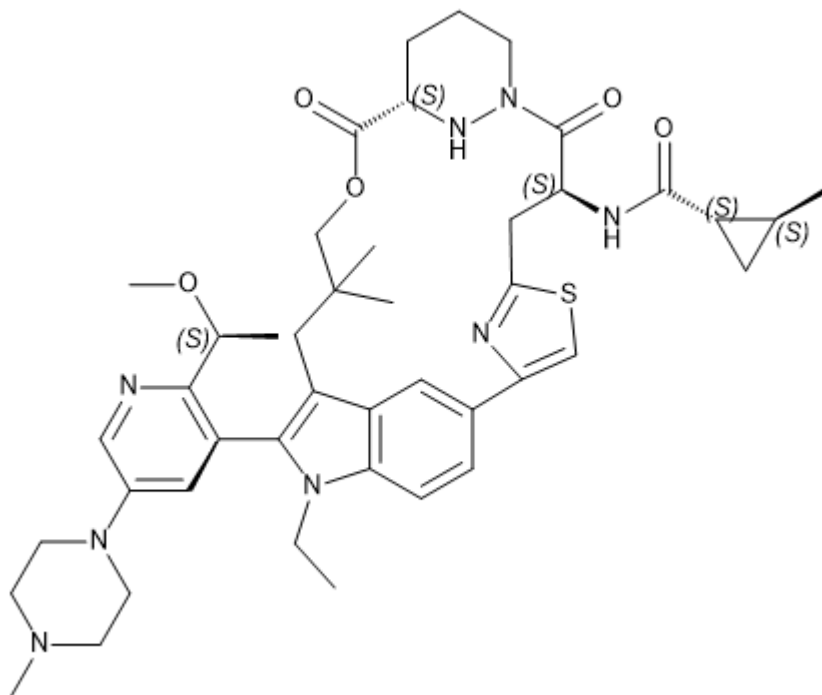


**Name: RMC-6236 cat#:** EX-A6631

Target:: Ras

Pathway: GPCR/G Protein; MAPK/ERK Pathway

Chemical Structure:



Chemical Name	(1S,2S)-N-((63S,4S,Z)-11-ethyl-12-(2-((S)-1-methoxyethyl)-5-(4-methylpiperazin-1-yl)pyridin-3-yl)-10,10-dimethyl-5,7-dioxo-6,1,6,2,6,3,6,4,6,5,6,6-hexahydro-11H-8-oxa-2(4,2)-thiazola-1(5,3)-indola-6(1,3)-pyridazinacycloundecaphane-4-yl)-2-methylcyclopropane-1-carboxamide		
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Molecular Weight	811.047	Storage	3 years -20°C powder
Formula	C44H58N8O5S		6 months -80°C in solvent Away from light
CAS No.	2765081-21-6	Synonyms	RMC 6236; RMC6236

Solubility (25°C) *	In vitro	DMSO	Soluble, >100mg/mL (Need ultrasonic)
		Ethanol	N/A
		Water	N/A
	In vivo (should be freshly prepared each time)		

\* <1 mg/ml means slightly soluble or insoluble.

\* Please note that Selleck tests the solubility of all compounds in-house, and the actual solubility may differ slightly from published values. This is normal and is due to slight batch-to-batch variations.

Preparing Stock Solutions:

Concentration \ Volume	Mass	1 mg	5 mg	10 mg
	Volume			
1 mM		1.2330 mL	6.1649 mL	12.3297 mL
5 mM		0.2466 mL	1.2330 mL	2.4659 mL
10 mM		0.1233 mL	0.6165 mL	1.2330 mL

\*The above data is based on the product molecular weight 811.047.

Biological Activities:

<b>Description</b>	RMC-6236 is a first-in-class, RAS-selective, orally bioavailable, tri-complex RAS <sup>MULTI</sup> (ON) inhibitor.
<b>IC<sub>50</sub> &amp; Target</b>	KRas G12D
<b>In Vitro</b>	RMC-6236 (Compound A122) (5 days) inhibits cell viability with an IC <sub>50</sub> of 1-10 μM against AsPC-1 (K-Ras G12D).

<b>References</b>	<p>[1]. <a href="#">Gustafson, W. Clay et al. Direct targeting of RAS in pancreatic ductal adenocarcinoma with RMC-6236, a first-in-class, RAS-selective, orally bioavailable, tri-complex RAS<sup>MULTI</sup> (ON) inhibitor. Journal of Clinical Oncology 2022 40:4_suppl, 591-591.</a></p> <p>[2]. <a href="#">Ethan AHLER, et al. Use of sos1 inhibitors to treat malignancies with shp2 mutations. Patent WO2022060583A1.</a></p>
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