

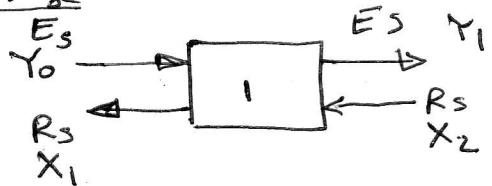
Calculations Based on total immiscibility of carrier solvents (A and B).

For simple ternaries on solute free basis:

Y_1 : mass ratio of solute in Extract (E phase).

X_1 : mass ratio of solute in Raffinate (R phase).

Simple single stage



$$MBI: R_s(X_2 - X_1) = E_s(Y_1 - Y_0)$$

$$R_s(X_2 - X_1) = E_s Y_1$$

$$Y_1 = m X_1$$

$$\therefore \frac{X_2}{X_1} = m \frac{E_s}{R_s} + 1$$

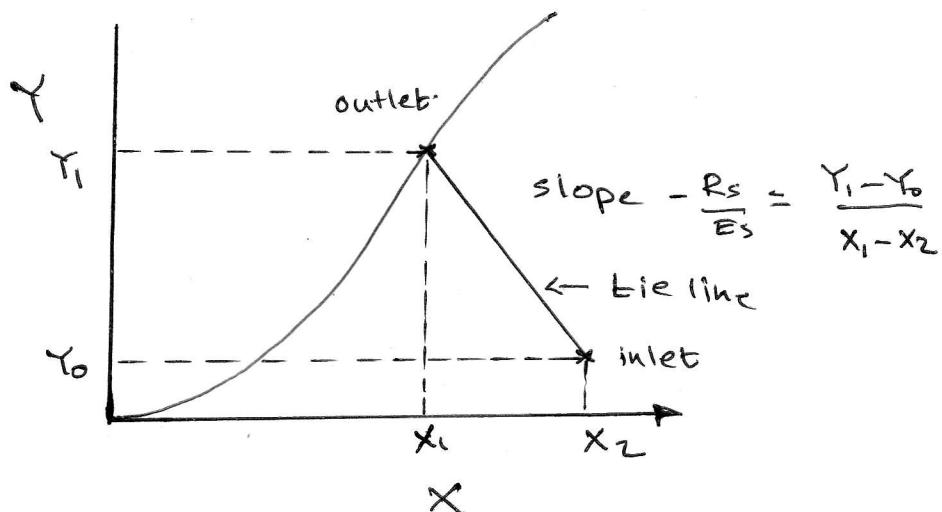
Extraction Ratio

for fresh solvent $Y_0 = 0$

At eqm $Y = mX$

m : partition coefficient

: at any value of $X = k$ tangent to curve at that point



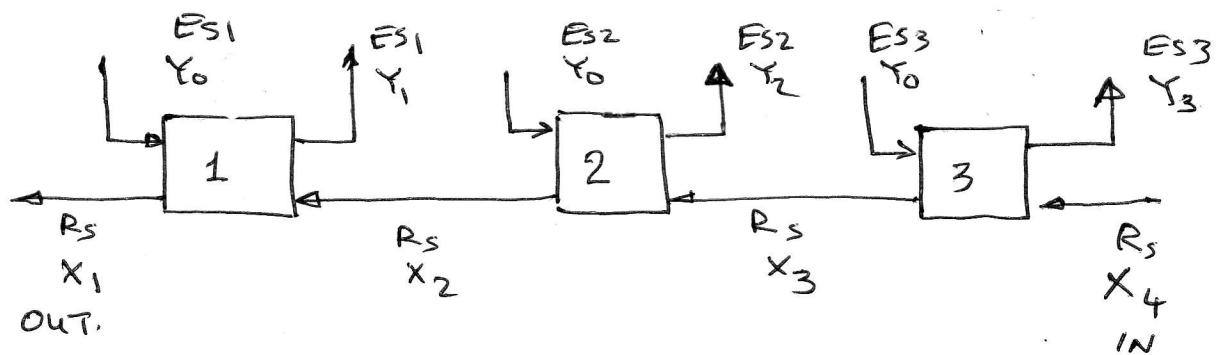
$$\% \text{ Recovery } (\% \text{ Extraction}) = \frac{X_2 - X_1}{X_2} \times 100 \quad X_1 : \text{out} \quad X_2 : \text{in.}$$

$$E = m \frac{E_s}{R_s} \quad \text{Extraction factor.}$$

$$\therefore \frac{X_2}{X_1} = (E + 1)$$

$$\Rightarrow \frac{X_1}{X_2} = \frac{1}{(E + 1)} = \frac{x_{in}}{x_{out}}$$

Simple multiple contact (Repeated application of Solvent. Crossflow)
consider 3 contacts for example.



