

**INSTRUCTIONS**

1. This question paper consists of two parts. Part A is close book and Part B is open **(only text)** book.
2. Part-B answer book will be supplied after you return Part-A answer book.
3. Make and state suitable, logical and justifiable assumptions if necessary.
4. Give just 2 iterations for iterative procedure(s).  
**Be to the point. Do not be descriptive. Use as less words as possible.**

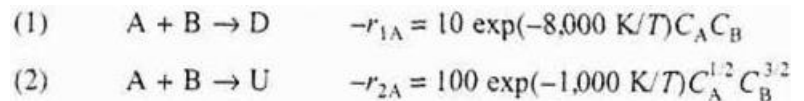
**PART A (CLOSE BOOK)**

**Q1 [Marks 25]** (a) Give the CRE algorithm [1]; (b) List 5 criteria used to evaluate laboratory reactors [5]; (c) We can use membrane reactors to increase the selectivity in multiple reactions (*True or False*) [1]; (d) Justify: “*in no other type of reaction is exactness in the calculation of the time needed to carry out the reaction more important than in series reactions*” [3]; (e) Define turnover frequency and dispersion in the context of catalysis [2]; (f) Use well-labelled schematics to describe 3 surface reaction models [5]; (g) What do 3 main moments of RTD signify? [3]; (h) Which 5 models help us to predict the conversion (in non-ideal reactors) from RTD data? Categorize them in 3 groups [5].

**PART B (ONLY OPEN TEXT BOOK)**

**Q2 [Marks 20]** Compound A undergoes a reversible isomerization reaction,  $A \rightleftharpoons B$ , over a supported metal catalyst. Under pertinent conditions, A and B are liquid, miscible, and of nearly identical density; the equilibrium constant for the reaction (in concentration units) is 5.8. In a fixed-bed isothermal flow reactor in which back mixing is negligible, a feed of pure A undergoes a net conversion to B of 55%. The reaction is elementary. If a second, identical flow reactor at the same temperature is placed downstream from the first, what overall conversion of A would you expect if the reactors are directly connected in series?

**Q3 [Marks 20]** For the given (to your RHS) set of reactions, describe 3 reactor systems and conditions to maximize the selectivity to the desired product D. Make sketches (schematics) to support your choices. The rates are in (mol/dm<sup>3</sup>.s), and concentrations are in (mol/dm<sup>3</sup>). Say: T<sub>1</sub> and T<sub>2</sub> = 300 and 1000 K.



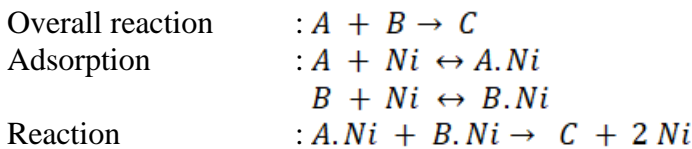
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**Q4 [Marks 15]** The endothermic liquid-phase elementary reaction  $A + B \rightarrow 2C$  proceeds, substantially, to completion in a single steam-jacketed, continuous-stirred reactor. From the following data, calculate the steady-state reactor temperature: Reactor volume: 125 gal; Steam jacket area: 10 ft<sup>2</sup>; Jacket steam: 150 psig (365.9°F saturation temperature); Overall heat-transfer coefficient of jacket,  $U$ : 150 Btu/h.ft<sup>2</sup>.°F; Agitator shaft horsepower: 25 hp; Heat of reaction,  $\Delta H^0_{R_x} = +20,000$  Btu/lb mol of A (independent of temperature).

	Component		
	A	B	C
Feed (lbmol/hr)	10.0	10.0	0
Feed temperature (°F)	80	80	—
Specific heat (Btu/lb mol·°F)*	51.0	44.0	47.5
Molecular weight	128	94	222
Density (lb/ft <sup>3</sup> )	63.0	67.2	65.0

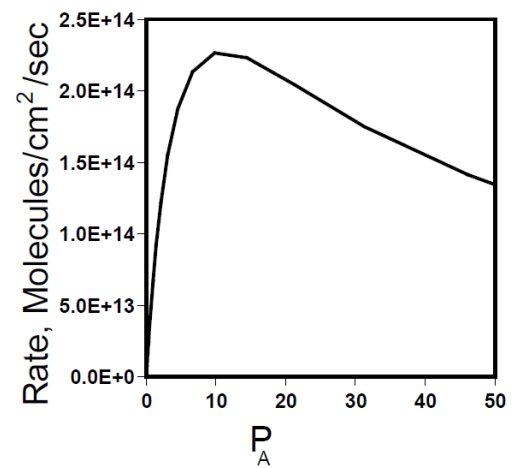
\*Independent of temperature.

**Q5 [Marks 20]** The following reactions take place on a nickel catalyst (spherical particles):



Determine the rate determining step (RDS) and rate of formation of C ( $r_C$ ) in consistent with the experimental data. (1 mole = 6.023 x 10<sup>23</sup> molecules).

**Figure (RHS):** A plot of the experimental (formation) rate of C ( $r_C$ ) with  $K_B P_B = 10$ . ( $K_B$  = Adsorption constant of B,  $P_B$  = partial pressure of B,  $P_A$  = partial pressure of A)



**Q6 [Marks 20]** The flow through a reactor is 10 dm<sup>3</sup>/min. A pulse test gave the concentration measurements at the outlet as given in the table to your right:

- (a) Plot the external age distribution  $E(t)$  as a function of time.  
 (b) What are the mean residence time  $t_m$ , and the variance,  $\sigma^2$ ?

$t$ (min)	$c \times 10^5$	$t$ (min)	$c \times 10^5$
0	0	15	238
0.4	329	20	136
1.0	622	25	77
2	812	30	44
3	831	35	25
4	785	40	14
5	720	45	8
6	650	50	5
8	523	60	1
10	418		

**END**