

Product Name: Gap 27 Revision Date: 11/18/2024

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# **Product Data Sheet**

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# Gap 27

	tion.
Cat. No.:	A1045
CAS No.:	198284-64-9
Formula:	C60H101N15O17
M.Wt:	1304.55
Synonyms:	Ser-Arg-Pro-Thr-Glu-Lys-Thr-Ile-Phe-Ile-Ile
Target:	Neuroscience
Pathway:	Gap Junction
Storage:	Desiccate at -20°C

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### Solvent & Solubility

	insoluble in EtOH; ≥	≥5 mg/mL in H2O;  ≥65.25 mg/mL in DMSO			
In Vitro	Preparing Stock Solutions	Solvent Concentration	1mg	5mg	10mg
		1 mM	0.7665 mL	3.8327 mL	7.6655 mL
		5 mM	0.1533 mL	0.7665 mL	1.5331 mL
		10 mM	0.0767 mL	0.3833 mL	0.7665 mL

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

## **Biological Activity**

Shortsummary

Selective gap junction blocker

#### IC<sub>50</sub> & Target

In Vitro

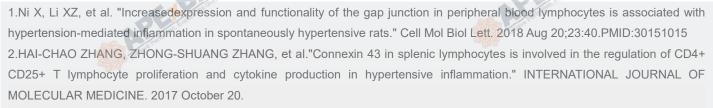
Cell	Viab	ility	Assav

Cell Viability Assay	SELECTION TO THE OWNER
Cell Line; and the	Rat osteoclasts
Preparation method:	The solubility of this peptide in sterile water is >10 mM. Stock solution should be splited and stored at -80°C for several months.
Reacting conditions:	500 μM, 48 hours
Applications:	Heptanol-treated cells acted as positive controls for gap-junctional inhibition. A

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		significant decrease could be seen in the number of both TRAP-positive
		mononuclear and multinucleated cells with Gap 27 compared to controls. The
		numbers of TRAP-positive mononuclear and multinucleated cells with both
		treatments were very similar. After the 48-hour incubation, survival of
	Burrow	osteoclasts was clearly reduced in the groups where gap-junctional
	EROOF UPS S	communication was blocked either by heptanol or Gap 27.
	Animal experiment	Align man particular
	Animal models:	Female Sprague-Dawley rats
	Dosage form:	300 μM, 45 min
	Applications:	The rats were prepared with closed cranial windows 24 h before the study. A
		10-mm-diameter craniotomy was performed over the skull midline. The dura
		was removed carefully to keep the sagittal sinus intact. An 11-mm-diameter
		glass window outfitted with three ports was glued to the skull using
	.0	cyanoacrylate. The skin overlying the window was sutured, and the animals
	Bas Unicoun	were permitted to recover. On the day of study, three stainless steel screws
	APEN BROWN	were inserted into the skull, along the periphery of the cranial window, fo
	Coneve Person	electroencephalogram (EEG) recording. Cannulae were then connected to the
		three ports. The rats were subjected to one of two neuronal activation
n Vivo		paradigms: SNS or bicuculline-induced seizure. Following the initia
		measurement of pial arteriolar diameter changes during SNS or during
		bicuculline exposure, baseline conditions were reestablished. After 20 min, a
		suffusion of gap-27 was initiated. Forty-five minutes later, the neural activation
		was repeated. Application of gap-27 peptide attenuated bicuculline-induced
		pial arteriolar dilation (by ~ 50%), without altering neuronal activation. A simila
	<b>O</b>	result was obtained with the SNS-associated pial arteriolar response, althoug
	soce ne une	the degree of reduction in the vasodilating response (~ 75%) was somewha
	Perfection En	greater.
	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 experimenta
		system error and it is normal.
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### **Product Citations**



3.Ni X, Wang A, et al."Up-regulation of gap junction in peripheral blood T lymphocytes contributes to the inflammatory response in essential hypertension." PLoS One. 2017 Sep 14;12(9):e0184773.PMID:28910394

2 www.apexbt.com

4.Koenen, Anna, et al. "Effects of renal denervation on renal pelvic contractions and connexin expression in rats." Acta Physiologica (2015).PMID:26436542

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[1] Ilvesaro J, Tavi P, Tuukkanen J. Connexin-mimetic peptide Gap 27 decreases osteoclastic activity. BMC musculoskeletal disorders, 2001, 2(1): 10.

[2] Xu H L, Mao L, Ye S, et al. Astrocytes are a key conduit for upstream signaling of vasodilation during cerebral cortical neuronal activation in vivo. American Journal of Physiology-Heart and Circulatory Physiology, 2008, 294(2): H622-H632.

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 of the product, follow the storage recommendations on the product data sheet.



#### APExBIO Technology

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