# BAY 61-3606. hydrochloride

Syk inhibitor

Cell permeable, potent, ATP-competitive and reversible inhibitor of spleen tyrosine kinase (Syk) (IC $_{50}$ =10nM). Highly selective if compared to other kinases; shows no inhibitory effect against Btk, Fyn, Itk, Lyn, and Src up to concentrations of 4.7 $\mu$ M. Anti-inflammatory and orally active. Exhibits good *in vivo* efficacy in the treatment of various allergy and asthma conditions in rat models.

Citations: 5

**View Online »** 

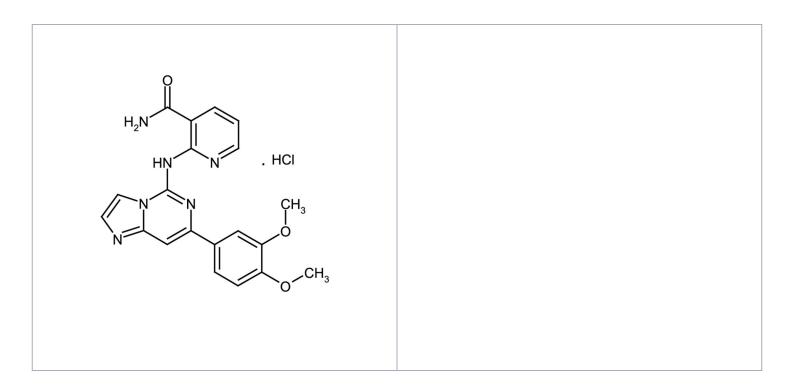
## **Ordering Information**

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ALX-270-479-M001	1mg
ALX-270-479-M005	5mg
ALX-270-479-M025	25mg

Manuals, SDS & CofA

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### **Handling & Storage**

**Use/Stability** As indicated on product label or CoA when stored as recommended.

**Handling** Protect from light. Hygroscopic.

Long Term Storage -20°C

**Shipping** Ambient Temperature

#### Regulatory Status RUO - Research Use Only

#### **Product Details**

Alternative Name 2-[[7-(3,4-Dimethoxyphenyl)imidazo[1,2-c]pyrimidin-5-yl]amino]pyridine-3-carboxamide .

HCI

Appearance Yellow solid.

**CAS** 648903-57-5

Couple Target Syk

Couple Type Inhibitor

Formula  $C_{20}H_{18}N_6O_3$  . HCI

**Identity** Determined by MS.

**MW** 390.4 . 36.5

Purity ≥98% (HPLC(UV))

**Solubility** Soluble in water (30mg/ml), DMSO (2mg/ml) or methanol (1mg/ml).

