## STOCK2S-26016

Cat. No.: HY-112143 CAS No.: 332922-63-1 Molecular Formula:  $C_{20}H_{19}N_{3}O_{2}$ Molecular Weight: 333.38

Target: Ser/Thr Protease

Pathway: Metabolic Enzyme/Protease Storage: Powder -20°C 3 years In solvent -80°C 6 months

> -20°C 1 month

**Product** Data Sheet

## **BIOLOGICAL ACTIVITY**

Description	STOCK2S-26016 is a WNK signalling inhibitors. STOCK2S-26016 inhibits WNK4 and WNK1 with IC $_{50}$ s of 16 $\mu$ M and 34.4 $\mu$ M, respectively. STOCK2S-26016 has potential for antihypertensive research <sup>[1]</sup> .
IC <sub>50</sub> & Target	IC50: 16 μM (WNK4); 34.4 μM (WNK1) <sup>[1]</sup>
In Vitro	STOCK2S-26016 (10 and 30 minutes) blocks the effect of L-NAME on phosphorylated sodium-chloride cotransporter (NCC) in mDCT cells $^{[2]}$ . STOCK2S-26016 (25-200 $\mu$ M) drastically and dose-dependently reduces the phosphorylation of STE20/SPS1-related proline/alanine-rich protein kinase (SPAK) and NCC in mpkDCT cells $^{[1]}$ . STOCK2S-26016 (50-200 $\mu$ M) drastically and dose-dependently reduces the phosphorylation of SPAK and Na/K/Cl cotransporter 1 (NKCC1) in MOVAS cells $^{[1]}$ . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **REFERENCES**

[1]. Takayasu M, et, al. Chemical library screening for WNK signalling inhibitors using fluorescence correlation spectroscopy. Biochem J. 2013 Nov 1; 455(3): 339-45.

[2]. Conghui W, et, al. Low dose L-NAME induces salt sensitivity associated with sustained increased blood volume and sodium-chloride cotransporter activity in rodents. Kidney Int. 2020 Jun 24; S0085-2538(20)30703-1.

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA