

Saxagliptin

Catalog No: tcsc0650



Available Sizes

Size: 10mg

Size: 50mg

Size: 100mg



Specifications

CAS No:

361442-04-8

Formula:

$C_{18}H_{25}N_3O_2$

Pathway:

Metabolic Enzyme/Protease

Target:

Dipeptidyl Peptidase

Purity / Grade:

>98%

Solubility:

DMSO : ≥ 34 mg/mL (107.80 mM)

Alternative Names:

BMS-477118

Observed Molecular Weight:

315.41

Product Description

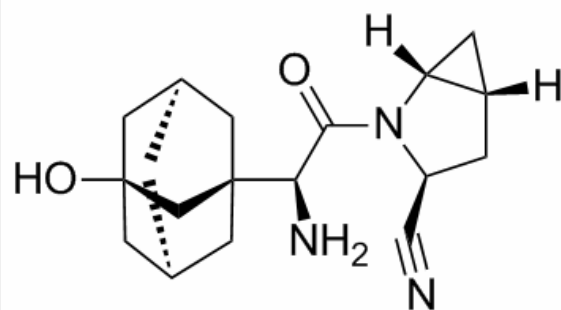
Saxagliptin(BMS477118) is a selective and reversible DPP4 inhibitor with IC₅₀ of 26 nM and K_i of 1.3 nM.

IC₅₀ value: 26 nM [1]

Target: DPP4

in vitro: Saxagliptin has an inhibition constant K_i of 1.3 nM for DPP4 inhibition, which is 10-fold more potent than either vildagliptin or sitagliptin (another two DPP4 inhibitors) with K_i of 13 and 18 nM. In addition, Saxagliptin demonstrates greater specificity for DPP4 than for either the DPP8 or DPP9 enzymes (400- and 75- fold, respectively). The active metabolite of saxagliptin is two-fold less potent than the parent. Both Saxagliptin and its metabolite are highly selective (>4000-fold) for the prevention of DPP4 compared with a range of other proteases (selectivity of sitagliptin and vildagliptin for DPP4 is >2600 and

in vivo: Saxagliptin is highly effective at eliciting marked dose-dependent enhancements in glucose clearance in the dose range 0.13-1.3 mg/kg in ob/ob mice relative to controls. Saxagliptin dose-dependently elevate plasma insulin significantly at 15 min post-oGTT, with concomitant improvement in the glucose clearance curves at 60 min post-oGTT [4].



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