

B.M.S. College of Engineering, Bengaluru-560019

Autonomous Institute Affiliated to VTU

May 2023 Semester End Make-Up Examinations

Programme: B.E.

Branch: Biotechnology

Course Code: 19BT7DE5MTE

Course: Metabolic Engineering

Semester: VII

Duration: 3 hrs.

Max Marks: 100

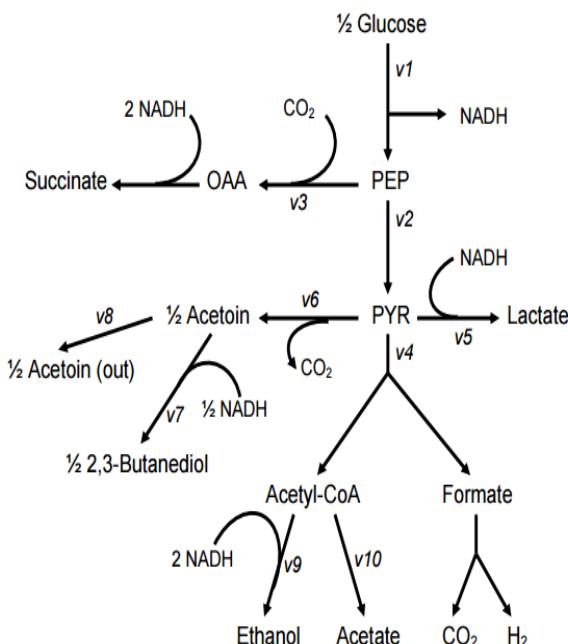
Date: 17.05.2023

Instructions: 1. Answer any FIVE full questions, choosing one full question from each unit.
2. Missing data, if any, may be suitably assumed.

Important Note: Completing your answers, compulsorily draw diagonal cross lines on the remaining blank pages. Revealing of identification, appeal to evaluator will be treated as malpractice.

1 a)

10



Shown above is the flux pattern for the anaerobic metabolism of *Enterobacter aerogenes*. Give the rate equation in matrix notation for the labelled reaction rates v1-v10.

- b) Differentiate between metabolome, fluxome & proteome. 06
- c) Illustrate the mechanism of group translocation taking an example of PTS systems. 04

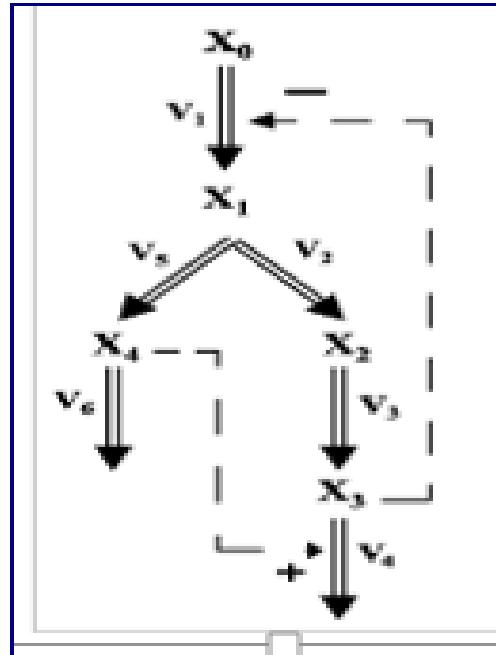
UNIT - II

2 a) What is branch point effect? Explain taking a suitable example.

06

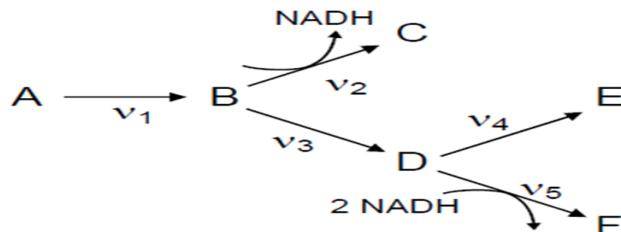
b)

04



In the given pathway, identify the mode of inhibition. Justify your answers.

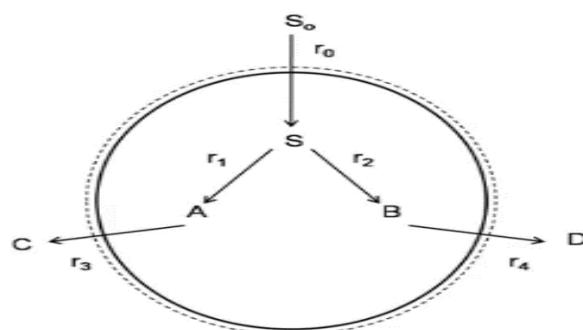
c) In the following pathway, assuming pseudo steady state for metabolites B, D and NADH, calculate fluxes v_2 , v_3 , & v_4 , given that v_1 is 8 mm/g/sec and v_5 is 2 mm/g/sec.



d) Cross regulation (cross talk) helps in control of pathways of central carbon metabolism. Justify this statement taking the example of Pho regulon.

