

part of.

6. DEAMINATION essay & mark

The removal of amino group from amino acids as NH_3 .

- oxidative deamination.
- Non-oxidative deamination.

oxidative deamination.

An amino acid is converted into the corresponding keto acid by the removal of the amino functional group as ammonia.

and.

The amino functional group is replaced by the ketone group.

• The ammonia eventually goes into urea cycle.

- Oxidative deamination occurs primarily on glutamic acid

Because,

glutamic acid is the end product of many transamination reactions.

- ^(mark) Only liver mitochondria contains glutamate dehydrogenase (GDH).

- all amino acids are first transaminated to glutamate and then deaminated.

- Amino acids are deaminated at the rate of 50 - 70 g/day.

Minor pathway of deamination:

(a) L-amino acid oxidase. (FMN is the co-enzyme)

- It acts on all amino acids except hydroxy and dicarboxylic amino acids.

(b) D-amino acid oxidase. (FAD is the coenzyme)

- It can oxidise glycine & any D-amino acids that forms from bacterial metabolism.

• Action of MAO (mono amino oxidase)

Non-oxidative Deamination.

(A) Dehydratases - acts on hydroxy amino acids.

(a) Ser forms pyruvate.

(b) Thr forms ~~α-amino butyrate~~
α-Ketobutyrate

B) Desulphydrases. - cysteine undergoes decamination and simultaneous ions-sulphurization to form pyruvate.

- Histidine converted to uracanic acid by histidase.

- Ammonia is formed in GIT by bacterial putrefaction.

Transdeamination

- The amino gp of most of the aa is released by a coupled reaction, transdeamination

- transamination followed by oxidative deamination.

- Transamination takes place in cytoplasm of all cells, ammonia is transported to liver as glutamate.

• Glu is finally oxidatively deaminated in liver mitochondria.

• Any amino acid $\xrightarrow{\alpha\text{-Ketoglutarate}}$ Transamination in all tissues \rightarrow Glutamate.

~~Too~~