Data Sheet (Cat.No.T23028)



MRT67307 HCl (1190378-57-4 free base)

Chemical Properties

CAS No.:

Formula: C26H36N6O2· xHCl

Molecular Weight:

Appearance: no data available

Storage: Powder: -20°C for 3 years | In solvent: -80°C for 1 year

Biological Description

Description	MRT67307 is an inhibitor for TBK1, IKKE, MARK1-4 and NUAK1 with IC50 value of 19, 160,
	27-52 and 230nM, respectively [1]. It is an inhibitor for ULK1 and ULK2 with IC50 value of
	45 and 38nM, respectively [2]. Also, it is a salt inducible kinase (SIK) inhibitor with IC50
	value of 250, 67 and 430nM for SIK1, SIK2 and SIK3, respectively. siKs prevent the
	formation of regulatory macrophages and their inhibition induces increasing in some
	markers of regulatory macrophages, such as IL-10 and other anti-inflammatory
	molecules. IKKs and TBK-1 mediate the phosphorylation of interferon regulatory factor 3
	(IRF3). MARK1 is a Serine/threonine-protein kinase. In macrophages, MRT67307
	prevented the production of IFNβ and the phosphorylation of IRF3 without suppressing
	the activation of NF-kB, which showed that MRT67307 blocked the induction of Pellino 1
	through inhibiting TBK1/IKKs kinase activity [1] [3]. Also, MRT67307 completely blocked
	the TBK1- or IKKε-induced decrease in the mobility of Pellino 1 [3]. Exposed
	macrophages to MRT67307 increased the levels of the anti-inflammatory cytokines IL-
	1ra and IL-10 and decreased the levels of proinflammatory cytokines (such as IL-6, IL-
	12, and TNF) in response to bacterial lipopolysaccharide (LPS) [4].
Targets(IC50)	Others

Solubility Information

Solubility	H2O: > 52.5 mg/mL,Heating is recommended.	
	Ethanol: ≥30.67 mg/mL,	
	DMSO: ≥23.25 mg/mL,	
	(< 1 mg/ml refers to the product slightly soluble or insoluble)	

Inhibitor · Natural Compounds · Compound Libraries · Recombinant Proteins

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