

Review of: "Synthesis of 1, 2-Disubstituted Benzimidazoles at Ambient Temperature Catalyzed by 1-Methylimidazolium Tetraflouroborate ([Hmim] BF\_4) and Investigating Their Anti-ovarian Cancer Properties Through Molecular Docking Studies and Calculations"

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Potential competing interests: No potential competing interests to declare.

The manuscript developed a new method for the synthesis of 1,2-disubstituted benzimidazoles by the reaction of aromatic aldehydes and o-phenylenediamines (OPD) in the presence of 1-methylimidazolium tetrafluoroborate ([Hmim] BF4) at ambient temperature under green conditions. There are some major revisions before publication:

- 1. In the experimental section, mass spectrum or elemental analysis data needs to be mentioned.
- 2. In the Results and Discussion section, in Table 1: Have you ever tried another ratio between water and EtOH? Have you explained that the reaction time increased when the temperature accelerated?
- 3. The docking study is well performed. However, the antiproliferative assay needs to be performed to confirm the results.
- 4. English checking is necessary before online publication.