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## **Product Information**

Product ID G3552 CAS No. 14197-60-5

**Chemical Name** 

Synonym Ginsenoside Rg3

Formula C<sub>42</sub>H<sub>72</sub>O<sub>13</sub> Formula Wt. 785.01 **Melting Point** 

Purity ≥98% Solubility

## Pricing and Availability

Bulk quanitites available upon request

Product ID	Size	List Price
G3552	5 mg	\$171.30
G3552	10 mg	\$296.00
G3552	25 mg	\$591.70

Store Temp 4°C Ship Temp Ambient

**Description** Ginsenoside Rg3 is a triterpene saponin found in *Panax* that exhibits neuromodulatory, cognition enhancing, anti-inflammatory, antioxidative, anti-angiogenic, and anticancer activities. Ginsenoside Rg3 activates KCNQ1/Kv7.1 K+ channels and the y2 subunit of GABA-A receptors and inhibits the  $\alpha 10$  subunit of nicotinic acetylcholine receptors (nAChRs). In animal models, ginsenoside Rg3 decreases expression of TNF-a, IL-1B, and COX-2 in the hippocampus, improving learning and memory deficits. In vitro, ginsenoside Rg3 prevents LPS-induced upregulation of TNF-a, IL-1B, and IL-6 levels and decreases activation of microglia. In other animal models, this compound decreases oxidative stress by increasing activity of catalase, superoxide dismutase (SOD), and lysozyme and decreasing levels of NO and malondialdehyde. Ginsenoside Rg3 also exhibits anti-angiogenic benefit, inhibiting VEGF/p38/ERK signaling to inhibit tubular formation and migration of endothelial progenitor cells. In osteosarcoma cells, this compound increased DNA damage by inducing strand breaks into double-stranded DNA.

References Zhang YH, Li HD, Li B, et al. Ginsenoside Rg3 induces DNA damage in human osteosarcoma cells and reduces MNNG-induced DNA damage and apoptosis in normal human cells. Oncol Rep. 2014 Feb;31(2):919-25. PMID: 24337872.

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