

Catalog # 10-1048 SCH-202676

CAS# 265980-25-4 N-(2,3-Diphenyl-1,2,4-thiadiazol-5-(2H)-ylidene)methanamine hydrobromide Lot # X101310



SCH-202676 is a reversible inhibitor of both agonist and antagonist binding to G protein-coupled receptors (GPCRs).¹ IC_{50} 's = 0.1-1.7 μ M for nine GPCRs.¹ Modification of GPCRs is accomplished *via* sulfhydryl modification.^{2,3}

- 1) Fawzi et al. (2001) SCH-202676: an allosteric modulator of both agonist and antagonist binding to G protein-coupled receptors. Mol.Pharmacol. **59** 30.
- 2) Lewandowicz et al. (2006) The 'allosteric modulator' SCH-202676 disrupts G protein-coupled receptor function via sulphydryl-sensitive mechanisms Br.J.Pharmacol. **147** 422
- Goblyos et al. (2005) Synthesis and biological evaluation of a new series of 2,3,5-substituted[1,2,4]thiadiazoles as modulators of adenosine A1 receptors and their molecular mechanism of action J.Med.Chem. 48 1145

PHYSICAL DATA

Molecular Weight:	348.26
Molecular Formula:	C15H13N3S·HBr
Purity:	>98% by TLC (10% Methanol/methylene chloride + 0.1% NH4OH; Rf = 0.64)
	NMR: Conforms
Solubility:	DMSO (up to 10 mg/ml)
Physical Description:	Beige solid
Storage and Stability:	Store as supplied at room temperature for up to 1 year from the date of purchase. Protect from
	exposure to moisture. Solutions in DMSO may be stored at -20°C for up to 3 months.

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