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产品名称: **VAL-083**

产品别名: 去水卫矛醇; **Dianhydrodulcitol; Dianhydrogalactitol**

生物活性:																				
Description	VAL-083 is an alkylating agent that creates N7 methylation on DNA, with antitumor activity.																			
IC₅₀ & Target	DNA Alkylator[1]																			
In Vitro	VAL-083 is an alkylating agent that creates N7 methylation on DNA. VAL-083 suppresses U251 and SF188 cell growth and induces apoptosis after 72 h. VAL-083 (5 μM) inhibits the growth of SF188 by ~95%. VAL-083 inhibits T98G cells growth in a dose-dependent manner (IC ₅₀ <5 μM)[1]. VAL-083 (Dianhydrogalactitol) inhibits the proliferation of HUVEC and U251 cells at doses of more than 12.5 μg/mL. VAL-083 (3.125, 6.25, 12.5 μg/mL) also suppresses the migration and invasion, and reduces MMP2, VEGF, VEGFR2, and FGF2 expression in HUVEC and U251 cells[2]. VAL-083 (1,2:5,6-dianhydrogalactitol, 1, 2, 5 μM) dose-dependently induces cell cycle arrest at G2/M phase in the 3 glioma cell lines. VAL-083 activates two parallel signaling cascades, the p53-p21 and the CDC25C-CDK1 cascade. In addition, VAL-083 significantly enhances the radiosensitivity of LN229 cells[3].																			
In Vivo	VAL-083 (Dianhydrogalactitol; 25, 50, 100 μg/mL) dose-dependently inhibits angiogenesis in zebrafish model. VAL-083 considerably reduces VEGF, VEGFR2, and FGF2 expression at 25 μg/mL, and further causes reduction in FGFR2 expression at 50 μg/mL[2]. VAL-083 (1,2:5,6-dianhydrogalactitol; 5 mg/kg, iv, twice per week for 6 weeks) significantly blocks the growth of LN229 cells in mice with the relative tumor growth rate (T/C) of 22.38%, and the tumor growth inhibitory rate (TGI) of 83.58%. Moreover, VAL-083 dramatically activates the CDC25C-CDK1 cascade in the xenografted tumor model[3].																			
Solvent&Solubility	<p>In Vitro:</p> <p>DMF : ≥ 100 mg/mL (684.28 mM)</p> <p>H₂O : 50 mg/mL (342.14 mM; Need ultrasonic)</p> <p>* "≥" means soluble, but saturation unknown.</p>																			
	<table border="1"> <thead> <tr> <th rowspan="2">Solvent</th> <th rowspan="2">Mass Concentration</th> <th rowspan="2">1 mg</th> <th rowspan="2">5 mg</th> <th rowspan="2">10 mg</th> </tr> </thead> <tbody> <tr> <td>Preparing</td> <td>1 mM</td> <td>6.8428 mL</td> <td>34.2138 mL</td> <td>68.4275 mL</td> </tr> <tr> <td rowspan="2">Stock Solutions</td> <td>5 mM</td> <td>1.3686 mL</td> <td>6.8428 mL</td> <td>13.6855 mL</td> </tr> <tr> <td>10 mM</td> <td>0.6843 mL</td> <td>3.4214 mL</td> <td>6.8428 mL</td> </tr> </tbody> </table>	Solvent	Mass Concentration	1 mg	5 mg	10 mg	Preparing	1 mM	6.8428 mL	34.2138 mL	68.4275 mL	Stock Solutions	5 mM	1.3686 mL	6.8428 mL	13.6855 mL	10 mM	0.6843 mL	3.4214 mL	6.8428 mL
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<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>																				
References	<p>[1]. Kaiji Hu, et al. Abstract 811: VAL083, a novel N7 alkylating agent, surpasses NSC 362856 activity and inhibits cancer stem cells providing a new potential treatment option for glioblastoma multiforme. Cancer Research. 2012 Mar 31-Apr 4.</p> <p>[2]. Jiang X, et al. Dianhydrogalactitol, a potential multitarget agent, inhibits glioblastoma migration, invasion, and angiogenesis. Biomed Pharmacother. 2017 Jul;91:1065-1074.</p>																			



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	[3]. Peng C, et al. 1,2:5,6-dianhydrogalactitol inhibits human glioma cell growth in vivo and in vitro by arresting the cell cycle at G2/M phase. Acta Pharmacol Sin. 2017 Apr;38(4):561-570.
实验参考:	
Cell Assay	The effects of VAL-083 in HUVEC and U251 cellHUVEC and U251 cell proliferation are measured by the CCK8 assay. Cells are seeded into 96-well plates at a density of 1×10^4 cells per well. After overnight incubation, cell attachment is followed by the addition of VAL-083 in various concentrations for 24 h; then 10 μ L CCK8 is added to each well and incubated at 37°C for 2 h. Optical density is measured at 450 nm[2].
Animal Administration	Mice[3] LN229 cells are suspended in MEM, and 2×10^6 cells per mouse are subcutaneously injected into the flank of BALB/c nude mice at 6-8 weeks old. The tumor volume is calculated as follows: $0.5 \times L \times W^2$. Tumor-bearing mice are divided into two groups (n = 8) with similar average volumes (vehicle: $108 \pm 4 \text{ mm}^3$ vs VAL-083: $107 \pm 4 \text{ mm}^3$). Then, both groups undergo the following treatment: The VAL-083 treatment group receives VAL-083 at 5 mg/kg or 10 μ L/g, iv, twice per week for 6 weeks. The vehicle group receives saline at 10 μ L/g, iv, three times per week for 6 weeks. Tumor volumes are measured twice per week[3].
References	[1]. Kaiji Hu, et al. Abstract 811: VAL083, a novel N7 alkylating agent, surpasses NSC 362856 activity and inhibits cancer stem cells providing a new potential treatment option for glioblastoma multiforme. Cancer Research. 2012 Mar 31-Apr 4. [2]. Jiang X, et al. Dianhydrogalactitol, a potential multitarget agent, inhibits glioblastoma migration, invasion, and angiogenesis. Biomed Pharmacother. 2017 Jul;91:1065-1074. [3]. Peng C, et al. 1,2:5,6-dianhydrogalactitol inhibits human glioma cell growth in vivo and in vitro by arresting the cell cycle at G2/M phase. Acta Pharmacol Sin. 2017 Apr;38(4):561-570.

源叶生物