

Oving 4

1

In a heat exchanger with constant heat capacities and constant UA , we have the following simple formula for the total heat transfer from the hot to the cold stream [J/s]

$$Q = UA\Delta T_M$$

where $\Delta T_M = \frac{\Delta T_1 - \Delta T_2}{\ln(\Delta T_1 / \Delta T_2)}$ is the log-mean temperature difference.

Countercurrent absorption is similar to a heat exchanger, but the driving force is the concentration difference between y in the gas phase and $y^*(x)$ in equilibrium with the liquid phase. With constant slopes for the equilibrium and operating lines (which is reasonable for dilute mixtures), the total mass transfer of A from the gas to liquid stream is [mol A/s] (as derived in the lecture)

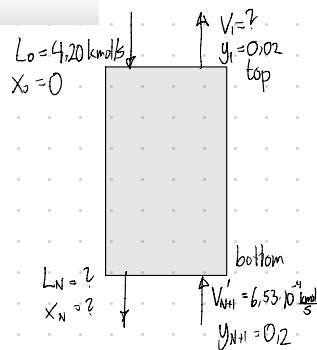
$$N_A = K_y(aS_z)(y-y^*)_M$$

where (aS_z) [m²] is the total mass transfer area inside the column and $(y-y^*)_M$ is the log mean vapor concentration difference between the phases, $(y-y^*)_M = \frac{(y-y^*)_1 - (y-y^*)_2}{\ln((y-y^*)_1 / (y-y^*)_2)}$ (1=top of column, 2=btm of column).

*it should be easy to remember this formula at the exam and possibly for the rest of your life!

We want to apply this easy-to-remember formula to calculate the absorption of SO₂ using water. A tray tower is to be designed to absorb SO₂ (A) from an air stream by using pure water at 293 K (680F). The entering gas contains 20 mol % SO₂, and that leaving 2 mol % at a total pressure of 101.3 kPa. The inert air flow rate is 150 kg air/h • m², and the entering water flow rate is 6000 kg water/h • m². Assuming an overall tray efficiency of 25%, **how many theoretical trays and actual trays are needed?** Assume that the tower operates at 293 K (20°C). It is not really a dilute mixture, so we expect some error.

Given data: The mole fraction of SO₂ drops from 0.2 (btm) to 0.02 (top) in air. The airflow (inert) is $6.53 \cdot 10^{-4}$ kmol/s. The entering liquid is $4.20 \cdot 10^{-2}$ kmol/s of pure water. The cross-sectional area of the column is 0.0929 m².



Find the total mass transfer of A from the vapor to liquid,

$$N_A = y_{\text{btm}}V_{\text{btm}} - y_{\text{top}}V_{\text{top}} \text{ [mol A/s]}$$

Oppgitt: ~~Column efficiency = $\frac{N_t - 1}{N_a} = 0.25$~~

$$A = 0.0929 \text{ m}^2$$

$$V_{N+1} = \frac{V_{N+1}}{1-0.2} = 8.1625 \cdot 10^{-4} \text{ kmol/s}$$

$$V_1 = \frac{V_1}{1-0.02} = 6.6632 \cdot 10^{-4} \text{ kmol/s}$$

$$\Rightarrow N_A = 0,281625 \cdot 10^{-4} - 0,02 \cdot 6,632 \cdot 10^{-4} \text{ kmol/s}$$

$$\underline{N_A = 1,499 \cdot 10^{-4} \text{ kmol/s}}$$

ii)

The column operates at 1 atm and 20°C. Use the equilibrium data to find the slope m of the equilibrium line for the concentration range of interest.

Linearregresjon av doten gir.

$$y = 31,62 \cdot x - 0,01$$

$$\Rightarrow \underline{m = 31,62}$$

A.3-19 Equilibrium Data for SO₂-Water System

Mole Fraction SO ₂ in Liquid, x_A	Partial Pressure of SO ₂ in Vapor, P_A (mm Hg)		Mole Fraction SO ₂ in Vapor, y_A , $P = 1$ Atm	
	20°C (293 K)	30°C (303 K)	20°C	30°C
0	0	0	0	0
0.0000562	0.5	0.6	0.000658	0.000790
0.0001403	1.2	1.7	0.00158	0.00223
0.000280	3.2	4.7	0.00421	0.00619
0.000422	5.8	8.1	0.00763	0.01065
0.000564	8.5	11.8	0.01120	0.0155
0.000842	14.1	19.7	0.01855	0.0259
0.001403	26.0	36	0.0342	0.0473
0.001965	39.0	52	0.0513	0.0685
0.00279	59	79	0.0775	0.1040
0.00420	92	125	0.121	0.1645
0.00698	161	216	0.212	0.284
0.01385	336	452	0.443	0.594
0.0206	517	688	0.682	0.905
0.0273	698		0.917	

