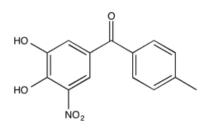


Catalog # 10-3075 Tolcapone

CAS# 134308-13-7 (3,4-Dihydroxy-5-nitrophenyl)(4-methylphenyl)methanone Ro 40-7592 Lot # X106523



Catechol O-methyltransferase inhibitor (COMT), inhibiting both brain and peripheral enzymes.¹ Potent inhibitor of alpha-synuclein and beta-amyloid oligomerization and fibrillogenesis protecting against extracellular toxicity.² Binds to transthyretin (TTR) with high affinity (21 to 58 nM) and inhibits TTR aggregation in human plasma and prevents TTR-induced cytotoxicity *in vitro*. Stabilizes TTR in mice and humans *in vivo*.³ Inhibits O-methylation of exogenous polyphenols such as EGCG.⁴ Cell permeable. Orally bioavailable

- 1) Manisto et al. (1992), *Different in vivo properties of three new inhibitors of catechol O-methyltransferase in the rat*; Br, J. Pharmacol., **105** 569
- Giovanni et al. (2010), Entacapone and tolcapone, two catechol O-methyltransferase inhibitors, block fibril formation of alpha-synuclein and beta-amyloid and protect against amyloid-induced toxicity; J. Biol. Chem., 285 14941
- 3) Sant'Anna et al. (2016), Repositioning tolcapone as a potent inhibitor of transthyretin amyloidogenesis and associated cellular toxicity; Nat. Commun., **7** 10787
- 4) Forester and Lambert (2015), *The catechol-O-methyltranseferase inhibitor, tolcapone, increases the bioavailability of unmethylated (-)-epigallocatechin-3-gallate in mice*; Funct. Foods, **17** 183

PHYSICAL DATA

Molecular Weight:	273.24
Molecular Formula:	C ₁₄ H ₁₁ NO ₅
Purity:	98% by TLC
	NMR: (Conforms)
Solubility:	Soluble in DMSO (up to 25 mg/ml) or in Ethanol (up to 25 mg/ml)
Physical Description:	Yellow solid
Storage and Stability:	Store as supplied desiccated at -20°C for up to 1 year from the date of purchase. Solutions in DMSO or ethanol may be stored at -20°C for up to 2 months.

Materials provided by Focus Biomolecules are for laboratory research use only and are not intended for human or veterinary applications.