

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

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- Trockeneiszuschlag
- Gefahrgutzuschlag
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PRODUCT INFORMATION



PTIQ

Item No. 33540

CAS Registry No.: 1032822-42-6

Formal Name: 1-(3,4-dihydro-7-hydroxy-6-methoxy-

2(1H)-isoquinolinyl)-1-propanone

MF: $C_{13}H_{17}NO_3$ FW: 235.3 **Purity:** ≥95% Supplied as: A solid Storage: -20°C Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

PTIQ is supplied as a solid. A stock solution may be made by dissolving the PTIQ in the solvent of choice, which should be purged with an inert gas. PTIQ is slightly soluble in DMSO and methanol.

Description

PTIQ is a neuroprotective compound.¹ It inhibits tetrahydro-L-biopterin-induced cytotoxicity and increases in matrix metalloproteinase-3 (MMP-3) levels in CATH.a cells (IC $_{50}$ s = 1.8 and 0.06 μ M, respectively). PTIQ inhibits LPS-induced production of MMP-3, TNF-α, and COX-2 in BV-2 microglia (IC₅₀s = 2.4, 6.5, and 9.3 μM, respectively). *In vivo*, PTIQ (3 and 30 mg/kg) increases survival of striatal nigral dopaminergic neurons, reduces microglial activation, and alleviates motor deficits in a mouse model of MPTP-induced Parkinson's disease.

Reference

1. Son, H.J., Lee, J.A., Shin, N., et al. A novel compound PTIQ protects the nigral dopaminergic neurones in an animal model of Parkinson's disease induced by MPTP. Br. J. Pharmacol. 165(7), 2213-2227 (2012).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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