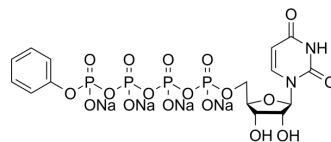


## MRS2768 tetrasodium salt

Cat. No.:	HY-108649A
CAS No.:	2567869-47-8
Molecular Formula:	C <sub>15</sub> H <sub>16</sub> N <sub>2</sub> Na <sub>4</sub> O <sub>18</sub> P <sub>4</sub>
Molecular Weight:	728.14
Target:	P2Y Receptor
Pathway:	GPCR/G Protein
Storage:	-80°C



### BIOLOGICAL ACTIVITY

<b>Description</b>	MRS2768 tetrasodium salt is a moderately potent and selective P2Y2 receptor agonist. MRS2768 tetrasodium salt has a protective effect on cardiomyocytes from ischemic damage in vivo and in vitro <sup>[1][2]</sup> .								
<b>IC<sub>50</sub> &amp; Target</b>	P2Y2 Receptor								
<b>In Vitro</b>	<p>MRS2768 (0.01-10000 μM; 24 hours) significantly increases the proliferation of PANC-1 cells<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only. Cell Proliferation Assay<sup>[1]</sup></p> <table border="1"> <tr> <td>Cell Line:</td> <td>Human pancreatic duct epithelial cells PANC-1</td> </tr> <tr> <td>Concentration:</td> <td>0.01, 0.1, 1, 10, 100, 1000, 10000 μM</td> </tr> <tr> <td>Incubation Time:</td> <td>24 hours</td> </tr> <tr> <td>Result:</td> <td>The effect on proliferation of PANC-1 cells was dependent on concentration (0.1 μM to 1 mM). The concentration that elicited a half-maximal response (EC<sub>50</sub>) in the stimulation of proliferation was 0.8±1.7 μM. Resulted in a dose- and time-dependent increase of proliferation in PANC-1 cells</td> </tr> </table>	Cell Line:	Human pancreatic duct epithelial cells PANC-1	Concentration:	0.01, 0.1, 1, 10, 100, 1000, 10000 μM	Incubation Time:	24 hours	Result:	The effect on proliferation of PANC-1 cells was dependent on concentration (0.1 μM to 1 mM). The concentration that elicited a half-maximal response (EC <sub>50</sub> ) in the stimulation of proliferation was 0.8±1.7 μM. Resulted in a dose- and time-dependent increase of proliferation in PANC-1 cells
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<b>In Vivo</b>	<p>MRS2768 (4.44 μg/kg i.v.) pretreatment reduces myocardial damage in mice. MRS2768 has a protective effect on cardiomyocytes from ischemic damage in vivo<sup>[2]</sup>. 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>Male wild-type mice (C57BL)<sup>[2]</sup></td> </tr> <tr> <td>Dosage:</td> <td>4.44 μg/kg</td> </tr> <tr> <td>Administration:</td> <td>Injected i.v.; 1 h before myocardial infarct (MI)</td> </tr> <tr> <td>Result:</td> <td>Pretreatment reduced myocardial damage. The damage was significantly smaller in mice compare to the untreated mice (25.6±4.5% vs. 39.2±6.3%).</td> </tr> </table>	Animal Model:	Male wild-type mice (C57BL) <sup>[2]</sup>	Dosage:	4.44 μg/kg	Administration:	Injected i.v.; 1 h before myocardial infarct (MI)	Result:	Pretreatment reduced myocardial damage. The damage was significantly smaller in mice compare to the untreated mice (25.6±4.5% vs. 39.2±6.3%).
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## REFERENCES

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[1]. Ji Hun Choi, et al. Uridine Triphosphate Increases Proliferation of Human Cancerous Pancreatic Duct Epithelial Cells by Activating P2Y2 Receptor. *Pancreas*. 2013 May;42(4):680-6.

[2]. Edith Hochhauser, et al. P2Y2 Receptor Agonist With Enhanced Stability Protects the Heart From Ischemic Damage in Vitro and in Vivo. *Purinergic Signal*. 2013 Dec;9(4):633-42.

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

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