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## Biological evaluation of New imidazole derivatives tethered with indole moiety as Potent $\alpha$ -Glucosidase Inhibitors

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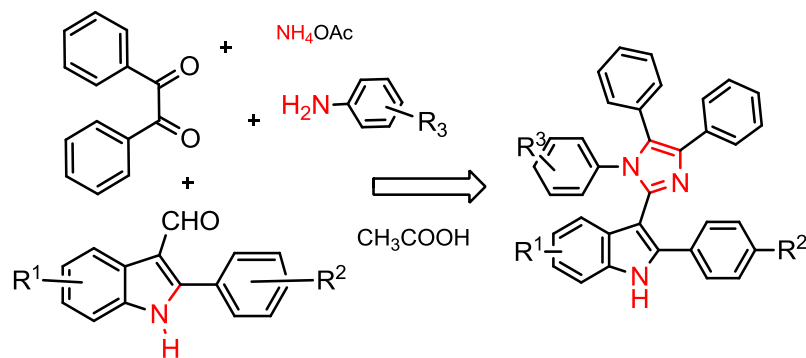
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### Abstract

A series of triarylimidazoles substituted with 2-arylindoles (**4a-4j**) were prepared and evaluated for their *in vitro*  $\alpha$ -Glucosidase inhibition.  $\alpha$ -Glucosidase inhibition assay displayed a new class of highly potent agents. The new compounds showed significant  $\alpha$ -glucosidase inhibitory activity as compared to the standard inhibitor acarbose. Structures of synthesized compounds were determined by using Mass spectrometry FT-IR, <sup>1</sup>H NMR and <sup>13</sup>C NMR.

### Graphical Abstract:

$\alpha$ -Glucosidase inhibition by triarylimidazoles substituted with 2-arylindoles



R<sup>1</sup> = H, Cl, F

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