Inhibitors

Balsalazide sodium hydrate

Cat. No.: HY-B0667A CAS No.: 150399-21-6 Molecular Formula: C₁₇H₁₇N₃Na₂O₈

Molecular Weight: 437.31

Interleukin Related; STAT Target:

Pathway: Immunology/Inflammation; JAK/STAT Signaling; Stem Cell/Wnt

Storage: 4°C, sealed storage, away from moisture and light

* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture

and light)

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 125 mg/mL (285.84 mM; Need ultrasonic) H₂O: 100 mg/mL (228.67 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.2867 mL	11.4335 mL	22.8671 mL
	5 mM	0.4573 mL	2.2867 mL	4.5734 mL
	10 mM	0.2287 mL	1.1434 mL	2.2867 mL

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description	Balsalazide sodium hydrate could suppress colitis-associated carcinogenesis through modulation of IL-6/STAT3 pathway.			
IC ₅₀ & Target	IL-6	IL-1	STAT3	
In Vivo	At the endpoint, the protein production of MIP-1 β , MCP-1, IL-6, and IL-10 in the colon tissues decrease in concordance with the plasma concentrations of the cytokines. The drug-treated groups reveal lower expression of p-STAT3 compared to the CAC group. In addition, BCL2 decreases and BAX increases markedly in the BSZ+VSL#3 group ^[1] . Balsalazide is a new 5-aminosalicylic acid (5-ASA) containing prodrug ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.			

REFERENCES

[1]. Do EJ, et al. Suppression of colitis-associated carcinogenesis through modulation of IL-6/STAT3 pathway by balsalazide and VSL#3. J Gastroenterol Hepatol. 2016 Aug;31(8):1453-61.

e balsalazide (1.5 g twice daily) and mesalazine (0.5 g three times daily) maintained re n preventing relapses. Gut, 2001. 49(6): p. 783-789.	ernission of uicerative colitis but high dose balsalazide (3
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