

Summary

Principle 1

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PRINCIPLES

* Chemical Engineering: ^{مواد أولية} Raw materials ^{معالجة} Process ^{منتج} → Products

* In engineering: the numerical values of two quantities may be added or subtracted only if the units are the same.

eg. $3\text{cm} + 5\text{mg} = ???$ wrong equation

* A ^{كمية} measured or ^{عددية} counted quantity has:

- 1) Numerical value ^{قيمة عددية}
- 2) Unit ^{وحدة}

* A dimension is a property that can be:

1) Measured such as: time, mass, length ...

2) Calculated by multiplying or dividing other dimensions

such as: $\text{velocity} = \frac{\text{length}}{\text{time}}$, $\text{volume} = \text{length}^3$ and $\text{density} = \frac{\text{mass}}{\text{length}^3}$

* Units are important for effective communication and standardization of measurements: (Pounds vs. Kilograms)

* Base Units: 1) Mass = m 2) Length = L

3) Time = t 4) Temperature = T 5) Mole = n

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* Multiple Units: 1) Area = L^2 2) Volume = L^3

3) Velocity = L/θ 4) Acceleration = L/θ^2 5) Force = $m \cdot (L/\theta^2)$

6) Pressure = $m \cdot (L/\theta^2) / L^2 = m \cdot (L^3/\theta^2)$ 7) Energy = $m \cdot (L/\theta^2) \cdot L = m \cdot (L^2/\theta^2)$
compound units

* Derived Units:

1) Force: $1 \text{ kg} \cdot \text{m}/\text{s}^2 = 1 \text{ N} \xrightarrow{\text{Am}} 32.174 \text{ lbm} \cdot \text{ft}/\text{s}^2 = 1 \text{ lbf}$

2) Pressure: $1 \text{ kg}/(\text{m} \cdot \text{s}^2) = 1 \text{ N}/\text{m}^2 = 1 \text{ Pa} \xrightarrow{\text{Am}} 32.174 \text{ lbm}/(\text{ft} \cdot \text{s}^2) = 1 \text{ lbf}/\text{ft}^2$
 $= (1/144) \text{ lbf}/\text{in}^2 (\text{psi})$

3) Energy: $1 \text{ kg} \cdot (\text{m}^2/\text{s}^2) = 1 \text{ N} \cdot \text{m} = 1 \text{ J} \xrightarrow{\text{Am}} 32.174 \text{ lbm} \cdot (\text{ft}^2/\text{s}^2) = 1 \text{ ft} \cdot \text{lbf}$

* SI and American Engineering System Units:

Dimension	SI Unit	CGS Unit	Am. Eng. Unit
Mass	Kilogram (kg)	gram (g)	Pound mass (lbm)
Length	meter (m)	centimeter (cm)	foot (ft)
Time	Second (s)	Second (s)	Second (s)
Temperature	Kelvin (K)	Kelvin (K)	Rankine (R)
Mole	grammole (gmol)	gram mole (mol) (g-mol)	pound mole (lb-mol)

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Five Apple

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*) Conversion of Unit:

$$*) \text{ Old Unit} \cdot \left(\frac{\text{New unit}}{\text{old Unit}} \right) = \text{New Unit}$$

$$*) \frac{1}{\text{old Unit}} \left(\frac{\text{old unit}}{\text{New Unit}} \right) = \text{New unit}$$

eg: Express 300 cm^2 in mm^2 :

$$\text{ans: } \frac{300}{\text{cm}^2} \left(\frac{1 \text{ cm}}{10 \text{ mm}} \right)^2 = \frac{300}{\text{cm}^2} \left(\frac{1 \text{ cm}^2}{100 \text{ mm}^2} \right) = \frac{3}{\text{mm}^2} = 3 \text{ mm}^2$$

*) Force and Weight :

*) Newton's second Law: Force is proportional to the product of mass and acceleration ($\text{length}/\text{time}^2$)

*) Natural Force Units:

*) Derived Force Units:

$$1) \text{ SI} = \text{kg} \cdot \text{m}/\text{s}^2 =$$

1 Newton (N)] conversion

$$2) \text{ CGS} = \text{g} \cdot \text{cm}/\text{s}^2 =$$

1 dyne

$$3) \text{ Am. eng} = \text{lb} \cdot \text{ft}/\text{s}^2 =$$

$\frac{1}{32.174}$ lbf

Factors between natural and derived force units.

