

## **BAY-678**

ALTERNATE NAMES:	5-[(6R)-5-acetyl-4-methyl-2-oxo-3-[3-(trifluoromethyl)phenyl]-1,6-dihydropyrimidin-6-yl]pyridine-2- carbonitrile; (R)-5-(5-Acetyl-6-methyl-2-oxo-1-(3-(trifluoromethyl)phenyl)-1,2,3,4-tetrahydropyrimidin-4- yl)picolinonitrile; 5-[(4R)-5-Acetyl-1,2,3,4-tetrahydro-6-methyl-2-oxo-1-[3-(trifluoromethyl)phenyl]-4- pyrimidinyl]-2-pyridinecarbonitrile
CATALOG #:	B3061-5 5 mg B3061-25 25 mg
STRUCTURE:	
MOLECULAR FORMULA:	$C_{20}H_{15}F_{3}N_{4}O_{2}$
MOLECULAR WEIGHT:	400.35
CAS NUMBER:	675103-36-3
APPEARANCE:	Yellow solid
PURITY:	≥98%
SOLUBILITY:	~20 mg/ml in DMSO
DESCRIPTION:	BAY-678 is a potent, selective, cell-permeable inhibitor of human neutrophil elastase (HNE). The activity of HNE is increased in inflammatory diseases. BAY-678 shows $K_i$ values of 15 nM and 600 nM for human and rat neutrophil elastase respectively and IC <sub>50</sub> of 20 nM <i>in vitro</i> . It shows beneficial pulmonary hemodynamic and vascular effects in animal models of pulmonary arterial hypertension. The S-enantiomer BAY-677 is a negative control for BAY-678.
STORAGE TEMPERATURE:	-20°C
HANDLING:	Do not take internally. Wear gloves and mask when handling the product! Avoid contact by all modes of exposure.
REFERENCES:	<ol> <li>von Nussbaum, F., Li, V. MJ., Allerheiligen. S., et al. Freezing the Bioactive Conformation to Boost Potency: The Identification of BAY 85-8501, a Selective and Potent Inhibitor of Human Neutrophil Elastase for Pulmonary Diseases. Chem. Med. Chem., 10, 1163 – 1173 (2015).</li> <li>von Nussbaum, F., Li, V. MJ. Neutrophil elastase inhibitors for the treatment of (cardio) pulmonary diseases: Into clinical testing with pre-adaptive pharmacophores. Bioorganic &amp; Medicinal Chemistry Letters 25 4370–4381 (2015).</li> </ol>
RELATED PRODUCTS:	
BAY-677 (Cat. No. B3060) Elastase Inhibitor (Cat. No. 1922) Alvelestat (Cat. No. B1653) Elastase Inhibitor, SPCK (Cat. No. 1921) Sivelestat (Cat. No. B2144)	

DISCLAIMER:

FOR RESEARCH USE ONLY! Not to be used on humans.