adsorbers, filters and reactors on a

laboratory scale are significant in this respect. There has been a conceptual development of a semifluidized bed reactor in the form of the MT-reactor on a laboratory scale, which is advocated as being more efficient than either the fixed or the fluidized bed (for fast exothermic reactions). However, attempts to scale up and analyze its performance are few. Hence, there is need for investigations on bed semifluidized systems on a broader spectrum to judge their suitability to different process applications along the lines that have become common for other fluid-solid systems such as packed and fluidized beds.

*Paper received: 18.3.86 Revised: 19.7.86 Accepted: 25.7.86*

### NOMENCLATURE

a	specific surface area of the particles
C	pressure drop correction factor
$C_{pf}$	specific heat of fluid
$C_{ps}$	specific heat of solid
D	diffusivity
$D_c$	diameter of column
$d_p$	diameter of particle
f	friction factor
$g_c$	gravitational constant
G	mass velocity of fluid
$G_{mf}$	minimum fluidization mass velocity
$G_{ost}$	onset mass velocity of semifluidization
$G_{st}$	semifluidization mass velocity
$G_t$	free-settling mass velocity of the solid
h	height of semifluidized bed
$h_e$	height of empty section
$h_f$	height of fluidized bed
$h_{ff}$	free fluidized bed height
$h_{lbz}$	height of lean bed zone
$h_{pa}$	height of packed bed
$h_s$	static bed height
$h_{st}$	height of semifluidized bed
$H_e$	heat transfer coefficient of solids free section
$H_f$	heat transfer coefficient of fluidized section
$H_{pa}$	heat transfer coefficient of packed bed section
$H_{st}$	heat transfer coefficient of semifluidized bed section
$k_f$	thermal conductivity of the fluid
$k_L$	mass transfer coefficient
$\Delta P_T$	total pressure drop across a semifluidized bed
R	bed expansion ratio. $= h_{st}/h_{pa}$
$t_{fo}$	mean residence time for free fluidized bed
$t_{st}$	mean residence time for semifluidized bed
$V_o$	linear velocity of fluid
$W_p$	weight of solid in packed section in semifluidization
$W_s$	weight of solid in initial static bed
W	weight of the solid mixture

X	mass fraction of the material
$x_1$	$D_c/d_p$
$x_2$	$\rho_s/\rho_f$
$x_3$	$G_{st}/G_{mf}$
$x_4$	R
$x_5$	$h_s/d_p$
$X_1$	$(x_1 - 77.16)/(25.72)$
$X_2$	$(x_2 - 77.16)/(25.72)$
$X_3$	$(x_3 - 0.6)/(0.15)$
$X_4$	$(x_4 - 3.5)/(0.5)$
$X_5$	$(x_5 - 144.05)/(20.55)$
Y	$h_{lbz}/d_p$
$y_1$	$G_{st}/G_{mf}$
$y_2$	$\rho_s/\rho_f$
$y_3$	$C_{ps}/C_{pf}$
$y_4$	R
$Y_1$	$(y_1 - 0.575)/(0.175)$
$Y_2$	$(y_2 - 2.725 \times 10^3)/(0.275 \times 10^3)$
$Y_3$	$(y_3 - 0.87)/(0.207)$
$Y_4$	$(y_4 - 3.25)/(0.75)$
Z	$h_{dp}/k_f$

### Greek letters

$\rho_s$	density of solid material
$\rho_f$	density of fluid
$\beta$	constant
$\epsilon$	porosity of initial static bed
$\epsilon_{pa}$	porosity of the packed bed section of semifluidized bed
$\epsilon_f$	porosity of the fluidized bed section of semifluidized bed
$\mu_f$	viscosity of fluid
$\phi_s$	shape factor

### Subscripts

a	additional
av	average
f	fluidized bed
fo	free fluidized bed

msf	maximum semifluidization condition
osf	onset of semifluidization
pa	packed bed condition
sf	semifluidization condition

### Dimensionless groups/numbers

Ar	Archimedis number = $\frac{d_p^3 g_c \rho_s (\rho_s - \rho_f)}{\mu_f^2}$
Ga	Galileo number = $\frac{d_p^3 \rho_f (\rho_s - \rho_f) g}{\mu_f^2}$
J <sub>D</sub>	mass transfer factor
J <sub>H</sub>	heat transfer factor
Nu <sub>p</sub>	particle Nusselt number, = $\frac{H d_p}{k_f}$
N <sub>sh</sub>	Sherwood number, $k_L h/D$
N <sub>Re</sub>	
or Re <sub>p</sub>	particle Reynolds number, $\frac{d_p G}{\mu_f}$
N <sub>sc</sub>	Schmidt number, $\mu_f / \rho_f D$
Re <sub>mf</sub>	$D_p G / \mu_f (1 - \epsilon_f)$
Sf	Semifluidization number, $= (w_s - w_o) / (h - h_s) \rho_s$

### ACKNOWLEDGEMENT

The authors are thankful to Mr. P. K. Patra and Mr. G. N. Sahoo for their assistance in the preparation of this paper.