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*) Weight: force exerted on the object by gravitational attraction ($W = mg$) $g = 9.8066 \text{ m/s}^2 = 980.66 \text{ cm/s}^2 = 32.174 \text{ ft/s}^2$

*) Scientific Notation: a number is expressed as the product of another number (usually between 0.1 and 10) and a power of 10.

e.g: $123000000 = 1.23 \times 10^8$ or 0.123×10^9

e.g: $0.000028 = 2.8 \times 10^{-5}$ or 0.28×10^{-4}

*) Significant Figure: the digits from the first nonzero digit on the left to either:

a) the last digits (zero or nonzero) on the right if there is a decimal point, or (b) the last nonzero digit of the number if there is no decimal point.

e.g: $23,040 = 2.304 \times 10^4 = 4 \text{ Sig. fig}$

e.g: $0.035 = 3.5 \times 10^{-2} = 2 \text{ Sig. fig}$

*) In Multiplication / Division: The number of sig. fig in the product or quotient is equal to the lowest number of sig. fig of any of number.

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e.g: $3.23 \times 4.125429 = 13.32513567 = \underline{\underline{13.3}}$

$3 \text{ sig. fig} \times 7 \text{ sig. fig} = 3 \text{ sig. fig}$

*) In Addition / Subtraction:

1) Compare position of last sig. fig relative to the decimal point.

بالأرقام الأقرب للفاصلة مع تقريب باقي الأعداد قبل الفاصلة.

2) The position farthest to the left is the last permissible

sig. fig for the sum or difference.

e.g: $3.\underline{\underline{23}} + 4.\underline{\underline{125429}} = 7.\underline{\underline{355429}} = \underline{\underline{7.36}}$

تقريب
العدد 3
العدد 6

*) Dimensionless Quantities:

*) Reynolds Number = $\frac{\rho D u}{\mu}$ = $\frac{\text{Kg/m}^3 \cdot \text{m} \cdot \text{m/s}}{\text{Kg/m.s}} = \frac{\text{Kg} \cdot \text{m} \cdot \text{m} \cdot \text{m/s}}{\text{m}^3 \cdot \text{m.s}} = \frac{\text{Kg} \cdot \text{m} \cdot \text{m} \cdot \text{m/s}}{\text{Kg} \cdot \text{m} \cdot \text{s}} = 1$

*) Importance of Dimensionless Quantities:

*) Used in arguments of special functions such as exponential, logarithmic, or trigonometric functions.

e.g: e^{10} is ok but $e^{10\text{ft}}$ is undefined.

e.g: $\cos(10)$ is ok but $\cos(10\text{ft})$ is undefined.



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* Dimensionless Quantities:

* Consider the Arrhenius Equation: $K = A \cdot \exp(E/RT)$

Where: 1) E = Activation energy (cal/mol)

2) T = Temperature (K)

* What is the Unit of R ?

ans: $\frac{E}{RT}$ is exp $\rightarrow \frac{E}{RT} = 1 \rightarrow \frac{\frac{\text{cal}}{\text{mol}}}{R \cdot K} = 1 \rightarrow \frac{\text{cal}}{K \cdot \text{mol}} = R \cdot \frac{K \cdot \text{mol}}{K \cdot \text{mol}}$

$R = \frac{\text{cal}}{K \cdot \text{mol}}$ To make the argument of the exponential function dimensionless.