Increase range of extraction ratio and enables production of extracts of different concentration.

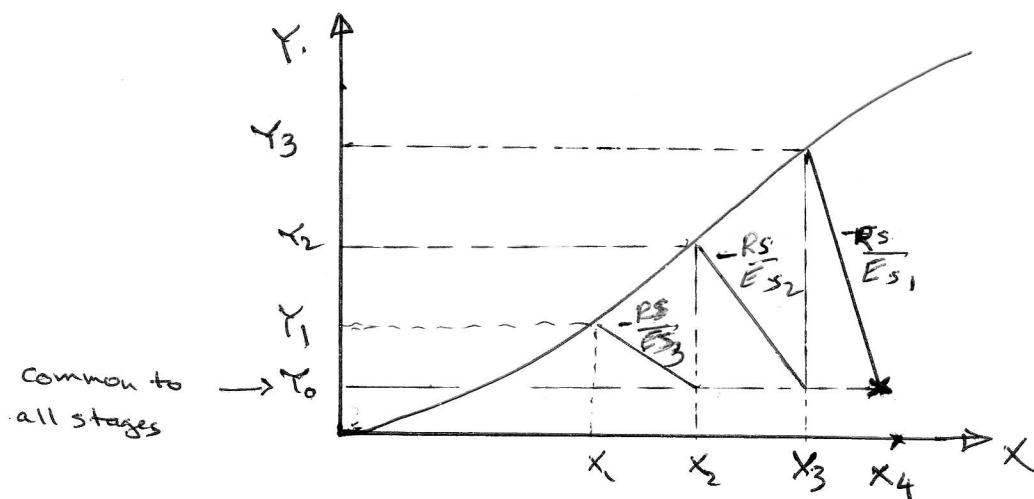
$$\text{Extraction ratio} = \frac{X_4}{X_1}$$

$$\Rightarrow \frac{X_4}{X_1} = \frac{X_4}{X_3} \times \frac{X_3}{X_2} \times \frac{X_2}{X_1}$$

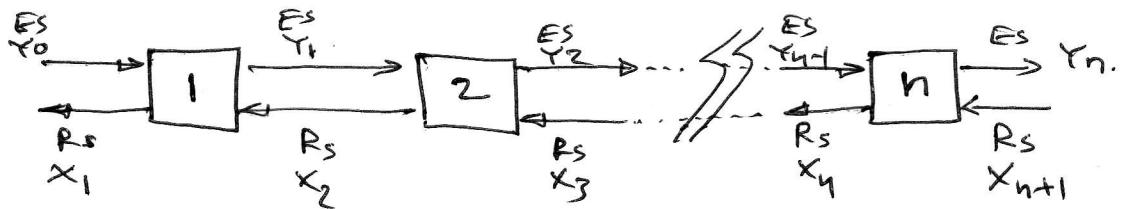
$$= \left(m \frac{E_{S1}}{R_s} + 1 \right) \left(m \frac{E_{S2}}{R_s} + 1 \right) \left(m \frac{E_{S3}}{R_s} + 1 \right).$$

Special case $E_{S1} = E_{S2} = E_{S3} = E_0$ = $\frac{\text{Total Solvent}}{\text{No. of stages}}$.
 $m = \text{const.}$