Source: T. K. Sherwood, Ind. Eng. Chem., 15, 740 (1923).

iii)

Compute K_{ya} using the data $k_{La} = 0.85 \text{ kmol/s, m}^3$ and $k_{Ga} = 0.037 \text{ kmol/s, m}^3$.

$$\frac{1}{K_{ya}} = \frac{1}{k_{ya}} + \frac{m}{k_{xa}}$$

$$\Rightarrow K_{ya} = \frac{1}{\frac{1}{k_{ya}} + \frac{m}{k_{xa}}} = \frac{1}{\frac{1}{0,037} + \frac{31,62}{0,85}} = \underline{0,016 \text{ kmol/s, m}^3}$$

iv)

Find the logarithmic mean driving force $(y - y^*)_M$

$$(y - y^*)_M = \frac{(y - y^*)_1 - (y - y^*)_2}{\ln \frac{(y - y^*)_1}{(y - y^*)_2}} = \frac{(0,02 - 0) - (0,2 - 0,1)}{\ln \frac{0,02}{0,1}} = \underline{0,05}$$

$$y_1 = y_{\text{top}} = 0,02$$

$$y_2 = y_{\text{bot}} = 0,2$$

$$x_1 = x_{\text{top}} = 0 \Rightarrow y_1^* = 0$$

y_2^* finnes ved interpolasjon fra tabellen, trengs x_N , mabalans SO_2 :

$$x_N = \frac{N_A}{L_0 + N_A} = \underline{0,00356}$$

$$y_2^* = 0,0775 + (0,00356 - 0,00275) \frac{0,121 - 0,0775}{0,082 - 0,00275}$$

$$y_2^* = 0,10$$

v)

Use the easy-to-remember formula to find the packing height z .

$$N_A = K_{ya} S z (y - y^*)_M \Rightarrow z = \frac{N_A}{K_{ya} S (y - y^*)_M} = \frac{1,499 \cdot 10^{-4} \text{ kmol/s}}{0,016 \text{ kmol/s, m}^3 \cdot 0,0094 \cdot 0,05}$$

$$\underline{z = 2,02 \text{ m}}$$

2

Absorption of CO₂ in water-amine solution (part 2)

This continues the previous exercise, but now you should use the rate-based method (mass transfer) rather than equilibrium stages. Note that since we do not really have stages anymore, we choose to use a different notation with "top" and "bun", so x_0 is now x_{top} , etc. (see Figure).

Let us first repeat the basic data for the column:

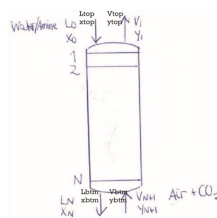
$$V_{bun} = 22 \text{ kmol/s}$$

$$y_{bun} = 0.1176 \text{ (CO}_2 \text{ mole fraction) (feed)}$$

$$y_{top} = 0.0116$$

$$x_{top} = 0.0219 \text{ (feed)}$$

$$x_{bun} = 0.0532$$



Assuming constant inert flows for air (V') and water-amine (L) through the column, we have

$$V' = V_{bun} (1 - y_{bun}) = 19.41 \text{ kmol/s}$$

$$L = 69.80 \text{ kmol/s}$$

The operating line is not quite linear, but it goes through the points (x_{top}, y_{top}) and (x_{bun}, y_{bun}) . To simplify, you may assume that the operating line is a straight line between these two points

(which corresponds to using a constant $\bar{K} = \frac{y_{top} - y_{bun}}{x_{top} - x_{bun}}$)

Table: Equilibrium data for CO₂ and water-MEA at 1.2 bar and about 60°C (inert air present).

$x_{CO_2}(\%)$	$y_{CO_2}(\%)$
2.227	1.162
3.001	2.917
3.339	3.783
4.101	6.057
4.700	8.448
5.324	11.60

$$\Rightarrow p = 1.2 \text{ bar}, T = 60^\circ\text{C}$$

The new part of the problem:

The overall mass transfer rate referred to the gas side is with pressure as the driving force (A=CO₂):

$$N_A = K_G \cdot a (p_A - p'_A)$$

where p'_A is the partial pressure of A that would result if we had gas in equilibrium with the liquid. The following data is given:

$$K_G \cdot a = 12.5 \text{ lbmol/(h} \cdot \text{ft}^3 \cdot \text{atm)}$$

a) Use the data to find $K_G a$ [kmol/(s·m³ packing)] as used in the book,

$$N_A = K_G a (y_A - y'_A)$$

where y'_A is the mole vapor mole fraction of a gas (imagined) in equilibrium with the liquid (which has mole fraction x_A).

