

## Library Indian Institute of Science Education and Research Mohali



## DSpace@IISERMohali (/jspui/)

- / Publications of IISER Mohali (/jspui/handle/123456789/4)
- / Research Articles (/jspui/handle/123456789/9)

Please use this identifier to cite or link to this item: http://hdl.handle.net/123456789/5153 Title: Synthesis, in vitro anti-plasmodial potency, in-silico-cum-SPR binding with inhibition of PfPyridoxal synthase and rapid parasiticidal action by 3,5-bis{(E) arylidene}-N-methyl-4-piperidones Authors: Joshi, Mayank (/jspui/browse?type=author&value=Joshi%2C+Mayank) Choudhury, Angshuman Roy (/jspui/browse? type=author&value=Choudhury%2C+Angshuman+Roy) 3,5-bis{(E) arylidene}-N-methyl-4-piperidones Keywords: anti-plasmodial potency Issue Date: 2021 Publisher: RCS Citation: New Journal of Chemistry, 45(47), 22150-22165. Abstract: Twenty-five (Ia-Iu, IIa-IIb, IIIa, and IVa) diarylidene-N-methyl-4-piperidones (DANMPs) were synthesized and characterized via UV, FT-IR, NMR, and MS while Id was characterized also by single crystal XRD. Twenty-one compounds shortlisted after initial in vitro anti-plasmodial activity successive screenings at 100 µM and 10 µM were evaluated for their IC50s against chloroquinesensitive Pf3D7, chloroquine-resistant PfINDO, and artemisinin-resistant PfMRA-1240 strains. The four most promising compounds were le (IC50s  $\mu$ M 0.35MRA, 1.39INDO, 1.923D7), If (IC50s  $\mu$ M 1.07MRA, 1.36INDO, 3.393D7), Ir (IC50s  $\mu$ M 0.74MRA, 2.45INDO, 1.443D7), and In (IC50s  $\mu$ M 1.27MRA, 1.8INDO, 1.73D7), Resistance indices as low as 0.2 to 0.5 for these potent compounds and <1 for most other compounds suggest their greater potency against drug resistant strains than the drug sensitive strain. The parasiticidal action of Ir was seen within 4 h against the trophozoite stage of the parasite, which is known to express the highest levels of PLP synthase. In silico docking scores of -7.0 to -8.0 kcal mol-1 between potent DANMPs and PfPLP synthase, the direct binding of Ir studied by SPR to recombinantly expressed and purified PfPdx-1 and inhibition of Pdx1 enzymatic activity by Ir suggest this vital enzyme to be a probable target for the DANMPs. The non-hemolytic nature of Ir and conformity of most DANMPs to Lipinski's parameters indicate their potential as new anti-plasmodial leads with PfPLP synthase as one of their targets. Only IISER Mohali authors are available in the record. Description: URI: https://pubs.rsc.org/en/content/articlelanding/2021/NJ/D1NJ04604G (https://pubs.rsc.org/en/content/articlelanding/2021/NJ/D1NJ04604G) http://hdl.handle.net/123456789/5153 (http://hdl.handle.net/123456789/5153) Appears in Research Articles (/jspui/handle/123456789/9) Collections:

File	Description	Size	Format	
Need To AddFull Text_PDF (/ispui/bitstream/123456789/5153/1/Need%20To%20Add%e2%80%a6Full%20Text_PDF)		15.36 kB	Unknown	View/Open (/jspui/t

Show full item record (/jspui/handle/123456789/5153?mode=full)

■ (/jspui/handle/123456789/5153/statistics)

Items in DSpace are protected by copyright, with all rights reserved, unless otherwise indicated.