* Process Data Representation and Analysis:

* Experiment: Collection of (measurements) data, each represented by number which give us information about the quantity being determined. (The number may occur once or it may be repeated)

* Errors: Personal / insomantal

* This is due to Variability of measurements arising

form inaccuracies i.e. errors

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* We display measurements as: Frequency Distribution. توزيع التردد

e.g: What is the true value of the quantity being measured call it x (Random variable) N measurements?

ans: We have estimate of the true value X Sample mean

(arithmetic mean): $\bar{X} = \frac{1}{N} \sum_{i=1}^N x_i$ مجموع القراءات تقسيم عددهم

* Scatter of data (Variable X) $i=1$ with respect to their mean value can be characterized by: مقدار رقة القيم الناتجة

1) Range 2) Sample Variance 3) Sample Standard Deviation

$$R = X_{\max} - X_{\min} \quad \downarrow \quad \begin{matrix} \text{أقصى} - \text{أدنى} \\ \text{(الحد أعلى - الحد أسفل)} \end{matrix}$$

$$s_x^2 = \left(\frac{1}{N-1} \right) \left[\sum_{i=1}^N (x_i - \bar{x})^2 \right] \quad \begin{matrix} \text{مجموع مربع} \\ \text{الفرق بين} \\ \text{القراءة والناتج} \\ \text{مقسوم} \\ \text{على عدد القراءات} \\ \text{- واحد} \end{matrix}$$

$$s_x = \sqrt{s_x^2} \quad \begin{matrix} \text{نسبة} \\ \text{مستخدمة لإيجاد Error} \end{matrix}$$

* Value of measured value: $X = \bar{X} \pm \text{Error} \rightarrow \begin{matrix} = (X_{\max} - X_{\min}) \\ \text{or} \\ = (s_x + 2s_x + 3s_x) \end{matrix}$

* Process Data Representation And Analysis:

* The operation of any chemical process is ultimately based on the measurement of Process variables. متغيرات العملية

Such as: Temperatures * pressures * Flow rates * Concentrations

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* It is sometimes possible to measure these variables Directly, but as a rule Indirect techniques must be used.

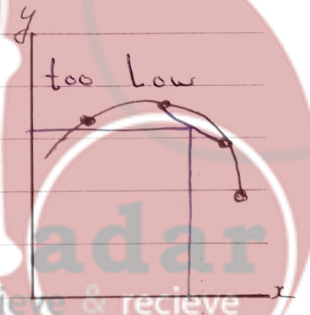
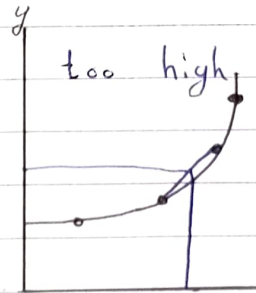
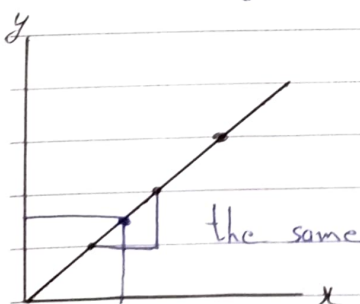
e.g. To measure the concentration (C) of a solute in a solution, we normally measure a quantity (X) such as thermal or electrical conductivity, a light absorbance or the volume of titration, and then calculate (C) from the measured value of (X) Calibration Curve

(* مستخرج قيمة التركيز من قيمة (X) عن طريق العلاقة الخطية على شكل نقاط)

* Our object is to use the calibration data to estimate the value of for a value of between tabulated points (interpolation) or outside the range of the table data (extrapolation).

* Two point linear interpolation: $y = y_1 + \frac{x - x_1}{x_2 - x_1} (y_2 - y_1)$

* If the function graph $y(x)$ contains an inclination, the values of y become:



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* Linear Regression: $y = mx + b$, Where:

y = dependent x = independent m = slope b = intercept

$$\text{e.g.: } y = \frac{1.3 \times 10^7 \sin(2x)}{x^{\frac{1}{2}} + 58.4} \rightarrow \frac{1}{y} = \frac{x^{\frac{1}{2}} + 58.4}{1.3 \times 10^7 \sin(2x)}$$

$$= \frac{\overset{y\text{-axis}}{\sin(2x)}}{\underset{y}{y}} - \frac{\overset{x\text{-axis}}{x^{\frac{1}{2}}} + 58.4}{1.3 \times 10^7}$$

* What is regression analysis?

