



NH

ONa

# **Rigosertib sodium salt**

Cat. No.:	A1404
CAS No.:	1225497-78-8
Formula:	C21H24NNaO8S
M.Wt:	473.47
Synonyms:	ON-01910,Estybon
Target:	Cell Cycle/Checkpoint
Pathway:	PLK
Storage:	Store at -20°C

# Solvent & Solubility

≥23.65mg/mL in DMSO

Preparing In Vitro Stock Solut		Solvent Concentration	1mg	5mg	10mg
	Stock Solutions	1 mM	2.1121 mL	10.5603 mL	21.1207 mL
		5 mM	0.4224 mL	2.1121 mL	4.2241 mL
	10	10 mM	0.2112 mL	1.0560 mL	2.1121 mL

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

## **Biological Activity**

Shortsummary	Plk1 inhibitor	
IC <sub>50</sub> & Target	9 nM (Plk1)	
	Cell Viability Assay	<b>B10</b>
In Vitro	Cell Line:	HeLa cells
	Preparation method:	The solubility of this compound in DMSO is > 10 mM. General tips for obtaining a higher concentration: Please warm the tube at 37 °C for 10 minutes and/or shake it in the ultrasonic bath for a while. Stock solution can be stored below - 20 °C for several months.
	Reacting conditions:	250 nM; 8, 12, 16, 20 or 28 hrs
	Applications:	In HeLa cells, Rigosertib significantly inhibited PLK1 activity at all stages of the

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		cell cycle. Moreover, the loss of PLK1 activity was not due to degradation of PLK1 or inhibition of PLK1 synthesis.
	Animal experiment	
Animal models: Dosage form:	Animal models:	Nude mice bearing Bel-7402, MCF-7 or MIA-PaCa cell xenografts
	250 mg/kg; i.p.	
In Vivo	Applications:	In nude mice bearing Bel-7402, MCF-7 or MIA-PaCa cell xenografts, Rigosertib (250 mg/kg) significantly inhibited tumor growth without obvious toxicity. In addition, Rigosertib completely inhibited PLK1 activity but partially reduced CDK1 activity.
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may slightly differ with the theoretical value. This is caused by an experimental system error and it is normal.

### **Product Citations**

1. Costa, D. C. S.; Forezi, L. S. M.; et al. "A Compendium of Tyrosine-kinase Inhibitors: Powerful and Efficient Drugs against Cancer." Rev. Virtual Quim., 2017, 9 (3), 974-1064.

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### References

[1]. Gumireddy K, Reddy MV, Cosenza SC, Boominathan R, Baker SJ, Papathi N, Jiang J, Holland J, Reddy EP. ON01910, a non-ATP-competitive small molecule inhibitor of Plk1, is a potent anticancer agent. Cancer Cell. 2005 Mar;7(3):275-86.



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#### NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

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