

Metabolism of phenylalanine and tyrosine

We classify Phenylalanine and tyrosine among '*aromatic amino acids*'. The metabolism of phenylalanine and tyrosine are linked directly to each other. The final products of their catabolism are fumarate and acetoacetate.

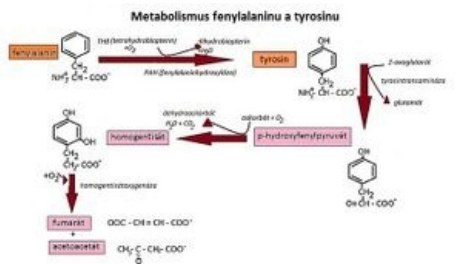
Reaction

- *Phenylalanine* is converted in a reaction catalyzed by PAH (*phenylalanine hydroxylase*), with the help of O₂ and THB (*tetrahydrobiopterin*) into **tyrosine** , releasing H₂O and DHB (*dihydrobiopterin*).
- *Tyrosine* is converted by *tyrosine transaminase* to p-hydroxyphenylpyruvate. 2-oxoglutarate enters the reaction and glutamate is released from the reaction.
- **p-hydroxyphenylpyruvate** in the presence of oxygen and with ascorbate gives rise to homogentisate. The reaction releases H₂O, CO₂ and the by-product dehydroascorbate.
- **Homogentisate** is cleaved under the influence of O₂ in a reaction catalyzed by *homogentisate oxygenase* and subsequently, in several steps, **fumarate and acetoacetate are formed**.

Links

Related Articles

- Disorders of aromatic and branched-chain amino acid metabolism
- Metabolism of AMK group pyruvate and oxaloacetate
- Ketones
- Amino acids



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Source

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