* A technique for modeling and analyzing the relationship between 2 or more variables. Usually, 1 variable is

* Purpose: Establish a mathematical relation between the variables.

* Fitting a straight line to a set of data points

calculating the slope m and intercept b using the method

$$\text{of least squares: } m = \frac{n(S_{xy}) - (S_x S_y)}{n(S_{xx}) - (S_x)^2} = \frac{\sum_{i=1}^n (x_i - \bar{x})(y_i - \bar{y})}{\sum_{i=1}^n (x_i - \bar{x})^2}$$

$$b = S_y - m \cdot S_x = \bar{y} - m \cdot \bar{x}$$

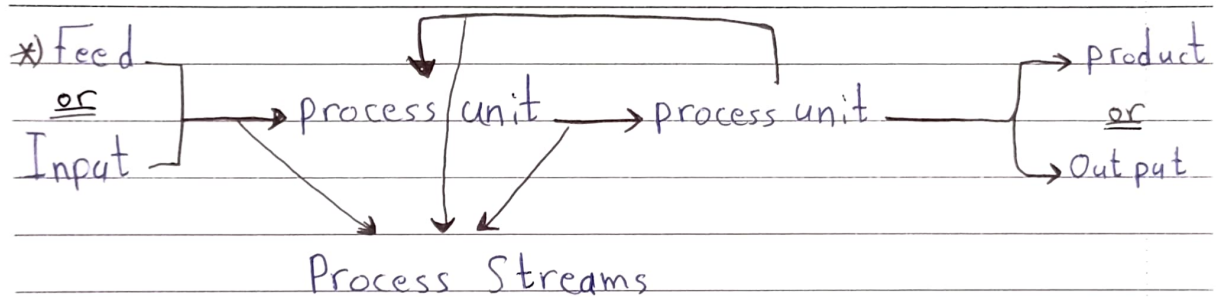
$$S_x = \sum_{i=1}^n x_i \quad S_y = \sum_{i=1}^n y_i \quad S_{xx} = \sum_{i=1}^n x_i^2 \quad S_{xy} = \sum_{i=1}^n x_i y_i$$

PRINCIPLES

*) Chemical Process:

*) A process is any operation or series of operations by which a particular objective is accomplished.

*) A chemical process is any operation or a series of operations by which a particular product is produced from raw materials.



*) What we want about these streams?

ans: Temperature * pressure * mass flow * volumetric flow * chemical composition.

*) Some of those variables will be given, others we measure

But many we have to calculate such as (mass flow and

material balance) CH_4 (energy balance, heating, cooling, etc.)

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*) The Process unit : modeling * designing * sizing
costing unit operations .

*) What is a (chemical) _{C.P} process variable ?

*) Any measurement used to characterize or describe (C.P)

we have : 1) measurements to quantify a material or
specify a chemical composition (Mass * Volume * Mole)
(لتقدير مادة أو تركيبة كيميائية)

2) measurements used to specify process conditions (Pressure * Temp.)
(لتحديد شروط العملية)

*) Density (P) : mass per unit volume

$$P = [Kg/m^3] = [lbm/ft^3] = [g/ml]$$

*) P : Used as conversion factor between (mass and volume) or
(يستخدم التركيز للتحويل بين الكتلة والحجم)
(سرعة تدفق الكتلة / سرعة تدفق الحجم)
(Flow rate and volumetric flow rate of a process stream)

*) Liquid and solid densities vary slightly with Temperature
(كثافة سائل / كثافة صلب)
(تغير بشكل قليل مع الحرارة)