OR

3 a) Differentiate between stimulon, regulon and modulon. **06**
 b) Give the equation which represents coordinated control of flux by energy charge & reduction charge. **06**
 c) Determine the intracellular metabolite flux assuming that the extracellular metabolite rates for the given pathway are known **08**

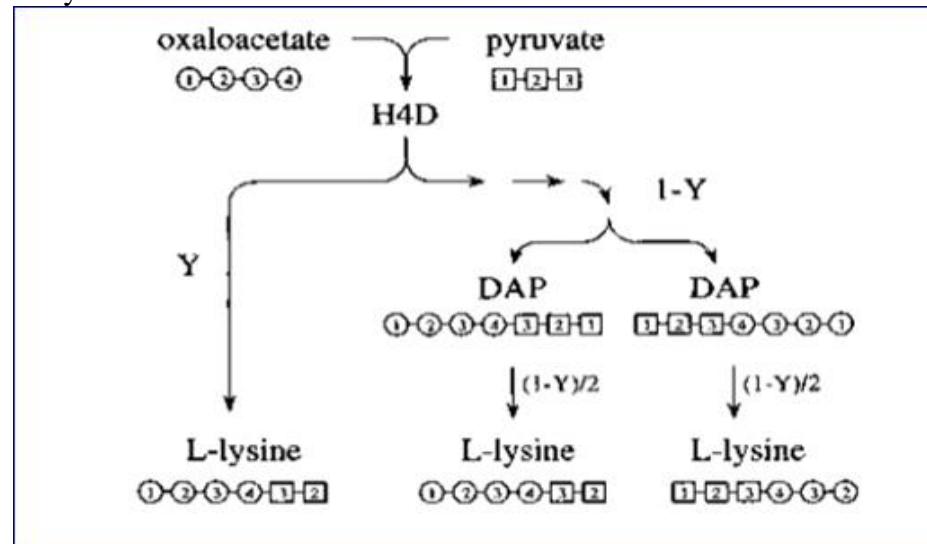


UNIT - III

4 a) Determine the theoretical yield of lysine biosynthesis by stoichiometric balancing and by considering the cofactor requirements. **08**

b) Construct the atom mapping matrices for the synthesis of lysine in the below pathway

08



c) What is endogenous metabolism? Give the relationship between specific substrate uptake rate and specific growth rate taking into consideration endogenous metabolism.

04

UNIT - IV

5 a) In an experiment w.r.t tryptophan metabolism, the effects of various levels of tryptophan 2,3, dioxygenase was measured. The enzyme activities (v_{max}) were found to be $30\mu\text{M}/\text{hr}$ & $58\mu\text{M}/\text{hr}$ and the corresponding fluxes were 4.6 and $6.7\mu\text{M}$. Given that pathway flux was $2.6\mu\text{M}$ under basal conditions with a measured activity of $13.7\mu\text{M}$ for the enzyme and an FCC of 0.77. Are the results predicted above consistent? 05

b) Define the theories of MCA. 04

c) Consider the pathway $X_0 \rightarrow Y \rightarrow Z \rightarrow X$. Xase, Zase and Yase are the enzymes respectively. Predict the changes in overall flux if additional amount of enzyme Yase is added. 06

d) What is Pathway Shortening? What are its drawbacks. 05

OR

6 a) With an example explain the approach “double modulation” for determination of the elasticity coefficients? List any four limitations of this method. 08

b) What is elasticity co-efficient? How is it measured graphically? Why is it an important parameter in MCA? 06

c) What is response coefficient? What are the factors that affect the response co-efficient? Consider the pathway, $x \rightarrow y \rightarrow z$, where an external metabolite “p” acts as a regulator for an enzyme “e1” which catalyzes the conversion of $x \rightarrow z$. what is the response co-efficient of “p”. 06

UNIT - V

7 a) “Multivariate-modular pathway engineering” is a strategy used for overproduction of complex natural products. Justify the statement with a suitable example. 10

b) Explain how metabolic engineering of *Escherichia coli* can be done to increase NADH availability. 10