### REFERENCES

1. Fan, L. T., Yang, Y. C. and Wen, C. Y., *A.I.Ch.E.J.*, **5**, 407 (1959)
2. Baburao, K., Mukherjee, S. P. and Doraiswamy, L. K., *A.I.Ch.E.J.*, **11**, 741 (1965)
3. Fan, L. T. and Hsu, E. H., Paper presented at the Seminar on Theoretical and Industrial aspects of Semifluidization and Fluidization, Rourkela, India, Nov. 26-30 (1979)
4. Hsu, E. H. and Fan, L. T., *Proceedings of the World Filtration Congress III*, Philadelphia, Pennsylvania, U.S.A. September 13-17 (1982)
5. Fan, L. T. and Hsu, E. H., *Proceedings of the VI International Fermentation Symposium*, London, Ontario, Canada, July 20-25 (1980)
6. Fan, L. T., Yang, Y. C. and Wen, C. Y., *A.I.Ch.E.J.*, **6**, 482 (1960)
7. Leva, M., *Fluidization*, McGraw-Hill, New York (1959)
8. Walas, M. S., *Reaction Kinetics for Chemical Engineers*, McGraw-Hill, New York (1959)
9. Fan, L. T. and Wen, C. Y., *A.I.Ch.E.J.*, **7**, 606 (1961)
10. Kurian, J. and Rajarao, M., *Ind. J. of Tech.*, **8**, 275 (1970)
11. Roy, G. K. and Sen Gupta, P., *Chem. Eng. J.*, **5**, 191 (1973)
12. Roy, G. K. and Sarma, K. J. R., *Ind. Chem. Engineer*, **XIII**, No. 4, 31 (1974)
13. Roy, G. K. and Sarma, K. J. R., *J. of Inst. Engineers (India)*, **57**, pt. Ch 3, 122 (1977)
14. Roy, G. K. and Sarma, K. J. R., *Ind. J. of Tech.*, **16**, 89 (1978)
15. Saha, M. and Dutt, D. K., *Ind. Chem. Eng.*, **XXIV**, No. 1, 30 (1982)
16. Poddar, S. K. and Dutt, D. K., *Ind. Chem. Eng.*, **11**, No. 3, 80 (1969)
17. Wen, C. Y. and Yu, Y. H., *Chem. Engg. Progr. Symp. Ser.*, **62**, 101 (1966)
18. Wilhelm, R. H. and Kwauk, R., *Chem. Engg. Prog.*, **44**, 201 (1948)
19. Lewis, W. K., Gilliland, E. R. and Baner, W. C., *Ind. Engg. Chem.*, **41**, 1104 (1969)
20. Richardson, J. F. and Zaki, W. N., *Trans. Inst. Chem. Engg.*, **32**, 35 (1954)
21. Roy, G. K. and Sarma, K. J. R., *Chem. Proc. and Eng.*, **5**, 23 (1971)
22. Roy, G. K. and Sarma, K. J. R., *Chem. Engg. Journal*, **4**, 1 (1972)
23. Roy, G. K., 'Studies on Certain Aspects of Semifluidization', Ph.D. thesis, Sambalpur University, India (1975)
24. Roy, G. K. and Sen Gupta, P., *Ind. J. of Tech.*, **10**, 397 (1972)
25. Roy, G. K. and Sarma, K. J. R., *Chem. Engg. J.*, No. 4 (1972)
26. Roy, G. K. and Sen Gupta, P., *Processing*, **10** (1975)
27. Roy, G. K., Sen Gupta, P. and Sarma, K. J. R., *Chemical Age of India*, **22**, 637 (1971)
28. Roy, G. K. and Sen Gupta, P., *J. Inst. of Eng. (India)*, **56**, Pt. CH 1, 40 (1975)
29. Roy, G. K. and Sharat Chandra, H. N., *Chem. Engg. J.*, **12**, 77 (1976)
30. Roy, G. K. and Dash, J., *Ind. Chem. J.*, **1**, Feb. (1977)
31. Roy, G. K. and Sarma, K. J. R., *Chemical Era*, **VIII**, 1 (1972)
32. Roy, G. K. and Sarma, K. J. R., *J. Inst. Engineers (India)*, **54**, 34 (1974)
33. Roy, G. K. and Sarat Chandra, H. N., *Ind. Chem. J.*, Aug. (1976)
34. Roy, G. K. and Dash, J., *Ind. Chem. J.*, Jan. (1976)
