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Product Information

Product ID CAS No.	V0147 1069-66-5		0	
Chemical Name				
Synonym	Sodium 2-propylpentanoate, Sodium valproate, 2-Propylpentanoic acid sodium salt		Olva	
Formula	C ₈ H ₁₅ NaO ₂		~	
Formula Wt.	166.19			
Melting Point	300°C	Bulk quanitites available upon request		
Purity	/ Water (50 mg/ml) Ethanol (30 mg/ml) DMSO (5 mg/ml)	Product ID	Size	
Solubility		V0147 V0147 V0147 V0147	10 g 25 g 100 g	
Store Temp	Ambient			
Ship Temp	Ambient			
Description	Valproic acid acts as an antagonist at T-type voltage-gated Ca2+ channels and voltage-gated Na+ channels; it also inhibits GABA transaminase, potentiating GABA signaling. Valproic acid is used clinically as an antiepileptic/anticonvulsant, although it also exhibits anti-inflammatory, anti-angiogenic, and anticancer chemotherapeutic activities. Valproic acid may also display antihypertensive benefit. In lung tissue, this compound prevents LPS-induced increases in TNF-α, IL-1β, NF-κB, NO, and iNOS. Valproic acid is an inhibitor of class I histone deacetylases (HDACs), primarily active against HDAC1, and downregulates			

expression of HDAC, VEGF, VEGFR2, and FGF, inhibiting tumor growth and angiogenesis in animal models.

References Zhang ZH, Hao CL, Liu P, et al. Valproic acid inhibits tumor angiogenesis in mice transplanted with Kasumi 1 leukemia cells. Mol Med Rep. 2014 Feb;9(2):443-9. PMID: 24297248.

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Zhao L, Chen CN, Hajji N, et al. Histone deacetylation inhibition in pulmonary hypertension: therapeutic potential of valproic acid and suberoylanilide hydroxamic acid. Circulation. 2012 Jul 24;126(4):455-67. PMID: 22711276.

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Caution: This product is intended for laboratory and research use only. It is not for human or drug use.