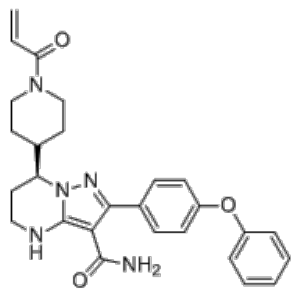


Product Name	: (S)-Zanubrutinib	
Synonyms	: S)-Zanubrutinib; BGB3111; BGB 3111	
Cat No.	: M12579	
CAS Number	: 1691249-45-2	
Molecular Formula	: C ₂₇ H ₂₉ N ₅ O ₃	
Formula Weight	: 471.56	
Chemical Name	(7S)-4,5,6,7-Tetrahydro-7-[1-(1-oxo-2-propen-1-yl)-4-piperidinyl]-2-(4-phenoxyphenyl)pyrazolo[1,5-a]pyrimidine-3-carboxamide	
Description	(S)-Zanubrutinib (BGB3111) is a potent, selective and orally available Btk inhibitor; shows much more restricted off-target activities against a panel of kinases, including ITK, compared with Ibrutinib; demonstrates nanomolar BTK inhibition activity, inhibits BCR aggregation-triggered BTK autophosphorylation, blocks downstream PLC-γ2 signaling, and potently inhibits cell proliferation in several MCL and DLBCL cell lines; demonstrates better anti-tumor activity than ibrutinib in TMD-8 subcutaneous xenograft model.	
Pathway	: Tyrosine Kinase	
Target	: BTK	
Receptor	: BTK	
Solubility	: DMSO : ≥ 56.75 mg/mL 120.35 mM; Ethanol : < 1 mg/mL	
SMILES	: <chem>C=CC(=O)N1CCC(CC1)C2CCNC3=C(C(=NN2)C4=CC=C(C(=C4)OC5=CC=CC=C5)C(=O)N</chem>	
Storage	: (-20°C)	
Stability	: ≥ 2 years	
Reference	:	

1. Na Li, et al. Abstract 2597: BGB-3111 is a novel and highly selective Bruton's tyrosine kinase (BTK) inhibitor. AACR.