

Product Name: Elacridar Revision Date: 04/25/2023

Product Data Sheet

Elacridar

Cat. No.: A3384

CAS No.: 143664-11-3 **Formula:** C34H33N3O5

M.Wt: 563.64

Synonyms: GF120918;GW0918;GG918;GF-120918;GF

120918

Target: Others
Pathway: Others

Storage: Store at -20°C

Solvent & Solubility

≥56.4 mg/mL in DMSO; insoluble in EtOH; insoluble in H2O

Mass Solvent 1mg 5mg 10mg Preparing Concentration In Vitro Stock Solutions 8.8709 mL 1 mM 1.7742 mL 17.7418 mL 5 mM 0.3548 mL 1.7742 mL 3.5484 mL 10 mM 0.1774 mL 0.8871 mL 1.7742 mL

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

Biological Activity

Shortsummary	BCRP inhibitor	
IC ₅₀ & Target	SIO TO THE PROPERTY OF THE PRO	agic tre Difference
	Cell Viability Assay	June 2 at action.
	Cell Line:	Human renal carcinoma cell lines 786-O and human breast cancer cell line
In Vitro		MCF-7
III VIIIO	Preparation method:	The solubility of this compound in DMSO is > 56.4mg/mL. 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:	5 μM, 24 h
Applications:	Elacridar is a P-glycoprotein inhibitor that also block ABC Sub-family B Member
.0	2 (ABCG2). Elacridar significantly enhanced sunitinib-induced cytotoxicity in
Unstaun	786-O cells. Confirmed by P-glycoprotein function assay, P-glycoprotein
O Expore the	activity was inhibited by elacridar.
Animal experiment	
Animal models:	10-14-week wild-type, Abcb1a/1b-/-, 32Abcg2-/-27 and Abcb1a/1b/Abcg2-/-
	mice, all of a >99% FVB genetic background
Dosage form:	Oral administration, 100 mg/kg
Applications:	Elacridar significantly increased sunitinib brain accumulation in wild-type mice
	(12-fold), to levels equal to those in Abcb1a/1b/Abcg2-/- mice. The sunitinib
	brain concentrations were not significantly affected by elacridar treatment in
40.	Abcb1a/1b/Abcg2-/- mice.
Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may
Paston Erdon	slightly differ with the theoretical value. This is caused by an experimental
Active Par	system error and it is normal.
	Applications: Animal experiment Animal models: Dosage form: Applications:

Product Citations

See more customer validations on www.apexbt.com.

References

- [1]. Sato H, Siddig S, Uzu M, et al. Elacridar enhances the cytotoxic effects of sunitinib and prevents multidrug resistance in renal carcinoma cells[J]. European journal of pharmacology, 2015, 746: 258-266.
- [2]. Tang S C, Lagas J S, Lankheet N A G, et al. Brain accumulation of sunitinib is restricted by P glycoprotein (ABCB1) and breast cancer resistance protein (ABCG2) and can be enhanced by oral elacridar and sunitinib coadministration[J]. International journal of cancer, 2012, 130(1): 223-233.

Caution

FOR RESEARCH PURPOSES ONLY.

NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

Specific storage and handling information for each product is indicated on the product datasheet. Most APExBIO products are stable under the recommended conditions. Products are sometimes shipped at a temperature that differs from the recommended storage temperature. Shortterm storage of many products are stable in the short-term at temperatures that differ from that required for long-term storage. We ensure that the product is shipped under conditions that will maintain the quality of the reagents. Upon receipt



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