35. Roy, G. K. and Sarat Chandra, H. N., *Trans. Ind. Chem. Eng.*, **XIX**, No. 1, 33 (1977)
36. Wen, Y. G., Cheng, S. W. and Fan, L. T., *A.I.Ch.E.J.*, **9**, 316 (1963)
37. Poddar, S. K. and Dutt, D. K., *Ind. Chem. Eng.*, **XII**, No. 1, 38a (1970)
38. Beranek, I. J., *British Chem. Engg.*, **3**, 368 (1958)
39. Purna Chandra Rao, S. and Ramalingam, K., *Ind. Chem. Eng.*, **XI**, No. 2, 43 (1969)
40. Murthy, J. S. N., Narayana, A. S., Roy, G. K. and Sarma, K. J. R., *Ind. Chem. Eng.*, **XXIII**, No. 4, 26 (1981)
41. Sunkoori, N. R., Moinuddin, S. and Kaparthi, R., *Trans. Ind. Chem Eng.*, **XI**, No. 3, 89 (1969)
42. Sexena, D. N. and Pandey, G. N., *Ind. J. of Technol.*, **15**, 455 (1977)
43. Sexena, D. N. and Pandey, G. N., *Ind. Chem. Eng.*, **XXII**, No. 1, 40 (1980)
44. Beranek, I. J. and Klumpar, T., *Collection Czechoslov. Chem. Commun.*, **23**, 1 (1958)
45. Roy, G. K. and Sarma, K. J. R., *Ind. J. of Tech.*, **11**, 237 (1973)
46. Roy, G. K. and Biswal, K. C., *J. of Chem. Engg. of Japan*, **10**, 330 (1977)
47. Baburao, K. and Doraiswamy, L. K., *A.I.Ch.E.J.*, **13**, 397 (1967)
48. Roy, G. K., Sen Gupta, P., *Ind. Engg. Chem. Process Design Dev.*, **13**, 219 (1974)
49. Churn, S. H., Fan, L. S. and Muroyama, K., *A.I.Ch.E.J.*, **30**, 288 (1984)
50. Murthy, J. S. N., Roy, G. K. and Sarma, K. J. R., *Chem. Engg. World*, **XI**, 55 (1975)
51. Leva, M., Grummer, M. and Clark, R. R., *Ind. Eng. Chem.*, **40**, 747 (1948)
52. Urie, R. W., M. S. Thesis, Cambridge, M.I.T., (1948)
53. Kern, D. Q., *Process Heat Transfer*, McGraw-Hill (1950)
54. Varma, R. L., Pandey, G. N. and Tripathi, G., *Ind. J. of Technol.*, **19**, 11 (1972)
55. Murthy, J. S. N., Suryanarayana, A., Roy, G. K. and Sarma, K. J. R., *J. of Inst. Eng. (India)*, **63**, Pt. CH 3 (1983)
56. Rai, H. S., Pandey, G. N. and Tripathi, G., paper presented at the 23rd Annual Session of Indian Institute of Chemical Engineers at Hyderabad (India) (1970)
57. Govindarajan, P. N. and Sen Gupta, P., *Trans. of Ind. Chem. Eng.*, **XVIII**, 51 (1976)
58. Baburao, K., *A.I.Ch.E.J.*, **16**, 273 (1970)
59. Murthy, J. S. N., Patnaik, G. and Sarma, K. J. R., *Chem. Age of India*, **28**, 777 (1977)
60. Upadrashta, K. R., Sreerangam, V. and Kaparthi, R., *Chem. Ind. Dev.*, Jan. 9 (1979)
61. Roy, G. K., *Chem. Engg. World*, **VII**, 47 (1972)
62. Roy, G. K. and Sen Gupta, P., *Chemical Industry Developments*, May (1973)
63. Lewatit Company Brochure (1972)
64. Chementator, *Chem. Engg.*, Nov. 26 (1962)
65. Chementator, *Chem. Engg.*, July, 24 (1972)
66. Cholette, A. and Blanchet, J., *Can. J. Chem. Eng.*, **39**, 192 (1961)
67. Cholette, A. and Cloutier, L., *Can. J. Chem. Eng.*, **37**, 105 (1959)
68. Aris, R., *Can. J. Chem. Engg.*, **40**, 87 (1962)
69. Roy, G. K. and Sen Gupta, P., *Ind. Chem. Eng.*, **XVII**, No. 4, 40 (1975)