$$\therefore \frac{X_4}{X_1} = \left(m \frac{E_0}{R_s} + 1 \right)^3$$



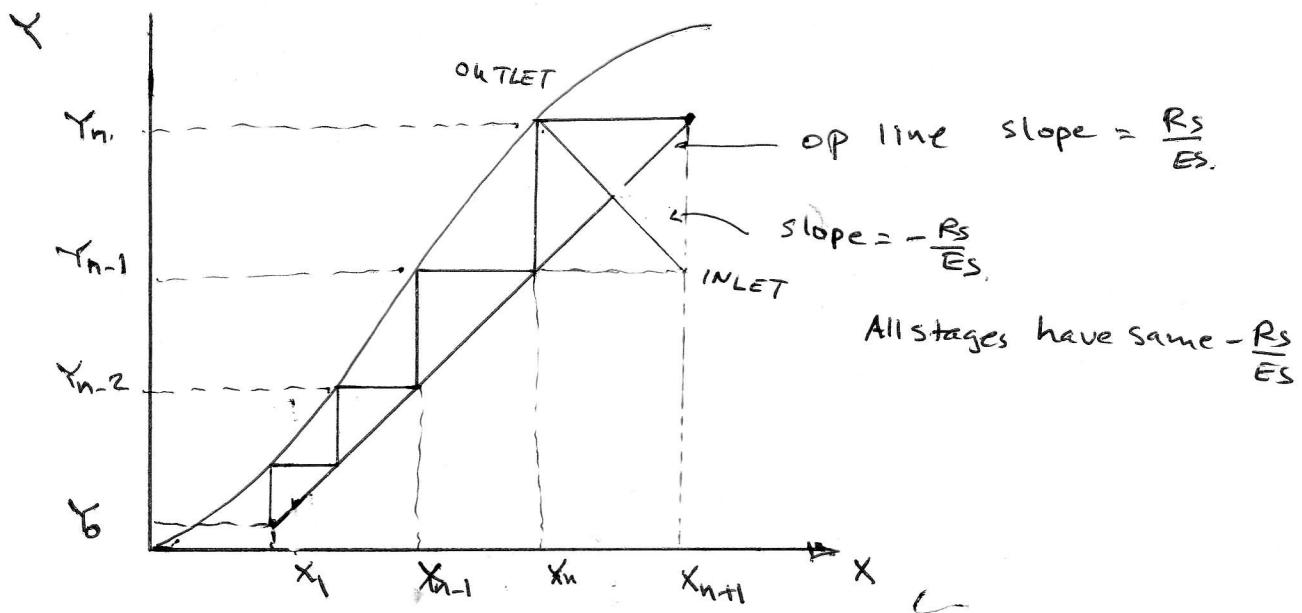
Continuous Counter Current Extraction:



From solute mass balances on each stage.

$$\text{Extraction ratio} = \frac{X_{n+1}}{X_1}$$

$$\left| \frac{X_{n+1}}{X_1} = \frac{\left(\frac{mES}{RS}\right)^{n+1} - 1}{\frac{mES}{RS} - 1} \right|$$

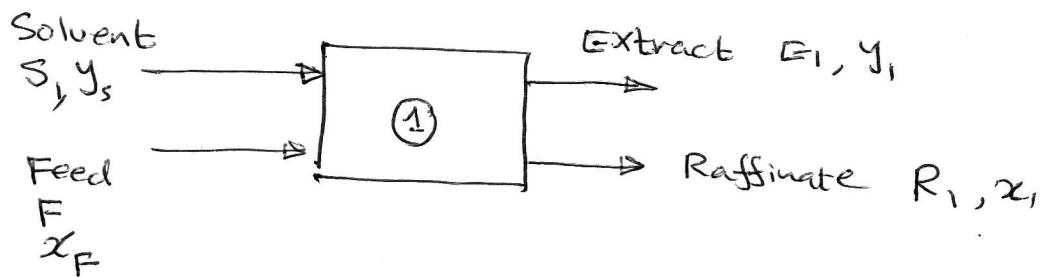


Impure solvent ie recycled solvent

$$Y_0 \neq 0$$

$$\text{Extraction ratio} = \frac{X_{n+1}}{X_1} = \frac{\left[\left(\frac{mES}{RS}\right)^{n+1} - 1\right]}{\left[\left(\frac{mES}{RS}\right) - 1\right]} - \frac{ES}{RS} \frac{Y_0}{X_1} \left[\frac{\left(\frac{mES}{RS}\right)^n - 1}{\left(\frac{mES}{RS}\right) - 1} \right]$$

$$\text{if } \frac{ES}{RS} > 1$$

General Case:Single Stage contacting:

F: Feed stream - contains diluent (A) + solute (C)

S₁: Solvent stream to stage: generally contains small amounts of A and C dissolved in solvent (Recycled)

CASE:

Known variables: $F, S_1, \underbrace{x_{CF}, x_{AF}}_{\text{solute}}, \underbrace{y_{CS}, y_{AS}}_{\text{diluent}}$

Unknown or desired: E_1, R_1 and outlet stream compositions
8 variables.



Mixture M_1 is a two phase mixture

Equations:

Overall balance

$$F + S_1 = M_1 = E_1 + R_1 \quad (1)$$

Component balances:

C-balance:

$$x_{CF} F + y_{CS} S = y_{C1} E_1 + x_{C1} R_1 \quad (2)$$

A-balance

$$x_{AF} F + y_{AS} S = y_{A1} E_1 + x_{A1} R_1 \quad (3)$$

Equilibrium: Solute is distributed between two phases

$$y_{CE} = m x_{CR} \quad (4)$$

y_{AE} : Obtained from Extract saturation curve (Knowing y_{CE} from tie line data). (5)

