

## Short Communication

## Open Access

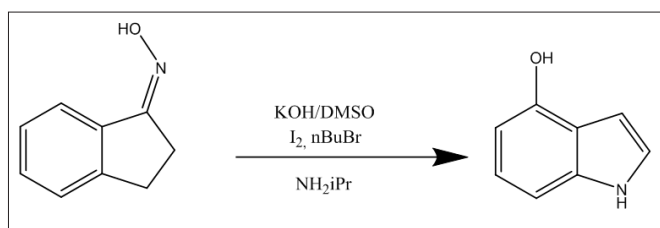
## High Throughput Synthesis of 4-Hydroxy Indole and its Derivatives

Mantas Jonušis

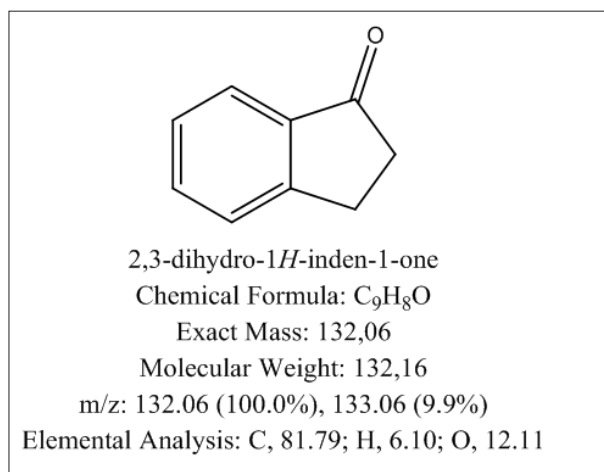
Head, Laboratory of Bioorganic Compounds Chemistry, VU LSC Institute of Biochemistry, Vilnius, Lithuania

**\*Corresponding author**

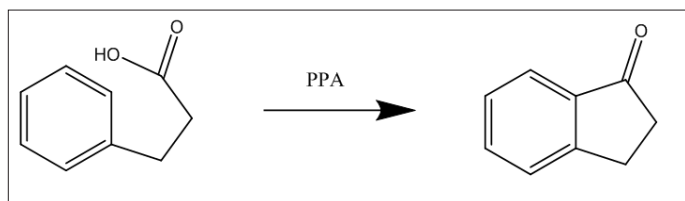
Mantas Jonušis, Head, Laboratory of Bioorganic Compounds Chemistry, VU LSC Institute of Biochemistry, Vilnius, Lithuania.

**Received:** March 07, 2024; **Accepted:** March 08, 2024; **Published:** March 22, 2024

Recently we have been working on 4-OH indole synthesis because it is very interesting heterocycle. For example it is used for the synthesis of indopan – drug for heart problems or anticancer agents. It is also very versatile building block in organic synthesis of other natural and artificial compounds.

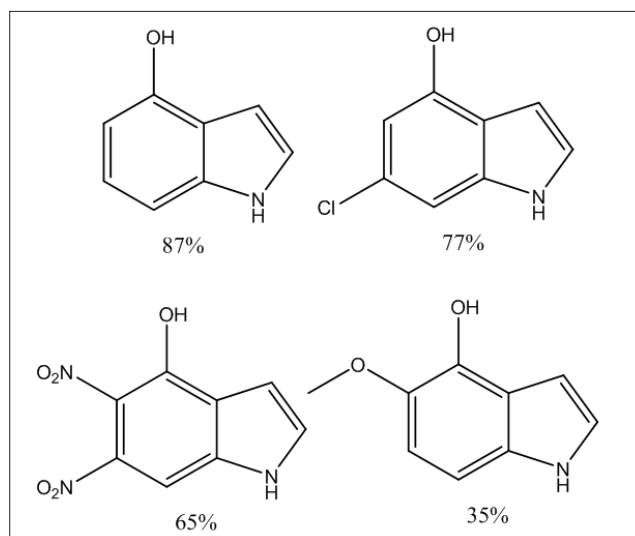


Synthesis starts from indenone or substituted indenones. These starting materials are available at Sigma-Aldrich or can be easily synthesized from phenylpropionic acids and polyphosphoric acid by the further reaction.



Our invention of 4-OH-indole synthesis from indenones is previously unpublished approach. We have studied reaction

conditions on broad scope of substrates and it seems to work on activated and deactivated indenones.



These findings suggest that it might be the invention of new named reaction. Some more research should be done on mechanism studies.

**Copyright:** ©2024 Mantas Jonušis. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.