

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.

In this study, the authors report a synthesis of 1,2-disubstituted benzimidazoles catalysed by 1-methylimidazolium tetrafluoroborate ([Hmim] BF4) and investigate their anti-ovarian cancer properties through molecular docking studies and calculations.

In my opinion, the manuscript needs revision, and my comments on the submission are as follows:

- 1. Table 3, Table 4, and Table 5 have the same title; they could be combined into one.
- 2. Figure 4 should be discussed in the text.
- 3. Was enough investigation done to conclude that all of the synthesized compounds possess beneficial effects for ovarian cancer treatment?
- 4. There are some small errors, e.g., orthophenylenediamine in some cases is marked as 2, and in other cases as 2a; this should be unified. In Table 1, the R, representing substituents, is unnecessary, etc.

Qeios ID: JX8JIP · https://doi.org/10.32388/JX8JIP