World Academy of Science, Engineering and Technology
International Journal of Chemical and Molecular Engineering
Vol:17, No:07, 2023

Enzyme Inhibition Activity of Schiff Bases Against Mycobacterium Tuberculosis Using Molecular Docking

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Abstract : The main cause of infectious disease in the modern world is Mycobacterium Tuberculosis (MT). To combat tuberculosis, new and efficient drugs are an urgent need in the modern world. Schif bases are potent for their biological pharmacophore activity. Thus we selected different Vanillin-based Schiff bases for their binding activity against target enzymes of Mycobacterium tuberculosis that is (DprE1 (decaprenyl phosphoryl-β-D-ribose 2′-epimerase), and DNA gyrase subunit-A), using molecular docking. We evaluate the inhibition potential, interaction, and binding mode of these compounds with the target enzymes.

Keywords : schiff bases, tuberculosis, DNA gyrase, DprE1, docking
Conference Title : ICFAC 2023 : International Conference on Flow and Analytical Chemistry
Conference Location : Toronto, Canada
Conference Dates : July 10-11, 2023

Open Science Index, Chemical and Molecular Engineering Vol:17, No:07, 2023 publications.waset.org/abstracts/168664.pdf