a)

$$\text{Setter } N_A = N_A \Rightarrow K_G a (y_A - y'_A) = K_G a (p_A - p'_A)$$

$$K_G a = \frac{K_G a (p_A - p'_A)}{(y_A - y'_A)}, \quad p_A = P \cdot y_A$$

$$\Rightarrow K_G a = K_G a \frac{P(y_A - y'_A)}{(y_A - y'_A)}$$

$$K_G a = 12.5 \frac{\text{lb} \cdot \text{mol}}{\text{h} \cdot \text{ft}^3 \cdot \text{atm}} \cdot 1.2 \text{ bar}$$

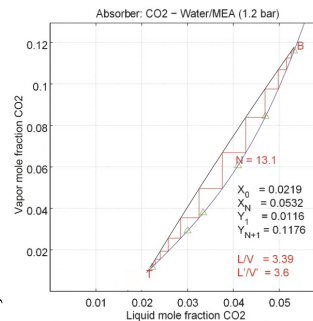
$$= 15 \cdot \frac{\text{lb} \cdot \text{mol} \cdot \text{bar}}{\text{h} \cdot \text{ft}^3 \cdot \text{atm}} \cdot 1.01325 \frac{\text{atm}}{\text{bar}} \cdot 0.95 \frac{\text{kmol}}{\text{lbmol}} \cdot \frac{1}{3600 \text{ s}} \cdot \frac{1}{(0.3048 \text{ m})^3}$$

$$K_G a = 0.067 \frac{\text{kmol}}{\text{s} \cdot \text{m}^3}$$

- b) Find the required packing height when $S = 253.5 \text{ m}^2$ (cross-sectional area of column) and you can assume constant molar flows in the column.

Note from the McCabe-Thiele diagram from the previous exercise that the equilibrium line is far from linear, so we cannot use the simple method from Problem 1.

(Hint: Make a table of $1/(y_A - y_A^*)$ as a function of y_A ; you can then integrate graphically, for example, by counting the number of squares on an mm-paper).



b) Massebalanse gir: $N_A dA = d(Vy_A)$ $V = \text{const}$

$$K y_A (y_A - y_A^*) S dz = V dy_A$$

$$\Rightarrow Z = \frac{V}{K y_A S} \int_{y_{A,\text{top}}}^{y_{A,\text{bot}}} \frac{dy_A}{y_A - y_A^*} = \frac{20.82 \text{ kmol/s}}{0.067 \text{ kmol/s.m}^2 \cdot 253.5 \text{ m}^2} \int_{y_{A,\text{top}}}^{y_{A,\text{bot}}} \frac{dy_A}{y_A - y_A^*}$$

Operating line: (fra forrige øving)

$$y_{N+1} = \frac{L_N}{V_{N+1}} x_N + \frac{y_1 V_1 - x_0 L_0}{V_{N+1}}$$

$$y = \frac{L}{V} x + y_{\text{top}} - \frac{L}{V} x_{\text{top}}$$

$$y = \frac{L}{V} (x - x_{\text{top}}) + y_{\text{top}}, \text{ gitt: } \frac{L}{V} = \frac{y_{\text{top}} - y_{\text{bot}}}{x_{\text{top}} - x_{\text{bot}}} = \frac{0.0116 - 0.1176}{0.0219 - 0.0532}$$

$$y = 3.387 x - 0.0626 \quad \frac{L}{V} = 3.387$$

Før denne tabellen:

Gitt i oppgaven

Operating line

x_A	y_A^*	y_A	$1/(y_A - y_A^*)$
0.02227	0.01162	0.01282849	827.47892
0.03001	0.02917	0.03904387	101.277412
0.03339	0.03783	0.05049193	78.97690163
0.04101	0.06057	0.07630087	63.56927494
0.047	0.08448	0.096589	82.58320258
0.05324	0.116	0.11772388	580.086781

$$Z = 1.226 \int_{y_{A,\text{top}}=0.0116}^{y_{A,\text{bot}}=0.1176} \frac{dy_A}{y_A - y_A^*} = 1.226 \cdot 13.5$$

↑
diagram neste side

$$Z = 16.6 \text{ m}$$

La $V = \frac{V_{\text{top}} + V_{\text{bot}}}{2}$

$$V = \frac{19.64 + 22}{2} \text{ kmol/s}$$

forrige øving

$$V = 20.82 \text{ kmol/s}$$