*) Density of gas depend heavily on temperature and pressure.
(كثافة غاز)

* Specific volume : volume occupied by unit mass = $\frac{1}{\text{density}}$

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*) Specific gravity: Ratio of a density of a substance to the density of a reference substance ($SG = \rho / \rho_{ref}$)

*) Typical Reference Density = water at 4°C and 1 atmosphere

*) Value: $\rho_{ref} = 1.000 \text{ g/cm}^3 = 1000 \text{ kg/m}^3 = 62.43 \text{ lbm/ft}^3$

*) SG is given at particular temperature (does not have to be the reference temperature)

*) To get density (ρ) from SG: $\rho_A = (SG_A)(\rho_{H_2O, ref})$

$$\rho_H = \frac{\rho}{\rho_{H_2O, ref}} \rightarrow \frac{m}{V} = \frac{m}{V}$$

*) The kinetic energy of substance produces:

- 1) Potential energy.
- 2) Temperatures.

*) Temperature units:

- 1) Kelvin (K)
- 2) Celsius ($^\circ\text{C}$)
- 3) Fahrenheit ($^\circ\text{F}$)
- 3) Rankine ($^\circ\text{R}$)



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* Temperature conversions:

$$1) K = C^{\circ} + 273.15$$

$$2) R = F + 459.67$$

$$3) R = 1.8 K$$

$$4) F = 1.8 C^{\circ} + 32$$

$$5) \Delta C = \Delta K$$

$$6) \Delta F = \Delta R$$

$$7) \Delta C = 1.8 \Delta F$$

* Pressure = Force / Area

Atmospheric
وزن عاصم الهواء
الساكن

Absolute
⊕ always

Gauge

$$P_{\text{Absolute}} = P_{\text{Atmospheric}} + P_{\text{Gauge}} \quad * \quad P V = R n T$$

* The average molecular weight:

$$1) \bar{M} = y_1 M_1 + y_2 M_2 + \dots$$

$$2) \frac{1}{\bar{M}} = \frac{x_1}{M_1} + \frac{x_2}{M_2} + \dots$$

* Fluid pressure in:

$$1) \text{Tank and a pipe} \Rightarrow P = P_0 + \rho g h$$

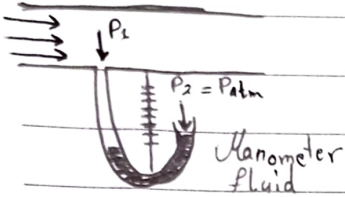
$$2) \text{At the base of a fluid column} \Rightarrow P\left(\frac{L}{A}\right) = \rho_{\text{fluid}} P_h (\text{head of fluid})$$

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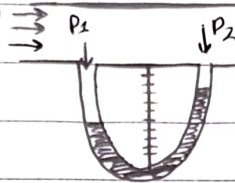
PRINCIPLES

* Manometers:

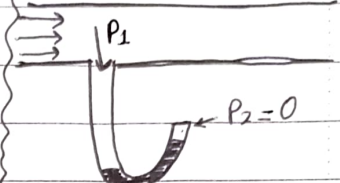
1) Open-end



2) Differential



3) Sealed-end



* General Manometer Equation: $P_1 + \rho_1 g d_1 = P_2 + \rho_2 g d_2 + \rho g h$

* Differential: $P_1 - P_2 = (\rho_f - \rho) g h$

* For gases: $P_1 - P_2 = h$

* The pressure in a manometers full of mercury is given by (The difference in length) mercury or any matter



PRINCIPLES

* Process classification :

- | | | |
|---|--|--|
| <p>1. Batch process.</p> <p>* inputs: ones (الشيء)</p> <p>* outputs: ones (الشيء)</p> <p>* process: in a close system (نظام مغلق)</p> | <p>2. Continuous Process.</p> <p>* inputs: continuously (مستمر)</p> <p>* outputs: continuously (مستمر)</p> <p>* process: in a open system (نظام مفتوح)</p> | <p>3. Semi batch Process.</p> <p>* inputs: (one of them continuously and the other one)</p> <p>* outputs: (one of them continuously and the other one)</p> <p>* process: semi open</p> |
|---|--|--|

