

产品名称: **ONO-4059 (analog)**

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生物活性:					
Description	ONO-4059 analog is the analog of ONO-4059, ONO-4059 is a highly potent and selective Btk inhibitor. IC50 value: sub-nM range [2] Target: Btk in vitro: ONO-4059 is a selective, once-daily, oral inhibitor of BTK, which has been shown to play a role in the survival and proliferation of malignant B-cells. ONO-4059 shows a favourable safety profile along with promising efficacy in this difficult-to-treat patient population. [1] in vivo: ONO-4059 has demonstrated anti-tumour activity in several pre-clinical models.[1] ONO-4059 potently and dose-dependently reverse clinical arthritis and prevented bone damage in the CIA model.[2]				
Solvent&Solubility	In Vitro: DMSO : ≥ 29 mg/mL (63.53 mM) * "≥" means soluble, but saturation unknown.				
		Mass	1 mg	5 mg	10 mg
	Preparing	Concentration			
	Stock Solutions				
	1 mM	2.1906 mL	10.9529 mL	21.9058 mL	
	5 mM	0.4381 mL	2.1906 mL	4.3812 mL	
	10 mM	0.2191 mL	1.0953 mL	2.1906 mL	
*请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液, 一旦配成溶液, 请分装保存, 避免反复冻融造成的产品失效。 储备液的保存方式和期限 -80°C, 6 months; -20°C, 1 month。 -80°C 储存时, 请在 6 个月内使用, -20°C 储存时, 请在 1 个月内使用。					
References	[1]. Simon Rule, A Phase I Study Of The Oral Btk Inhibitor ONO-4059 In Patients With Relapsed/Refractory B-Cell Lymphoma. November 15, 2013; Blood: 122 (21) [2]. Toshio Yoshizawa, et al. Development of a Bruton's Tyrosine Kinase (Btk) Inhibitor, ONO-4059: Efficacy in a Collagen Induced Arthritis (CIA) Model Indicates Potential Treatment for Rheumatoid Arthritis (RA). Washington, DC November 9-14, 2012. [3]. Yamamoto, et al. Preparation of purinone derivatives as selective Btk inhibitors. From PCT Int. Appl. (2011), WO 2011152351 A1 20111208.				