

Q.2 Answer the following (Any seven) (14)

1. Explain different types of nodes based on branch point rigidity.
2. Define metabolism.
3. What is singularity?
4. Briefly explain summations theorem.
5. Briefly explain E4P is rate limiting step.
6. Schematically present strategies for PHA production.
7. Briefly explain increasing product selectivity in antibiotic synthesis.
8. Briefly describe the properties of yeast as a host for expression of genes
9. What is central dogma?

- Q.3**
- a) Enlist various cellular transport process and give detailed note on active transport system. (06)
 - b) What is metabolic network? How cell is accessing the cellular conditions for regulation of metabolic network? (06)

OR

- b) State and derive Briggs-Haldane equation for steady state assumption. (06)

- Q.4**
- a) Give comparative account on change in flux distribution in *E. coli* for the production of DAHP from PTS and non-PTS sugar. (06)
 - b) What is metabolic control analysis? Explain in detail flux control coefficient. (06)

OR

- b) Explain the perturbation of nitrogen regulation in *E. coli* on over expression of PCK gene. (06)

- Q.5**
- a) Write a detailed note on metabolic engineering of β -lactam antibiotics biosynthesis by increasing enzyme activity. (06)
 - b) Discuss in detail about solventogenic pathway in *Cl. acetobutylicum* (06)

OR

- b) Explain in detail production of L-Cysteine with the metabolic engineering of sulfur incorporation (06)

- Q.6**
- a) Discuss the degradation of toluene by *Pseudomonas putida* mt-2. (06)
 - b) What is system biology? Explain the role of system biology in metabolic engineering. (06)

OR

- b) Write a detail note on DNA microarray fabrication. (06)

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