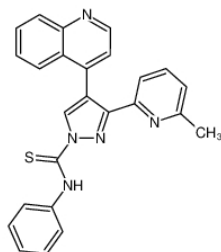


**A 83-01 (CAS: 909910-43-6)****Catalog #: EBC51012****Biological Activity**

<b>Synonyms</b>	ALK5 Inhibitor IV
<b>Chemical Name</b>	3-(6-Methylpyridin-2-yl)-N-phenyl-4-quinolin-4-ylpyrazole-1-carbothioamide
<b>Application</b>	A 83-01 is a small-molecule inhibitor of activin receptorlike kinase (ALK)-5
<b>CAS No.</b>	909910-43-6
<b>Purity</b>	≥98.0%
<b>Molecular Weight</b>	421.52
<b>Molecular Formula</b>	C <sub>25</sub> H <sub>19</sub> N <sub>5</sub> S
<b>Shipping</b>	Gel Pack
<b>Storage</b>	Store at -20° C
<b>Target &amp; IC<sub>50</sub></b>	ALK5 kinase: IC <sub>50</sub> = 12 nM ALK4: IC <sub>50</sub> = 45 nM ALK7: IC <sub>50</sub> = 7.5 nM

**Molecular Structure****Solubility**

DMSO: 10mg/mL (23.72 mM)

**PS:** < 1 mg/ml refers to the product insoluble

**Description**

A 83-01 is a selective inhibitor of TGF- $\beta$  type I receptor ALK5 kinase, type I activin/nodal receptor ALK4 and type I nodal receptor ALK7 (IC<sub>50</sub> values are 12, 45 and 7.5 nM respectively). A 83-01 blocks phosphorylation of Smad2 and inhibits TGF- $\beta$ -induced epithelial-to-mesenchymal transition. A 83-01 only weakly inhibits ALK-1, -2, -3, -6 and MAPK activity. A 83-01 is more potent than SB 431542. Fibroblasts can be reprogrammed by A 83-01 into alternative lineages.

**For Research Use Only. Not For Use In Diagnostic Procedures**

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