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# DATASHEET A83-01

#### **Product overview**

Name	A83-01
Cat No	HB3218
Alternative names	A83
Biological action	Inhibitor
Purity	>98%
Customer comments	Good quality and very affordable price. My lab are very happy with this product from Hello Bio, we will definitely purchase this product again. <b>Verified customer, UK</b>
Description	Selective TGF-βRI (ALK5), ALK4 and ALK7 inhibitor. Maintains human hiPSC self renewal, 3D growth matrix component and promotes long-term organoid growth. Aids fibroblast to NSC/cardiomyocyte reprogramming.

### Images



## **Biological Data**

Biological description	A 83-01 is a selective TGF- $\beta$ RI (ALK5), ALK4 and ALK7 inhibitor (IC <sub>50</sub> values are 12, 45 and 7.5 nM at ALK5, ALK4 and ALK7 respectively). It is more potent than SB431542
	Reprogramming
	A 83-01 aids reprogramming of fibroblasts into neural stem cells and cardiomyocytes.
	Differentiation
	A 83-01 blocks phosphorylation of SMAD2/3 to inhibit TGF- $\beta$ -induced epithelial-to-mesenchymal transition
	Maintenance / self-renewal
	A 83-01 helps to maintain homogeneity and long-term in vitro self-renewal of iPSCs
	Organoids
Application notes	3D growth matrix component and also promotes long-term organoid growth. Figures 1 - 3 show liver hepatocellular carcinoma tumouroids cultured with A 83-01 and forskolin

Storage instructions Solubility overview Handling -20 °C (protect from light) Soluble in DMSO (50mM)

- This compound is light sensitive; we therefore recommend protecting the solid material and solutions from exposure to light.
- It has been reported that A 83-01 can decompose to A 77-01 in solution. We therefore recommend that you make up solutions and use immediately. Do not store solutions.
- If storage of solutions is required, you should aliquot out the solution into tightly sealed vials and store at -20 °C and store these for up to one month.

Important

This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use.

## **Chemical Data**

3-(6-Methyl-2-pyridinyl)- <i>N</i> -phenyl-4 -(4-quinolinyl)-1 <i>H</i> -pyrazole-1-carbothioamide 421.52
$C_{25}H_{19}N_5S$
909910-43-6
16218924
CC1=CC=CC(=N1)C2=NN(C=C2C3=CC=NC4=CC=CC=C34)C(=S)NC5=CC=CC=C5
InChI=1S/C25H19N5S/c1-17-8-7-13-23(27-17)24-21(19-14-15-26-22-12-6-5-11-20(19)22)16-30(29-24)25(31)28-18-9-3-2-4-10-18/h2-16H,1H3,(H,28,31)
HIJMSZGHKQPPJS-UHFFFAOYSA-N
Pale yellow solid

#### References

The ALK-5 inhibitor A-83-01 inhibits Smad signaling and epithelial-to-mesenchymal transition by transforming growth fact beta.	
Tojo et al (2005) Cancer Sci 96(11 <b>PubMedID</b>	) 16271073
Brief report: combined chemica	Il treatment enables Oct4-induced reprogramming from mouse embryonic fibroblasts.
Yuan et al (2011) 2011 29(3) PubMedID	21425417
Generation of rat and human in	duced pluripotent stem cells by combining genetic reprogramming and chemical inhibitors.
Li et al (2009) Cell Stem Cell 4(1)	
PubMedID	19097958
Chemical compound-based dire	ect reprogramming for future clinical applications.
Takeda et al (2018) Biosci Rep 38 <b>PubMedID</b>	(3) 29739872