* Another classification of process :

- | | | |
|---|--|---|
| <p>1. Steady-state ثوابت</p> <p>* If the values of all variables in a process don't change with time.</p> | <p style="border: 1px solid black; border-radius: 50%; padding: 10px; display: inline-block;"> * Variables: volumes *
 pressure * Flow rates
 temperature </p> | <p>2. Unsteady-state متغير</p> <p>* If any of the process variables change with time.</p> |
|---|--|---|

* Unsteady state such as → Batch * Semi batch * Continuous.

* Steady state such as, Continuous.

* General balance equation :

$$\text{Input} + \text{generation} - \text{output} - \text{consumption} = \text{accumulation}$$

عزول
التوليد
مخرج
استهلاك
التراكم

* Types of balance :

Differential indicates what is happening in a system at an instant in time.

(Continuous steady system)

Per time

$$\Delta t \rightarrow 0$$

Integral describe what happens between two instants of time.

(Batch system)

(repeated)

$$t_i \quad t_f$$

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PRINCIPLES

*) Important Rules :

*) If the balanced quantity is the total mass: ^{إذا كانت الكمية المحسوبة هي الكلية}

1) generation = 0 2) consumption = 0

^{بالمستثنى = التفاعلات النووية}
except in nuclear reactions, mass can't be created or destroyed.

*) If the balanced substance is a non-reactive species (neither a reactant nor a product): 1) generation = 0 2) consumption = 0.

*) If a system is at steady state, accumulation = 0, regardless ^{بغض النظر}
of what is being balanced. By definition in a steady state system ^{لا يمكن أن يتغير شيء بمرور الوقت. يعني ذلك الكميات}
nothing can change with time including the amount of the quantity.

*) Balances on continuous steady state processes:

*) accumulation = 0 $\Rightarrow$ input + generation = output + consumption

^{مواد غير متفاعلة}
*) if it is on a non-reactive species or total mass $\Rightarrow$ input = output

*) Integral balances: initial input + gen = final output + consumption

*) Material balance calculation: the best way to solve it by drawing the system.

PRINCIPLE

* V.I always mass in = mass out (conservation of mass) ^{في الكمية (قانون)}

*) The best way to solve any question:

1) Find what is the question asking for such as ^{افتراضات} key words & phrases ^{عبارات} and be careful of assumptions.

2) Draw ^{ارسم} a picture for the system and locate the stem.

3) Label ^{حدد} Knowns + unknowns in the picture ^{* افترض شئ ب رسمه جديد و على نفس الرسمه}

as example put: $m \rightarrow \text{mass} * n \rightarrow \text{moles} * x \rightarrow \text{mass} . f * y \rightarrow \text{mole} . f$

4) Write ^{اكتب} a mass balance ^{تذكر} remember that you can

Write an 1 mass balance for every component ^{معادلة لكل مادة}

ex: $m_1 + m_2 + m_3 \dots \overset{\text{in}}{=} m_1 + m_2 + m_3 \dots \overset{\text{out}}{=}$
1stem 2stem 3stem 1stem 2stem 3stem

*) then write overall balance and substitute 1 overall balance for 1 component balance.

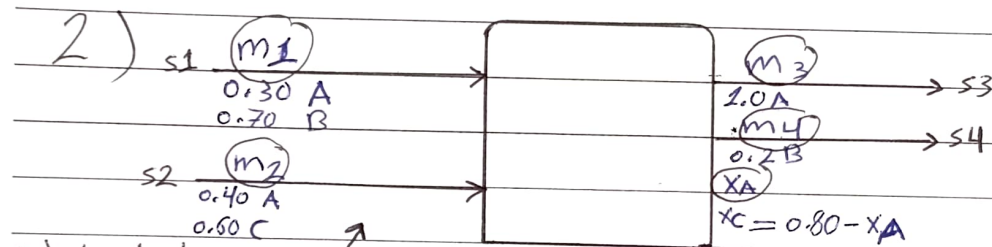
*) and be careful of when you do your mass balances ^{كل واحد} that each one of your balances is independent ^(مستقل)

