

Small Molecules

Minocycline

Antibiotic with anti-inflammatory and neuroprotective properties

Catalog # 74112
74114

25 mg
100 mg



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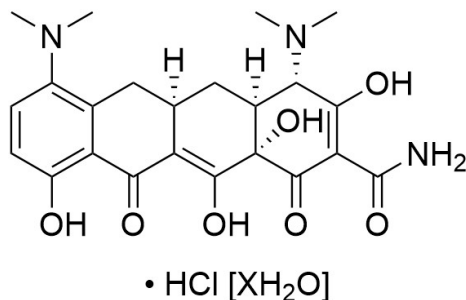
Product Description

Minocycline is a semi-synthetic tetracycline and antibiotic broadly used to treat various bacterial infections. It inhibits bacterial protein synthesis by binding to the 30S ribosomal subunit. It has also been reported to chelate with Ca^{2+} and Mg^{2+} (Lambs et al.). Reported to be anti-inflammatory, immunomodulatory, and neuroprotective (Garrido-Mesa et al.), some of its mechanism of actions include inhibiting enzymatic activities of inducible nitric oxide synthase (Amin et al.), matrix metalloproteinases (Golub et al.), and phospholipase A2 (Pruzanski et al.), inhibiting activation of caspase-1 and caspase-3 (Chen et al.), inhibiting activity of poly(ADP-ribose) polymerase 1 (Alano et al.), and reduction of p38 MAPK phosphorylation (Corbacella et al.). This product is supplied as the hydrochloride salt form of the molecule.

Molecular Name:	Minocycline (Hydrochloride Hydrate)
Alternative Names:	Minomycin; NSC 141993
CAS Number:	Not applicable*
Chemical Formula:	$\text{C}_{23}\text{H}_{27}\text{N}_3\text{O}_7 \cdot \text{HCl} \cdot \text{XH}_2\text{O}$
Molecular Weight:	493.9 g/mol
Purity:	$\geq 98\%$
Chemical Name:	4,7-bis(dimethylamino)-1,4,4 α ,5,5 α ,6,11,12 α -octahydro-3,10,12,12 α ,tetrahydroxy-1,11-dioxo-2-naphthacene-carboxamide, monohydrochloride hydrate*

*The degree to which this product is hydrated is unknown. The CAS number was previously provided but is now not applicable; 'hydrate' is now included in the Chemical Name. The product label for some lots may not be updated with these changes.

Structure:



Properties

Physical Appearance:	A crystalline solid
Storage:	Product stable at -20°C as supplied. Protect product from prolonged exposure to light. For long-term storage, store with a desiccant. Stable as supplied for 12 months from date of receipt.
Solubility:	\cdot DMSO ≤ 10 mM \cdot DMF ≤ 20 mM For example, to prepare a 5 mM stock solution in DMSO, resuspend 10 mg in 4.05 mL of DMSO.

Prepare stock solution fresh before use. Information regarding stability of small molecules in solution has rarely been reported, however, as a general guide we recommend storage in DMSO at -20°C . Aliquot into working volumes to avoid repeated freeze-thaw cycles. The effect of storage of stock solution on compound performance should be tested for each application.

Compound has low solubility in aqueous media. For use as a cell culture supplement, stock solution should be diluted into culture medium immediately before use. Avoid final DMSO concentration above 0.1% due to potential cell toxicity.

Published Applications

REPROGRAMMING

- Enables derivation of extended pluripotent stem cells from both humans and mice (Yang et al.).

IMMUNOLOGY

- Reduces the production of interleukin-8 elicited by activation of protease-activated receptor 2 in normal human epidermal keratinocytes, thus attenuating the proinflammatory reactions (Ishikawa et al.).

