PRODUCT INFORMATION



PS-1145

Item No. 14862

CAS Registry No.: 431898-65-6

Formal Name: N-(6-chloro-9H-pyrido[3,4-b]

indol-8-yl)-3-pyridinecarboxamide

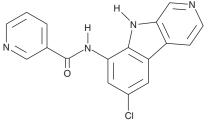
Synonyms: **IKK Inhibitor X** MF: $C_{17}H_{11}CIN_4O$

322.8 FW: **Purity:** ≥98%

UV/Vis.: λ_{max} : 239, 293, 364 nm A crystalline solid Supplied as:

Storage: -20°C Stability: ≥2 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.



Description

PS-1145 is an IkB kinase β (IKK β) inhibitor (IC₅₀ = 88 nM).¹ It selectively inhibits IKK β over a panel of 14 kinases (IC₅₀s = >100 nM for all). PS-1145 also inhibits the IKK complex (IC₅₀ = 100 nM).² It inhibits the proliferation of MM.1S, U266, and RPMI-8226 myeloma cells when used at concentrations ranging from 1.5 to 50 μ M. PS-1145 inhibits TNF- α -induced activation of NF- κ B in MM.1S cells in a concentrationdependent manner. It inhibits RANKL-induced differentiation of RAW 264.7 cells into osteoclasts when used at concentrations of 5 and 10 μM.³ PS-1145 (10 μM) inhibits RANKL-induced phosphorylation of IκBα, ERK, and JNK in RAW 264.7 cells.

References

- 1. Hideshima, T., Chauhan, D., Richardson, P., et al. NF-κB as a therapeutic target in multiple myeloma. The Journal of Biological Chemisty 277(19), 16639-16647 (2002).
- 2. Castro, A.C., Dang, L.C., Soucy, F., et al. Novel IKK inhibitors: β-carbolines. Bioorg. Med. Chem. Lett. 13(14), 2419-2422 (2003).
- 3. Tsunekage, Y., Takeiri, M., Yoshioka, Y., et al. Nasturtium officinale extract suppresses osteoclastogenesis in RAW 264 cells by inhibiting IκB-kinase β. Nat. Prod. Commun. 16(6), 1-9 (2021).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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