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## **Product Information**

Product ID G3556 CAS No. 38243-03-7

**Chemical Name** 

Synonym 20-(R)-Ginsenoside Rg3

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

Purity ≥98%

Solubility Soluble in methanol 0.1

mg/mL. DMSO to 5 mM.

## Bulk quanitites available upon request

Product ID	Size
G3556	5 mg
G3556	10 mg
G3556	25 mg

Store Temp 4°C Ship Temp Ambient

**Description** Ginsenoside Rg3 is a triterpene saponin originally found in species of *Panax* (ginseng) that exhibits neuromodulatory, cognition enhancing, anti-inflammatory, antioxidative, anti-angiogenic, and anticancer chemotherapeutic activities. Ginsenoside Rg3 activates KCNQ1 K+ channels and the γ2 subunit of GABA-A receptors and inhibits the α10 subunit of nicotinic acetylcholine receptors (nAChRs). In animal models, ginsenoside Rg3 decreases expression of TNF-α, 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 increases 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.