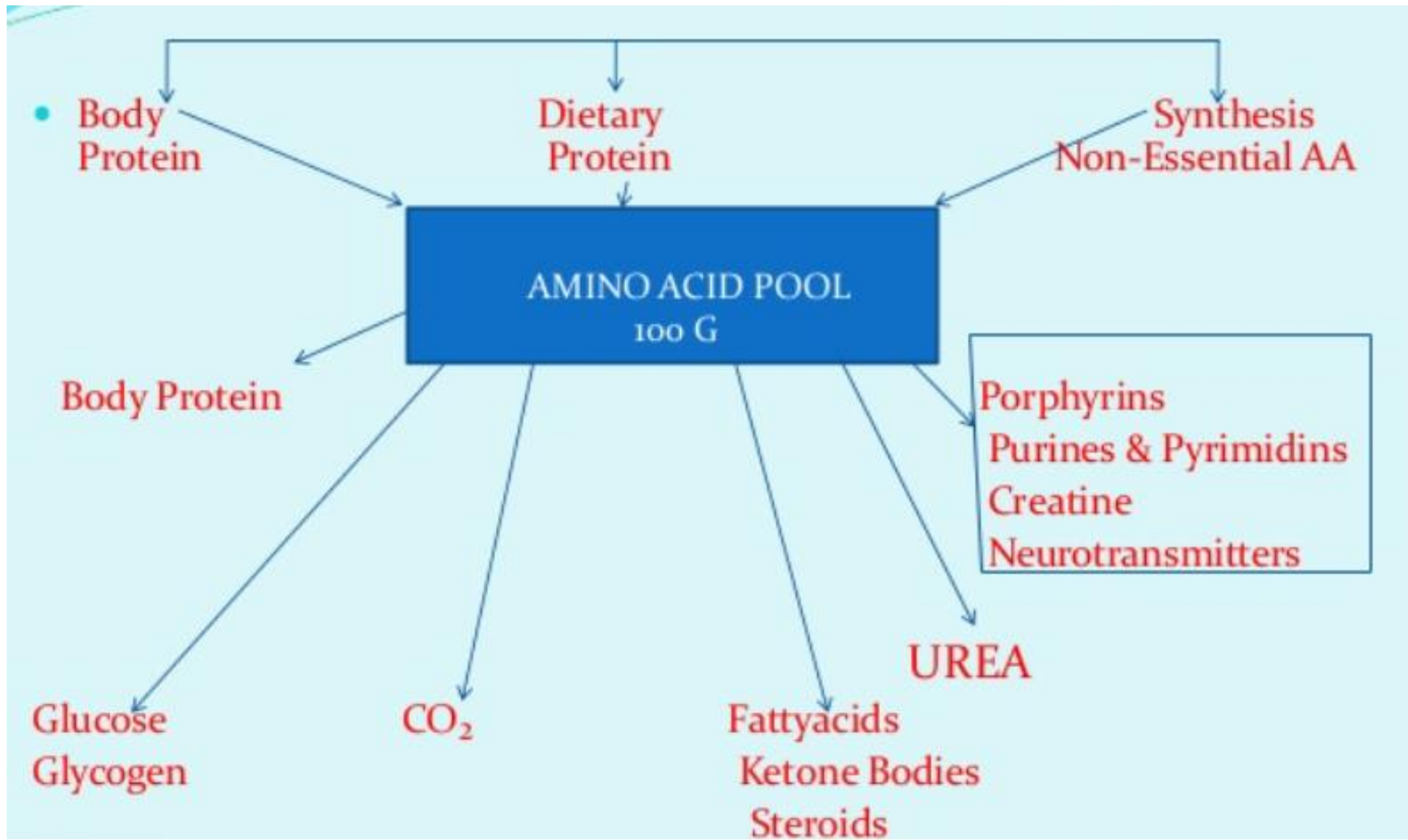


Metabolism of proteins



Deamination

- Removal of amino group in the form of ammonia from amino acid

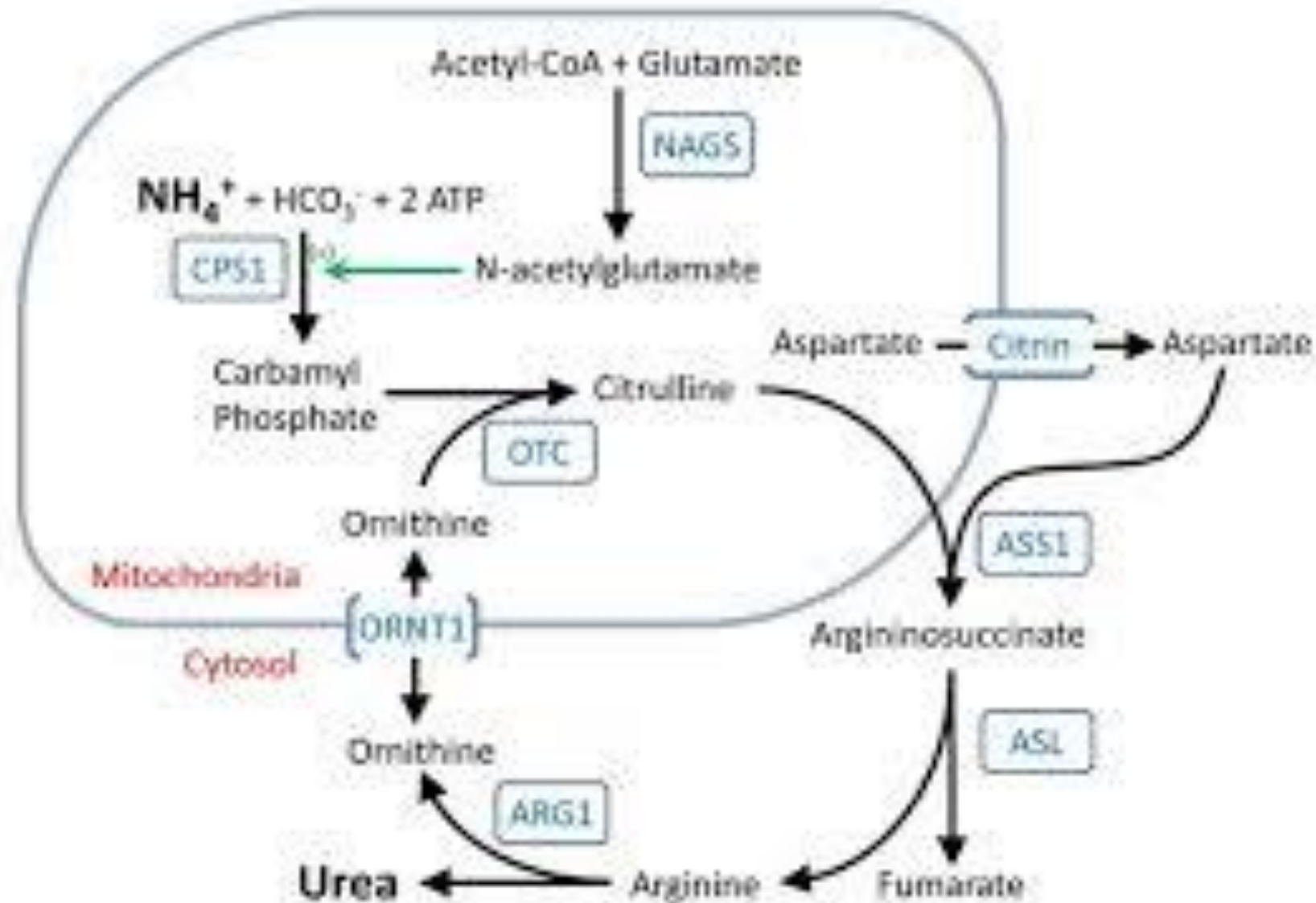
Metabolism of Ammonia:-

- SOURCES OF AMMONIA :-
 - Amino acids Synthesises Protein,
 - Protein degraded to Amino acids.
 - From Liver : a) Transamination
b) Oxidative deamination
 - From Kidney : Glutaminase reaction
 - From Intestine : By Bacterial action
 - From Diet : Amines
 - From Catabolism : Purines (Adenine)
Pyrimidines (Cytosine)
 - From Non-oxidative: Deamination : Aminoacids

UTILIZATION OF AMMONIA:-

- Glutamate+ Ammonia \longrightarrow Glutamine.
- **Glutamine synthetase**- Liver, Brain and Kidney.
- Brain :- Major mechanism for removal of Ammonia is Glutamate formation.
- $\alpha\text{KG} + \text{NH}_3 + \text{NADPH} + \text{H}^+ \longrightarrow \text{Glutamate} + \text{NADP}^+$
- Glutamate may be considered as a major transport form of NH_3 from tissue to liver. Concentration of Glutamate in blood is 10 times more than other amino acids.

Removal of ammonia = Urea cycle



Two nitrogen atoms of urea are derived from ammonia and alpha amino group of aspartic acid.

One mol of *urea* synthesis requires 4 mol of ATP.

Step 1 in Mitochondria:-



Carbamoyl phosphate synthase 1 (CPS-1)

It is a Mitochondrial enzyme,

Allosteric activator is N-Acetyl Glutamate.

- Step 2. Formation of Citrulline

Carbamoyl phosphate + Ornithine


Ornithine Transcarbamoylase

Citrulline + Pi

- Ornithine trans carbamoylase is also a Mitochondrial enzyme

This step onwards the reactions occurred in
CYTOPLASM

- Step3- Formation of Argininosuccinate.
- Citrulline + Aspartate + ATP

 **Argininosuccinate Synthase**

Argininosuccinate + AMP + PPi

- **Step 4. Formation of Arginine.**

Argininosuccinate



Argininosuccinase

Arginine + Fumarate

- **Step 5. Formation of Urea.**

Arginine + H_2O Arginase Ornithine + Urea.