



(ix) Plants do not possess  $\delta$  amino levulinic acid synthetase activity yet porphyrins are required for the synthesis of chlorophyll.

(x) Polyamines are required for cell proliferation. (2×9+1=19)

2. Write short notes on the following :

(i) Purine nucleotide cycle

(ii) Glucose alanine cycle

(iii) Regulation of biosynthesis of deoxyribonucleotide (4,5,5=14)

3. (a) Give the mode of action of the following inhibitors and their use in medicine :

(i) Azaserine

(ii) Sulphanilamide

(iii) Trimethoprim

(iv) Allopurinol

(v) Hydroxyurea (2×5=10)

(b) Show how different organisms help to recycle nitrogen in the atmosphere. (4)

4. (a) Differentiate between the following pairs :

(i) Positive and negative nitrogen balance

(ii) Carbamoyl phosphate synthetase I and II

(iii) Transamination and oxidative deamination (4×3=12)

(b) Which derivative of folate is a reactant in the conversion of :

(i) Glycine into serine

(ii) Homocysteine into methionine (1×2=2)

5. Give the biochemical basis of any four of the given metabolic disorders and name the defective enzyme :

(i) Lesch Nyhan syndrome

(ii) Orotic aciduria

(iii) Phenylketonuria

(iv) Maple syrup urine disease

(v) SCID

(3.5×4=14)

6. Write down the steps to accomplish the given conversions (**any 7**)

(i) Tyrosine to melanin

(ii) Histidine to N-formimino glutamate

(iii) Guanine to uric acid

(iv) dUMP to dTTP

(v) Arginine to creatine

(vi) Tryptophan to nicotinamide adenine dinucleotide

(vii) Ornithine to spermine

(viii) Methionine to homocysteine

(2×7=14)

7. (a) Give one significant contribution of the following scientists :

(i) P. Reicherd

(ii) John Buchanan

(iii) JoAnne Stubbe

(iv) A. Folling

(1×4=4)

*P.T.O.*

- (b) (i) What are the different pathways for breakdown and synthesis of glycine ?
- (ii) What are the steps involved in the degradation of heme ?
- (iii) Since dUTP is not a normal component of DNA, why do you suppose ribonucleotide reductase has the capacity to convert UDP to dUDP ?  
(4,3.3=10)
8. (a) Consider the regulation of *E. coli* glutamine synthetase (GS) and explain the metabolic rationale for each of the following effects :
- (i) Inhibition of GS by carbamoyl phosphate
- (ii) Inhibition of the deuridylylation of PII-UMP by  $\alpha$  keto glutarate
- (iii) Activation of uridylylation of PII by ATP (2×3=6)
- (b) Write the role of pyridoxal phosphate in amino acid metabolism. (4)
- (c) Arginine and proline catabolic pathways converge at the same molecule. Write down the steps and enzymes involved in complete degradation of this molecule. (4)