Technology bulletin

CSIR-Indian Institute of Chemical Biology



Improved synthetic process for the synthesis of 2-(4-chlorophenyl)-2-hydroxy-N-(3-methoxy-4-(prop-2-yn-1-yloxy)phenethyl)acetamide, a key intermediate for the synthesis of Mandipropamid.

INTRODUCTION: Name of Agrochemical: Mandipropamid Category: Fungicide

Mandipropamid is the class of <u>mandelic acid</u> amides and a <u>fungicide</u> from the family of CAA fungicides (carboxamides). Mandipropamid is an inhibitor of cellulose synthase and is effective against downy mildew and Phytophthora infestans. Mandipropamid is a very effective active ingredient: the EC 80 against Phythophthora infestans is 0.1 mg l⁻¹, that against <u>Plasmopara viticola</u> is 1.2 mg l⁻¹

CHALLENGE/APPLICATION DOMAIN:

- ✓ None of the toxic Cyanide sources like HCN, KCN and the expensive Pd or Bromine sources were employed in this synthetic process.
- ✓ The amide coupling was achieved using a relatively easily accessible coupling reagent at room temperature to deliver an intermediate for mandipropamid.
- ✓ In our quest to enhance the efficiency and practicality of this reaction, we've uncovered a valuable process; this newly devised process offers a straightforward and achievable alternative, boasting advantages such as low cost, high yield, and suitability for large-scale industrial production.

SCIENTIFIC MERIT: The process provides a mild, short (Four-step), hazardous, reagent-free strategy for synthesizing the key intermediate from Vanillin for Mandipropamid.

OPPORTUNITY: Competing technology is not available at present, and Pharma and the fine chemical industry can employ this cost-effective IICB- process for tilorone, an antiviral drug.

STAGE OF TECHNOLOGY DEVELOPMENT: TRL 4: Ready for transfer

REFERENCES/ PATENTS:

Patent Number: IN 202311081168 date 29 Nov 2023

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FUNDING: CSIR (HCP-049)

MARKET SIZE: Total import value in India: \$ 9,104,609

Collaborating Institute/Company if any (pls indicate if a separate MoU/agreement is in place.): None.