*) After no 4 you should calculate the DOF ^{يجب حساب قبل ذهاب} ^{ask :: before & recieve}

PRINCIPLE

* EX: Two streams enter a column. Stream 1 is 30% A and the rest is B. Stream 2 is 40% A and the rest is C. Two streams exit the column. Stream 3 is 100% A, while stream 4 is 20% B and the rest is A and C.

An: 1) we have 4 stream * 3 component * 1 column and asking for m.b. with 4 information



* we have 5 unknowns.

3) Labeling

4) A: $0.3m_1 + 0.40m_2 = m_3 + X_A m_4$

B: $0.7m_1 = 0.2m_4$

C: $0.60m_2 = X_C m_4$

$m_1 + m_2 = m_3 + m_4$

" by you because us
* if we add all of it together we will

Get overall balance
so we can use 3 of them to solve the system.

* In DOF we have 5 unknowns and 3 balances So we can't solve the system.

PRINCIPLE

*) DOF $\rightarrow$ Degree Of Freedom $\rightarrow$ It is show as if we have enough or to much information to solve problem.

* $\text{DOF} = n_{\text{unknown}} - n_{\text{eq}}$ $\leftarrow$ (Just for non-reactive)

$$= N_{\text{unknown}} - N_{\text{independent balance}} - N_{\text{other eq}} = N_{\text{species}}$$

If

↓
Process Specification

↓
Physical Properties
data

LO

$$> 0$$
$$= 0$$

* over specified

- * under specified

* The problem can be solved but there are ~~an~~ be solved.

* the problem solved

more than wenced of equ

* When we have more than 1 process or column in the problem we should calculate the DOF for each one of

then then start from $\text{DOF} = 0$

* Splitting point : a point have the same concentration like Produce in other stem
 (e.g. a cup of Juice)
 (e.g. a cup of Juice)

*) Mixing point: a point produce a new concentration in every stream. (in Recycle system)

PRINCIPLE

*) Chemical reaction stoichiometry (on reactive system):
النظام التفاعلي *قياس النسب الكيميائية للتفاعل*

*) The stoichiometric coefficients: tell us the mole
(not mass) relationships (stoichiometric ratios) of the
النسب المولية
compounds participating in the reaction.

eg: $2A + 3B \longrightarrow 5C + 7H \Rightarrow$ mean: 2 g mol of A
reacts with 3 g mol of B to yield 5 g mol of C and
7 g mol of H.

*) Mole $\Rightarrow$ is Avogadro's number of particles $= 6.23 \times 10^{23}$

$$\text{Mole} = \frac{\text{Mass}}{\text{Molecular Weight}} \rightarrow \frac{m}{M_w}$$
 and we use it because
it is fixed number and easy to deal with.

*) we use Mole instead of number of molecules.

*) In balanced chemical equation: Atoms in = Atoms out

*) Stoichiometric coefficient: The numbers that precede
the formulas for each species.

1) negative *سالب* for reactants *للتفاعلات* 2) Positive *إيجابي* for products *للمنتجات*

PRINCIPLE

*) Fractional excess: is the ratio of the amount by which the feed exceeds stoichiometric requirements in moles divided by the stoichiometric requirements in moles.

So Fractional excess of A = $\frac{(n_A)_{\text{feed}} - (n_A)_{\text{stoich}}}{(n_A)_{\text{stoich}}}$

*) Chemical reactions proceed quite slowly.

