

Catalog # 10-5630 Atazanavir

CAS# 198904-31-3

Methyl N-[(2S)-1-[2-[(2S,3S)-2-hydroxy-3-[[(2S)-2-(methoxycarbonylamino)-3,3-dimethylbutanoyl]amino]-4-phenylbutyl]-2-[(4-pyridin-2-ylphenyl)methyl]hydrazinyl]-3,3-dimethyl-1-oxobutan-2-yl]carbamate; BMS 232632; CGP 73547; Latazanavir Lot # X109534

Atazanavir is an azapeptide-based inhibitor of HIV protease (K_i = 2.66 nM).¹ Inhibits SARS-CoV-2 main protease (K_i = 703 nM), but not papain-like protease (K_i = 9.87 μ M).² Blocks SARS-CoV-2 replication and IL-6 / TNF α production *in vitro*.³ Also inhibits the liver enzyme UDP-glucuronyltransferase 1A1 (UGT1A1), IC₅₀ = 160 nM for recombinant UGT1A1.⁴

- 1) Robinson *et al.* (2000), *BMS-232632*, a highly potent human immunodeficiency virus protease inhibitor that can be used in combination with other available antiretroviral agents; Antimicrob. Agents Chemother. **44** 2093
- 2) Chaves et al. (2021), Atazanavir is a Competitive Inhibitor of SARS-CoV-2 M^{pro}, Impairing Variants Replication In Vitro and In Vivo; Pharmaceuticals (Basel) **15** 21
- 3) Fintelman-Rodrigues et al. (2020), Atazanavir, Alone or in Combination with Ritonavir, Inhibits SARS-CoV-2 Replication and Proinflammatory Cytokine Production; Antimicrob. Agents Chemother. **64** e00825
- 4) Mullapudi et al. (2021), UGT1A1 and UGT1A3 activity and inhibition in human liver and intestinal microsomes and a recombinant UGT system under similar assay conditions using selective substrates and inhibitors; Xenobiotica 51 1236

PHYSICAL DATA

 $\begin{tabular}{lll} Molecular Weight: & 704.87 \\ Molecular Formula: & $C_{38}H_{52}N_6O_7$ \\ Purity: & $>98\% \ by \ HPLC$ \\ NMR: (Conforms) \\ \end{tabular}$

Solubility: DMSO (45 mg/ml)
Physical Description: White to off-white solid

Storage and Stability: Store as supplied at -20°C for up to 2 years from the date of purchase. Solutions in

DMSO may be stored at -20°C for up to 3 months.

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