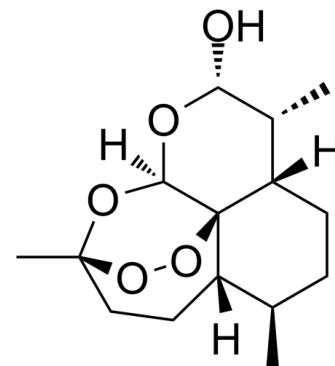


Data Sheet

Product Name:	Dihydroartemisinin
Cat. No.:	CS-5595
CAS No.:	71939-50-9
Molecular Formula:	C ₁₅ H ₂₄ O ₅
Molecular Weight:	284.35
Target:	Apoptosis; Autophagy; NF-κB; Parasite
Pathway:	Anti-infection; Apoptosis; Autophagy; NF-κB
Solubility:	DMSO : 41.67 mg/mL (146.54 mM; Need ultrasonic)



BIOLOGICAL ACTIVITY:

Dihydroartemisinin is a potent **anti-malaria** agent. IC₅₀ & Target: RelA/p65^[1]

Autophagy^[1] In Vitro: Dihydroartemisinin (DHA) is an antimalarial agent. Dihydroartemisinin treatment effectively up-regulates the cytosolic RelA/p65 protein level and down-regulates the nuclear RelA/p65 protein level. Dihydroartemisinin blocks the nuclear translocation of RelA/p65 from the cytosol rather than suppressing RelA/p65 protein synthesis. Dihydroartemisinin induces autophagy in RPMI 8226 cells. Dihydroartemisinin suppresses NF-κB activation in RPMI 8226 cells. The NF-κB Dihydroartemisinin -binding activity is examined by EMSA assay. RPMI 8226 cells are exposed to various concentrations of Dihydroartemisinin (10, 20 and 40 μM) for 12 h, and TNF-α is introduced as a positive control for NF-κB activation. Dihydroartemisinin suppresses NF-κB activation in a dose-dependent manner in contrast with TNF-α^[1]. Dihydroartemisinin (DHA) can enhance the anti-tumor effect of photodynamic therapy (PDT) on esophageal cancer cells, and cell viability is investigated using the MTT assay. Eca109 and Ec9706 cells are treated with Dihydroartemisinin (80 μM), PDT (25 and 20 J/cm², respectively) or their combination. Single treatment with Dihydroartemisinin or PDT causes a 37±5% or 34±6% reduction in viability in Eca109 cells and a 33±7% or 34±6% reduction in Ec9706 cells, respectively. However, when PDT is combined with Dihydroartemisinin, the cell viability is reduced 59±6% or 61±7% in the cell lines, respectively^[2]. **In Vivo:** Single oral doses of Dihydroartemisinin (at 200, 300, 400 or 600 mg/kg), given once on each of day 6-8 post-infection, reduce total-worm burdens by 69.2%-90.6% and female-worm burdens by 62.2%-92.2%, depending on dosage in the first experiment. Similar treatments given on day 34-36 post-infection reduce total-worm burdens by 73.9%-85.5% and female-worm burdens by 83.8%-95.3%^[3].

PROTOCOL (Extracted from published papers and Only for reference)

Kinase Assay: ^[1]To determine NF-κB Dihydroartemisinin-binding activity, an electrophoretic mobility shift assay (EMSA) is performed. Nuclear extracts are prepared and incubated with ³²P-end-labeled 45-mer double-stranded oligonucleotide (15 μg protein with 16 fmol DNA) from the HIV long terminal repeat, 5'-TTGTTACAAGGGACTTCCGCTG GGGACTTCCAGGGAGGCGTGG-3' (boldface indicates NF-κB binding sites), for 30 min at 37 °C. The Dihydroartemisinin-protein complex formed is separated from free oligonucleotide on 6.6% native polyacrylamide gels. A double-stranded mutated oligonucleotide, 5'-TTGTTACAA CTCACTTCCGCTGCTCACTTCCAGGGAGGCGTGG-3', is used to examine binding specificity of NF-κB to the DNA. The binding specificity is also examined by competition with the unlabeled oligonucleotide. Preimmune serum (PIS) is included as a negative control. The dried gels are visualized with a Storm 820, and radioactive bands are quantified using Imagequant software^[1].

Cell Assay: ^[2]**Eca109 (4×10³ cells/well) and Ec9706 (5×10³ cells/well) cells** are grown in 96-well plates and cultured overnight to allow for cell attachment. Eca109 and Ec9706 cells are treated with **Dihydroartemisinin (80 μM)**, PDT (25 and 20 J/cm², respectively) or their combination. After incubation for 24h, MTT (20 μL) is added to each well and incubated for 4 h at 37°C. Formazan crystals are dissolved in 150 μL of DMSO for 10 min with shaking. The absorbance is measured at 490 nm on a plate reader, and the experiment is repeated three times^[2].

Animal Administration: Dihydroartemisinin is prepared in DMSO, Tween-80 and distilled water to give aqueous solutions^[3]. Mice^[3] of the Kunming strain, each weighing 20-24 g, are used. In the first experiment, design to investigate the effect of multiple doses of Dihydroartemisinin on the schistosomula and adult worms of *S. japonicum*, mice are given **three daily doses, of 200, 300, 400 or 600 mg Dihydroartemisinin/kg (in dose volumes of 25 mL/kg)**, on days 6-8 or 34-36 post-infection, respectively. An additional group of mice, infected but not given the drug, serve as a control.

References:

- [1]. Hu W, et al. Dihydroartemisinin induces autophagy by suppressing NF-κB activation. *Cancer Lett.* 2014 Feb 28;343(2):239-48.
- [2]. Li YJ, et al. Dihydroartemisinin accentuates the anti-tumor effects of photodynamic therapy via inactivation of NF-κB in Eca109 and Ec9706 esophageal cancer cells. *Cell Physiol Biochem.* 2014;33(5):1527-36.
- [3]. Li HJ, et al. Dihydroartemisinin-praziquantel combinations and multiple doses of dihydroartemisinin in the treatment of *Schistosoma japonicum* in experimentally infected mice. *Ann Trop Med Parasitol.* 2011 Jun;105(4):329-33.

CAIndexNames:

3,12-Epoxy-12H-pyrano[4,3-j]-1,2-benzodioxepin-10-ol, decahydro-3,6,9-trimethyl-, (3R,5aS,6R,8aS,9R,10S,12R,12aR)-

SMILES:

O[C@@H]1[C@H](C)[C@]2([H])CC[C@@H](C)[C@]3([H])CC[C@@](O4)(C)OO[C@]32[C@]4([H])O1

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

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