

## Modzatinib

## Chemical Properties

CAS No. : 2411407-25-3

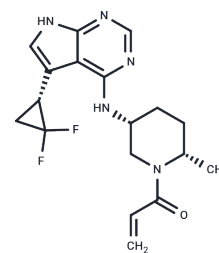
Formula: C<sub>18</sub>H<sub>21</sub>F<sub>2</sub>N<sub>5</sub>O

Molecular Weight: 361.39

Storage: Keep away from moisture, Keep away from direct sunlight, Store at low temperature, Store under nitrogen

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

Actual storage temperature shall be subject to the COA.



## Biological Description

Description	Modzatinib (ITKkinase-IN-1) (compound I) is an inhibitor targeting ITK kinase with an IC <sub>50</sub> of 8 nM in Jurkat cells, useful for researching inflammatory diseases. Modzatinib (ITKkinase-IN-1) (compound I) inhibits JAK3 in hPBMC cells with an IC <sub>50</sub> of 23 nM (based on IL-2/pSTAT5 levels) and shows no inhibitory effect on JAK2. Modzatinib (ITKkinase-IN-1) (compound I) is therefore applied in immunological signaling research to study T-cell receptor-associated kinase pathways, JAK/STAT signaling selectivity, and cytokine response modulation in primary immune cell systems.
Targets(IC <sub>50</sub> )	JAK, Tyrosine Kinases
In vitro	Methods: Whole blood was treated with modzatinib at concentrations of 0.01–1000 ng/mL for 2–18 hours, and its effects on related signaling pathways and gene expression were detected. Results: Modzatinib inhibited aCD3/CD28-induced mRNA expression of IL2 and IFN $\gamma$ , as well as IL15-mediated activation of pSTAT5 [1].
In vivo	Methods: Rat adjuvant-induced arthritis (AIA) model and mouse T cell transfer colitis model were established. Modzatinib was administered continuously via different routes, and changes in inflammation-related indicators were observed. Results: 1. In rat AIA model, intragastric administration of Modzatinib at 5–15 mg/kg twice daily for 14 consecutive days alleviated ankle swelling in a dose-dependent manner. 2. In mouse T cell transfer colitis model, dietary administration of the compound at 100–300 ppm once daily for 35 consecutive days ameliorated inflammatory lesions in colon and ileum dose-dependently [1].

## Solubility Information

Solubility	DMSO: 4.5 mg/mL (12.45 mM), Sonication is recommended. (< 1 mg/ml refers to the product slightly soluble or insoluble)
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### Preparing Stock Solutions

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	1mg	5mg	10mg
1 mM	2.7671 mL	13.8355 mL	27.6709 mL
5 mM	0.5534 mL	2.7671 mL	5.5342 mL
10 mM	0.2767 mL	1.3835 mL	2.7671 mL
50 mM	0.0553 mL	0.2767 mL	0.5534 mL

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Please select the appropriate solvent to prepare the stock solution, according to the solubility of the product in different solvents. Please use it as soon as possible.

Note: The dilution table applies only to solid products. For liquid products, please calculate the stock solution based on the stated concentration and/or density.

### Reference

David Randolph ANDERSON, et al. Pyrrolopyrimidine compositions for treatment of itk mediated conditions. WO2024102778A1. 2024-05-16.

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