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产品名称: **2'-C-甲基胞嘧啶核苷**
产品别名: **NM107 ; 2'-C-Methylcytidine; NM-107**

生物活性:																				
Description	NM107 (2'-C-Methylcytidine) is an nucleoside inhibitor of the hepatitis C virus (HCV) NS5B polymerase, the EC50 of NM107 in the wild-type replicon cells is 1.85 μM[1][2].																			
IC ₅₀ & Target	HCV[1]																			
In Vitro	NM107 (25 μM; 24 hours; Huh7-1 cells and cell culture-propagated HCV) treatment decreases extracellular viral titers and intracellular RNA levels. RT-PCR[2] <table border="1"><tr><td>Cell Line:</td><td>Huh7-1 cells and cell culture-propagated HCV (HCVcc)</td></tr><tr><td>Concentration:</td><td>25 μM</td></tr><tr><td>Incubation Time:</td><td>24 hours</td></tr><tr><td>Result:</td><td>Extracellular virus titers declined in parallel with intracellular RNA levels.</td></tr></table>	Cell Line:	Huh7-1 cells and cell culture-propagated HCV (HCVcc)	Concentration:	25 μM	Incubation Time:	24 hours	Result:	Extracellular virus titers declined in parallel with intracellular RNA levels.											
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Solvent&Solubility	<p>In Vitro:</p> <p>H₂O : ≥ 50 mg/mL (194.37 mM)</p> <p>* "≥" means soluble, but saturation unknown.</p> <table border="1"><thead><tr><th rowspan="2">Preparing Stock Solutions</th><th>Solvent</th><th>Mass</th><th>Concentration</th><th>1 mg</th><th>5 mg</th><th>10 mg</th></tr></thead><tbody><tr><td>1 mM</td><td>3.8874 mL</td><td>19.4371 mL</td><td>38.8742 mL</td></tr><tr><td>5 mM</td><td>0.7775 mL</td><td>3.8874 mL</td><td>7.7748 mL</td></tr><tr><td>10 mM</td><td>0.3887 mL</td><td>1.9437 mL</td><td>3.8874 mL</td></tr></tbody></table> <p>*请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液 一旦配成溶液, 请分装保存, 避免反复冻融造成的产品失效。</p> <p>储备液的保存方式和期限 -80°C, 6 months; -20°C, 1 month。 -80°C 储存时, 请在 6 个月内使用, -20°C 储存时, 请在 1 个月内使用。</p>	Preparing Stock Solutions	Solvent	Mass	Concentration	1 mg	5 mg	10 mg	1 mM	3.8874 mL	19.4371 mL	38.8742 mL	5 mM	0.7775 mL	3.8874 mL	7.7748 mL	10 mM	0.3887 mL	1.9437 mL	3.8874 mL
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References	[1]. Mathy JE, et al, Combinations of cyclophilin inhibitor NIM811 with hepatitis C Virus NS3-4A Protease or NS5B polymerase inhibitors enhance antiviral activity and suppress the emergence of resistance. Antimicrob Agents Chemother. 2008 Sep;52(9):3267-75. [2]. Guedj J, et al. Modeling shows that the NS5A inhibitor daclatasvir has two modes of action and yields a shorter estimate of the hepatitis C virus half-life. Proc Natl Acad Sci U S A. 2013 Mar 5;110(10):3991-6.																			