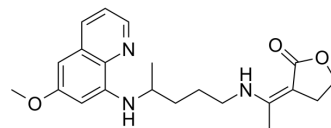


## Bulaquine

Cat. No.:	HY-106866
CAS No.:	79781-00-3
Molecular Formula:	C <sub>21</sub> H <sub>27</sub> N <sub>3</sub> O <sub>3</sub>
Molecular Weight:	369.46
Target:	Parasite
Pathway:	Anti-infection
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Bulaquine (CDRI 80/53) is a potent antimalarial agent which is an analogue of <a href="#">Primaquine</a> (HY-12651A). Bulaquine affects multiple metabolism pathways and shows inhibition effect on <i>Plasmodium cynomolgi</i> infection. Bulaquine can be used for the research of malaria <sup>[1][2][3]</sup> .								
<b>IC<sub>50</sub> &amp; Target</b>	Plasmodium								
<b>In Vivo</b>	<p>Bulaquine (40 mg/kg; p.o. once) affects multiple pathways in vivo<sup>[1]</sup>.</p> <p>Bulaquine (1.25 mg/kg; once daily; for 7 days) shows 100% curative anti-relapse activity with a primaquine index of 0.8 for rhesus monkeys with <i>Plasmodium cynomolgi</i> B infection<sup>[2]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>25-30 g male Swiss albino mice<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>40 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Oral gavage; 40 mg/kg once</td> </tr> <tr> <td>Result:</td> <td>Affected oxidative stress and fatty acid synthesis pathway, apoptosis, cell cycle, inflammatory response, glycogen metabolism, Krebs's cycle, electron transport chain, fatty acid β-oxidation, MAPK signalling, signalling of hepatocyte growth factor receptor, matrix metalloproteinases, steroid biosynthesis, TGF-β signalling, translation factors, Wnt signalling, regulation of actin cytoskeleton, ribosomal proteins, RNA transcription reactome, proteasome degradation and nuclear receptors in lipid metabolism according to GO results.</td> </tr> </table>	Animal Model:	25-30 g male Swiss albino mice <sup>[1]</sup>	Dosage:	40 mg/kg	Administration:	Oral gavage; 40 mg/kg once	Result:	Affected oxidative stress and fatty acid synthesis pathway, apoptosis, cell cycle, inflammatory response, glycogen metabolism, Krebs's cycle, electron transport chain, fatty acid β-oxidation, MAPK signalling, signalling of hepatocyte growth factor receptor, matrix metalloproteinases, steroid biosynthesis, TGF-β signalling, translation factors, Wnt signalling, regulation of actin cytoskeleton, ribosomal proteins, RNA transcription reactome, proteasome degradation and nuclear receptors in lipid metabolism according to GO results.
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### REFERENCES

[1]. Noel S, et al. Identification of differentially expressed genes after acute exposure to bulaquine (CDRI 80/53) in mice liver. *Basic Clin Pharmacol Toxicol*. 2008 Dec;103(6):522-9.

[2]. Dutta GP, et al. Radical curative activity of a new 8-aminoquinoline derivative (CDRI 80/53) against *Plasmodium cynomolgi* B in monkeys. *Am J Trop Med Hyg*. 1989 Dec;41(6):635-7.

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[3]. Wells TN, et al. Targeting the hypnozoite reservoir of Plasmodium vivax: the hidden obstacle to malaria elimination. Trends Parasitol. 2010 Mar;26(3):145-51. Wells TN, et al. Targeting the hypnozoite reservoir of Plasmodium vivax: the hidden obstacle to malaria elimination. Trends Parasitol. 2010 Mar;26(3):145-51.

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**Caution: Product has not been fully validated for medical applications. For research use only.**

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