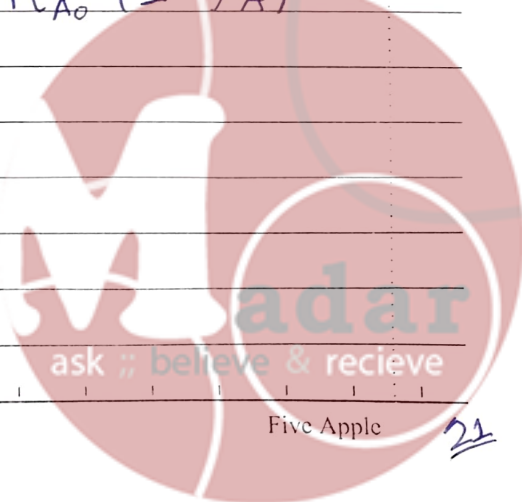
*) It is not practical to design a reactor for complete conversion of the limiting reactant. Instead, the reactant is separated from the reactor outlet stream and recycled back to the reactor inlet.

*) Fractional conversion (f) = moles reacted / moles fed.

*) In general, for $A + \frac{b}{a} B \rightarrow \frac{c}{a} C + \frac{d}{a} D$

$$f_A = \frac{n_{A0} - n_A}{n_{A0}} \rightarrow n_A = n_{A0} (1 - f_A)$$

$$n_C = n_{C0} + \frac{c}{a} n_{A0} f_A$$



PRINCIPLE

*) The extent of reaction (ξ or ξ) is the amount in mole that converted in a given reaction normalized to Stoichiometric

For batch process: $n_i = n_{i0} + \nu_i \xi$ where

- 1) n_i = mole of i (component)
- 2) n_{i0} = mole of i (species)
- 3) ν_i = stoichiometric coefficient

*) The (ξ) has the same unit of (n) (mole, kg-mol...)

2) For continuous processes, $\dot{n}_i = \dot{n}_{i0} + \nu_i \dot{\xi}$ where

- 1) $\dot{n}_i$ = molar flow (existing)
- 2) $\dot{n}_{i0}$ = molar flow (fed)
- 3) ν_i = stoichiometric coefficient

*) The ($\dot{\xi}$) has the same unit of ($\dot{n}$) (mole/time)

$$\text{So, } 1) \xi = \frac{n_i - n_{i0}}{\nu_i}$$

$$2) \dot{\xi} = \frac{\dot{n}_i - \dot{n}_{i0}}{\nu_i}$$

*) The final (equilibrium) composition of the reaction mixture is determined by Chemical Thermodynamic

*) The time needed for a system to reach a specified state short of equilibrium is determined by

Chemical Kinetic.

*) Reactions:

Reversible (Reactants $\rightleftharpoons$ Products)

*) Reactants from products and products undergo the reverse reactions to reform the reactants.

(Reactants $\rightarrow$ Products)

Irreversible

*) The reaction proceeds only in a single direction and the concentration of the limiting reactant eventually approaches Zero

PRINCIPLE

* In the reaction: $aA + bB \longrightarrow cC + dD$

if all compon is gases then the eqn constant (K) =

$$K = \frac{P_C^c \times P_D^d}{P_A^a \times P_B^b}$$

products
reactants

Where:

1) P is the partial pressure for gasses.

$$P_A = y_A P \longrightarrow K = \frac{y_C^c y_D^d}{y_A^a y_B^b}$$

total pressure
mole fraction

* In many processes, reactants are put together in order to make a product.

* Yield: $\frac{\text{mole of desired product formed}}{\text{mole formed at 100\% conversion}}$ (fraction) * عدد المولات الناتجة

* Selectivity: $\frac{\text{mole of desired product formed}}{\text{mole of undesired product formed}}$ (not fraction) * عدد المولات الناتجة في العملية
دون فقد أو تحويل
* عدد المولات الناتجة من المفاعل
* عدد مولات المواد الداخلة في التفاعل

* For multiple reactions, each independent reaction has its own extent.

* If a set of reaction takes place in a batch or continuous steady-state reactor: $n_j = n_{j0} + \sum \tau_j \nu_j$ where j is a reaction

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