



**VAS2870** 

Catalog No: tcsc0012525

Available Sizes		
Size: 5mg		
Size: 10mg		
Size: 25mg		
Size: 50mg		
Size: 100mg		
Specifications		
<b>CAS No:</b> 722456-31-7		
Formula: C <sub>18</sub> H <sub>12</sub> N <sub>6</sub> OS		
Pathway: Others		
<b>Target:</b> Others		
Purity / Grade: >98%		
Solubility: DMSO: 83.3 mg/mL (231.14	mM; Need ultrasonic)	
<b>Observed Molecular Weig</b> 360.39	ht:	





## **Product Description**

VAS2870 is a NADPH oxidase (NOX) inhibitor.

IC50 & Target: Target: NADPH oxidase<sup>[1]</sup>

In Vitro: VAS2870 is effective to suppress PDGF-BB-dependent activation of NADPH oxidase and subsequent production of intracellular ROS. Furthermore, VAS2870 suppresses PDGF-BB-dependent chemotaxis, but not DNA synthesis. Preincubation with VAS2870 (10 and 20  $\mu$ M) completely abolishes PDGF-mediated NADPH oxidase activation and ROS production. Preincubation with VAS2870 (0.1-20  $\mu$ M) does not affect PDGF-induced cell cycle progression. However, it abolishes PDGF-dependent chemotaxis of VSMC in a concentration-dependent manner (100% inhibition at 10  $\mu$ M)<sup>[1]</sup>. VAS2870 inhibits dose-dependently autocrine increase of cell number in FaO rat hepatoma cells, and almost completely blocked ROS production and thymidine incorporation when used at 25 mM. VAS2870 blocks serum-dependent cell growth of FaO rat hepatoma cells. VAS2870 inhibits proliferation of different human hepatocellular carcinoma (HCC) cell lines. VAS2870 pretreatment enhances TGF-b-mediated apoptosis of FaO rat hepatoma cells<sup>[2]</sup>.

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!