DISEASE MODELING

- Enhances survival of neural stem cells in rat model of Ischemia (Sakata et al.).
- Supports neuroprotective effects in animal models of Parkinson's disease (Du et al.), Huntington's disease (Chen et al.), amyotrophic lateral sclerosis (Zhu et al.), Alzheimer's disease (Choi et al.), multiple sclerosis (Brundula et al.), and spinal cord injury (Festoff et al.; Garrido-Mesa et al.).
- Reduces bone and cartilage damage in rheumatoid arthritis via inhibition of collagenase and proteinase activity (Arsenis et al.; Greenwald et al.).

References

- Alano CC et al. (2006) Minocycline inhibits poly(ADP-ribose) polymerase-1 at nanomolar concentrations. *Proc Natl Acad Sci USA* 103(25): 9685–90.
- Amin AR et al. (1997) Post-transcriptional regulation of inducible nitric oxide synthase mRNA in murine macrophages by doxycycline and chemically modified tetracyclines. *FEBS Lett* 410(2–3): 259–64.
- Arsenis C et al. (1992) Tetracyclines (TETs) inhibit the synthesis and/or activity of cartilage proteinases in vivo and in vitro. *Matrix Suppl* 1: 314.
- Brundula V et al. (2002) Targeting leukocyte MMPs and transmigration: minocycline as a potential therapy for multiple sclerosis. *Brain* 125(Pt 6): 1297–308.
- Chen M et al. (2000) Minocycline inhibits caspase-1 and caspase-3 expression and delays mortality in a transgenic mouse model of Huntington disease. *Nat Med* 6(7): 797–801.
- Choi Y et al. (2007) Minocycline attenuates neuronal cell death and improves cognitive impairment in Alzheimer's disease models. *Neuropsychopharmacology* 32(11): 2393–404.
- Corbacella E et al. (2004) Minocycline attenuates gentamicin induced hair cell loss in neonatal cochlear cultures. *Hear Res* 197(1–2): 11–8.
- Du Y et al. (2001) Minocycline prevents nigrostriatal dopaminergic neurodegeneration in the MPTP model of Parkinson's disease. *Proc Natl Acad Sci USA* 98(25): 14669–74.
- Festoff BW et al. (2006) Minocycline neuroprotects, reduces microgliosis, and inhibits caspase protease expression early after spinal cord injury. *J Neurochem* 97(5): 1314–26.
- Garrido-Mesa N et al. (2013) Minocycline: far beyond an antibiotic. *Br J Pharmacol* 169(2): 337–52.
- Golub LM et al. (1991) Tetracyclines inhibit connective tissue breakdown: new therapeutic implications for an old family of drugs. *Crit Rev Oral Biol Med* 2(3): 297–321.
- Greenwald RA et al. (1987) Tetracyclines inhibit human synovial collagenase in vivo and in vitro. *J Rheumatol* 14(1): 28–32.
- Ishikawa C et al. (2009) Tetracyclines modulate protease-activated receptor 2-mediated proinflammatory reactions in epidermal keratinocytes. *Antimicrob Agents Chemother* 53(5): 1760–5.
- Lambs L et al. (1984) Metal ion-tetracycline interactions in biological fluids. Part 3. Formation of mixed-metal ternary complexes of tetracycline, oxytetracycline, doxycycline and minocycline with calcium and magnesium, and their involvement in the bioavailability of these antibiotics in blood plasma. *Agents Actions* 14(5–6): 743–50.
- Pruzanski W et al. (1992) Inhibition of enzymatic activity of phospholipases A2 by minocycline and doxycycline. *Biochem Pharmacol* 44(6): 1165–70.
- Sakata H et al. (2012) Minocycline-preconditioned neural stem cells enhance neuroprotection after ischemic stroke in rats. *J Neurosci* 32(10): 3462–73.
- Yang Y et al. (2017) Derivation of pluripotent stem cells with in vivo embryonic and extraembryonic potency. *Cell* 169(2): 243–57.e25.
- Zhu S et al. (2002) Minocycline inhibits cytochrome c release and delays progression of amyotrophic lateral sclerosis in mice. *Nature* 417(6884): 74–8.

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