



Product Information

PARP inhibitor, RK-287107, Purity $\geq 98\%$

Cat. No.: X24-08-YM008

Size: 5 mg; 10 mg; 50 mg; 100 mg

CAS Number: 2171386-10-8

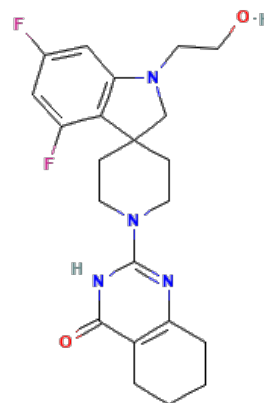
Compound CID: 137701512

Synonym: 2171386-10-8; RK287107; RK 287107;

2-[4,6-Difluoro-1-(2-hydroxyethyl)spiro[2H

-indole-3,4'-piperidine]-1'-yl]-5,6,7,8-tetrahydro-3H-quinazolin-4-one

This product is for research use only and is not intended for diagnostic use.



Product Information

Description	RK-287107, soluble in DMSO, water and ethanol, is an effective DNA damage inhibitor, that has the ability to prevent the pathway of epigenetics by inhibiting PARP. It targets the tankyrase-1 and tankyrase-2.
Molecular Weight	416.46
Molecular Formula	C ₂₂ H ₂₆ F ₂ N ₄ O ₂
Targets	Tankyrase-1: 14.3 nM (IC ₅₀); Tankyrase-2: 10.6 nM (IC ₅₀)
IUPAC Name	2-[4,6-Difluoro-1-(2-hydroxyethyl)spiro[2H -indole-3,4'-piperidine]-1'-yl]-5,6,7,8-tetrahydro-3H-quinazolin-4-one
InChI	InChI=1S/C22H26F2N4O2/c23-14-11-16(24)19-18(12-14)28(9-10-29)13-22(19)5-7-27(8-6-22)21-25-17-4-2-1-3-15(17)20(30)26-21/h11-12,29H,1-10,13H2,(H,25,26,30)
InChI Key	FZQYCOUBRJEYBC-UHFFFAOYSA-N
Canonical SMILES	C1CCC2=C(C1)C(=O)NC(=N2)N3CCC4(CC3)CN(C5=C4C(=CC(=C5)F)F)CCO
Form	Lyophilized powder
Purity	$\geq 98\%$
Titer	Free from inappropriate visible particulates, foreign matter, discoloration, or other defects
Solubility	<i>In vitro</i> : DMSO: 81 mg/mL (194.49 mM); Water: <1 mg/mL; Ethanol: <1 mg/mL
Stability	In its lyophilized form, the chemical remains stable for 36 months.
Quality Level	Research grade
Applications	RK-287107 is an epigenetic inhibitor that can be studied extensively for its potential therapeutic applications in the treatment of cancers and inflammatory diseases by disrupting the NF- κ B



signaling pathway, which plays a role in promoting cell survival and proliferation.

Storage

Store at -20°C, and keep desiccated.