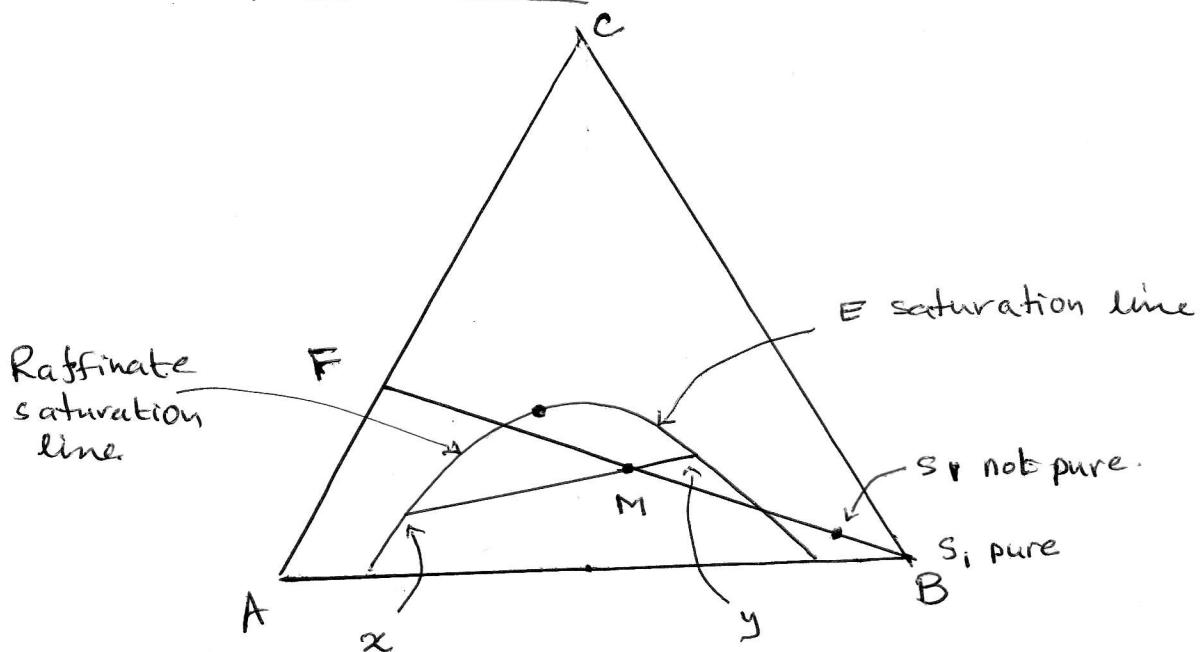
x_{AR} : Obtained from Raffinate saturation curve (Knowing y_{CR} from tie line). (6)

$$\sum x_i R = 1 \quad (7)$$

$$\sum y_i E = 1 \quad (8)$$

Solve system of equations (numerically).

Graphical Solution:



1. Locate points F and S (known compositions).
2. $F + S_1 \rightarrow M_1$. M_1 must be within the two phase region. It may be determined by:
 - lever rule (not recommended)
 - Calculating the composition of point M_1 , which falls on the line \overline{FS}_1 .

$$x_{M_1} = \frac{F \cdot x_F + S_1 y_S}{F + S_1}$$

If composition of M_1 is given, then read S_1 is

$$\frac{S_1}{F} = \frac{x_F - x_{M_1}}{x_{M_1} - y_S}$$

3. Mixture M_1 will separate into $E_1(y_1)$ in equilibrium with $R_1(x_1)$. Usually y_1 and x_1 are not known apriori; therefore a tie line that passes through M_1 must be found by trial and error; consequently y_1 and x_1 are determined.

