Product Data Sheet



Cat. No.: HY-103222 14756-24-2 CAS No.: Molecular Formula: C21H16O4 Molecular Weight: 332.35

Target: Aryl Hydrocarbon Receptor Pathway: Immunology/Inflammation Storage: Powder -20°C 3 years

4°C 2 years

-80°C 6 months In solvent

> -20°C 1 month

SOLVENT & SOLUBILITY

In Vitro

DMSO: 12.5 mg/mL (37.61 mM; ultrasonic and warming and heat to 70°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.0089 mL	15.0444 mL	30.0888 mL
	5 mM	0.6018 mL	3.0089 mL	6.0178 mL
	10 mM	0.3009 mL	1.5044 mL	3.0089 mL

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description DiMNF (3',4'-Dimethoxy-αNF) is a selective aryl hydrocarbon receptor (AHR) modulator. DiMNF is a competitive AHR ligand (

 $IC_{50} = 21 \text{ nM}$) with apparent antagonistic activity. DiMNF can be used as an anti-inflammatory agent^[1].

In Vitro DiMNF (3',4'-Dimethoxy- α NF) (10 μ M; 4 h) minimally influences AHR gene battery^[1].

DiMNF (10 μM; 1 h) effectively suppresses cytokine-mediated inflammatory gene expression^[1].

DiMNF adopts a unique orientation within the ligand binding pocket of $AHR^{[1]}$.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

RT-PCR^[1]

Cell Line:	Huh7 cell	
Concentration:	10 μΜ	
Incubation Time:	4 h or 1 h	
Result:	Decreased CYP1A1 mRNA expression (4 h). Exhibited repressive activity toward cytokine-	

mediated SAA1 induction (1 h). Significantly reduced IL1 β -mediated mRNA induction of C4 , C1S, C1R, and C3.

REFERENCES

[1]. Murray IA, et al. Suppression of cytokine-mediated complement factor gene expression through selective activation of the Ah receptor with 3',4'-dimethoxy- α -naphthoflavone. Mol Pharmacol. 2011 Mar;79(3):508-19.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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