Calculation of Tower Height

The physical absorption process can be carried out in countercurrent flow process, which may be carried out in packed or tray column:



HOG * NOG

Where:

HOG: is the height of transfer unit (HTU) based on gas phase, and it can be calculated from the equation

below:

$$HOG = \frac{G_s}{KoG. a} , in (meter)$$

NOG: is the number of transfer unit (NTU) based on gas phase, and it can be calculated based on equilibrium data:



If the equilibrium data are *linear*, then **NOG** will be calculated using a suitable equation. If the equilibrium data are *nonlinear*, then **NOG** will be calculated graphical method. If the equilibrium data are *linear*, then **N** will be calculated using a suitable equation.

If the equilibrium data are *nonlinear*, then N will be calculated graphical method.

H : is the distance between two trays, and it is given (0.3 - 0.7 m)

N : is the number of trays, and it can be calculated based on equilibrium data:



1. Packed tower:

Absorption and stripping are frequently conducted in packed columns, particularly when:

- (1) the required column diameter is less than 0.6 m.
- (2) the pressure drop must be low, as for a vacuum service.
- (3) corrosion consideration favor the use of ceramic or polymeric material.
- (4) low liquid holdup is desirable.

The gas liquid contact in a packed bed column is continuous, not stage-wise, as in a plate column. The liquid flows down the column over the packing surface and the gas or vapour, counter-currently, up the column. In some gas-absorption columns co-current flow is used. The performance of a packed column is very dependent on the maintenance of good liquid and gas distribution throughout the packed bed, and this is an important consideration in packed-column design.

Calculations of the packing height based on gas phase:

Overall material balance on the solute (A) over an element (∂z) based on gas phase:



 $G_S \; dY = L_S \; dX = N_{A^{\cdot}} \; A$

$$N_{A} = G_{s} Y - G_{s} \left(Y + \frac{dY}{dZ} \partial z \right) = (KoG)(a \ S \partial z)(Y - Y^{*})$$

Where:

The interficial area for transfer = $a dV = a S \partial z$

S: is the cross-sectional area of column (m^2) .

a: is the surface area of interface per unit volume of column (m^2/m^3) .

$$-G_{s}\left(\frac{dY}{dZ} \partial z\right) = (KoG. a)(S. \partial z)(Y - Y^{*})$$
$$G_{s}\frac{dY}{dZ} = -(KoG. a)(S. \partial z)(Y - Y^{*})$$

$$\int_{0}^{Z} dZ = \frac{-G_{s}}{(KoG. a).S} \int_{Y_{1}}^{Y_{2}} \frac{dY}{(Y - Y^{*})}$$
$$Z = \frac{(G_{s} / S)}{KoG.a} \int_{Y_{2}}^{Y_{1}} \frac{dY}{(Y - Y^{*})}$$
$$Z = \frac{G_{s}}{KoG.a} \int_{Y_{2}}^{Y_{1}} \frac{dY}{(Y - Y^{*})}$$

Z = HOG * NOG = HTU * NTU

Where:

 $HOG = \frac{\overline{G}_{s}}{KoG.a} : \text{heiht of transfer unit (HTU) based on gas phase, with the units of (m).}$ $NOG = \int_{Y_{2}}^{Y_{1}} \frac{dY}{(Y - Y^{*})} : \text{number of transfer unit (NTU) based on gas phase, without units.}$

Equation of the operating line:

Solute material balance between one end of the column and any point will give:

$$G_{s} (Y - Y_{2}) = L_{s} (X - X_{2})$$
$$Y = \frac{L_{s}}{G_{s}} X - X_{2} + Y_{2}$$

* The equation of operating line is a relation between mole ratio of solute in gas phase (Y) and the mole ratio of solute in liquid phase (X).



* The operating line can be draw from two points (X1, Y1) and (X2, Y2), or from its slope $(\frac{L_s}{G_s})$

and one of the two points.