4. Quantities E_1 and R_1 are then calculated as follows:

$$E_1 = M_1 \times \frac{(x_{M_1} - x_{E_1})}{(y_1 - x_1)}$$

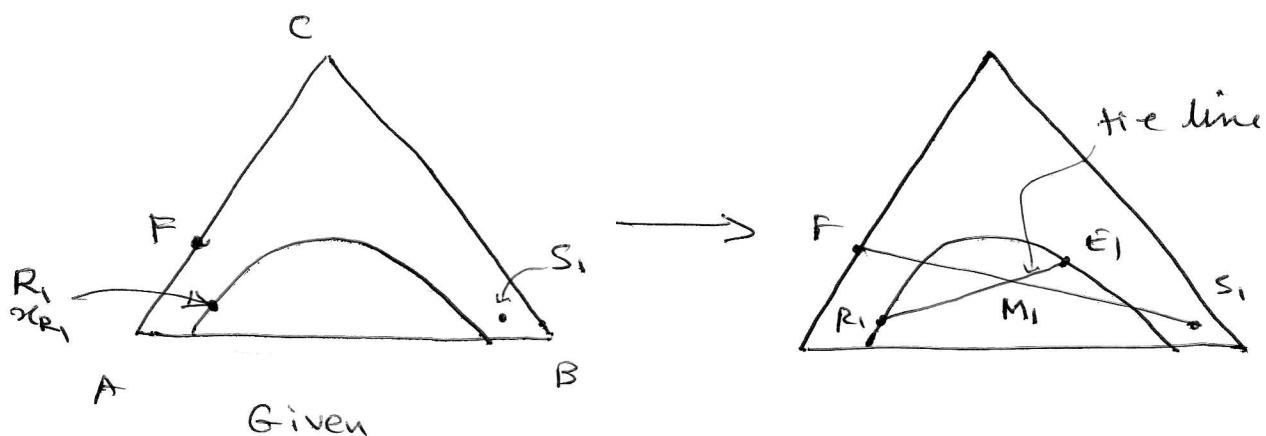
$$R_1 = M_1 - E_1$$

Case 2:

Known: F , $\overbrace{x_F}$ ^{solute}, y_S , $x_{A,F}$, $y_{A,S}$, x_{R_1}

Required S , E_1 , R_1 and outlet stream compositions.

Graphical solution!



Steps:- Draw tie line through R_1 ,

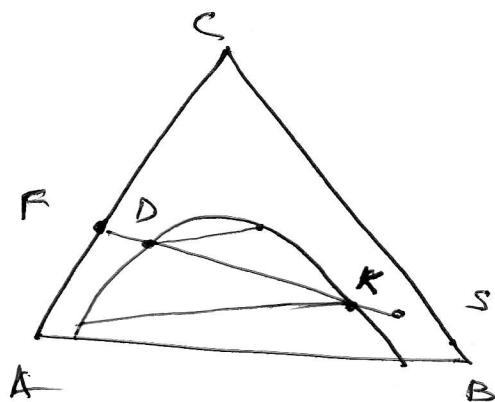
- tie line intersects with $\overline{FS_1}$ at M_1 ,
- Find S_1 (calculation or lever rule),
- E_1
- R_1
- compositions.

Notes:

- ① If solvent stream is pure then $y_{B,S_1} = 1$ and point S_1 would be at the B apex.
- ② For a given feed and solvent concentrations, the minimum amount of solvent required to form with F a 2φ mixture would be given by point "D", and the maximum amount of solvent allowable for 2φ mixture is given by point "K".

Min solvent \rightarrow very small amount of extract

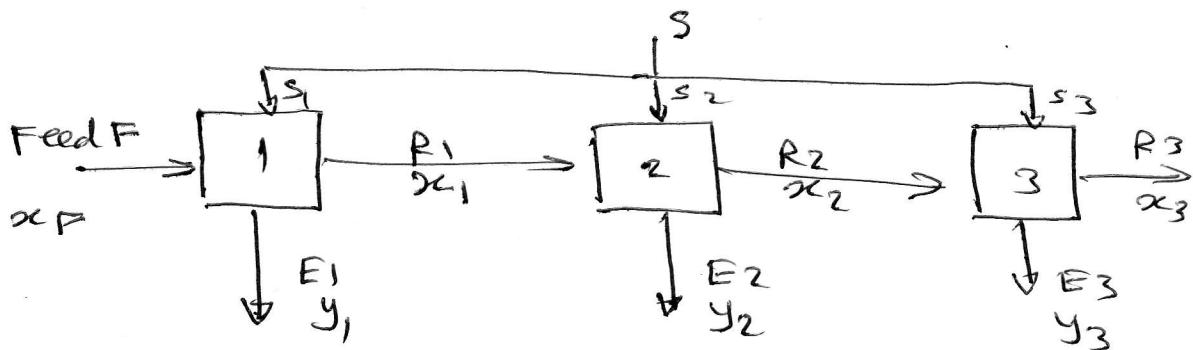
Max solvent \rightarrow very small amount of Raffinate



- ③ Same analysis can be carried out on ~~solute free~~^{solute} basis.

Cross Flow Contacting:

- Fresh solvent is fed to each stage
- Raffinate from one stage is fed to the subsequent stage
- Amount of solvent for each stage may be different
- Stages may be operated at different temperatures



The same analysis applied to single stage is used for each stage, thus:

- Each stage has an M point
- For each stage E and R are obtained by trial and error (in finding a tie line).
- All stages have a common S point, since the same solvent is used but feed is different amounts.
- Operating possibility:- Same $\frac{E}{S}$ ratio for each stage
 - Same amount of solvent
 - Different amounts of solvents

