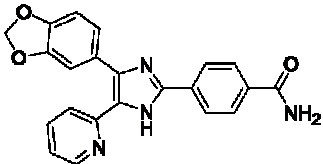


## Product Specification Sheet

<b>Product Name</b>	Stemolecule™ SB431542
<b>Description</b>	Stemolecule SB431542 is an inhibitor of the TGF-β1 activin receptor-like kinases (ALKs). It is a selective and potent inhibitor of ALK-4, -5 and -7. SB431542 inhibits endogenous activin and TGF-β signaling without affecting more divergent BMP signaling utilizing ALK-1, -2, -3, and -6 <sup>1,2</sup> . SB431542 stimulates proliferation, differentiation, and sheet formation of endothelial cells derived from embryonic stem cells <sup>3</sup> .
<b>Catalog Number</b>	AMS.04-0010
<b>Size</b>	5 mg
<b>Alternate Name</b>	4-[4-(1,3-benzodioxol-5-yl)-5-pyridin-2-yl-1H-imidazol-2-yl]benzamide
<b>Chemical Formula</b>	C <sub>22</sub> H <sub>16</sub> N <sub>4</sub> O <sub>3</sub>
<b>Structure</b>	
<b>Molecular Weight</b>	384.4
<b>CAS Number</b>	301836-41-9 (anhydrous)
<b>Purity</b>	Greater than 98% by HPLC analysis
<b>Formulation</b>	White solid
<b>Solubility</b>	For a 10 mM concentrated stock solution of SB431542, reconstitute the compound by adding 1.3 ml of DMSO to the entire contents of the vial. If precipitate is observed, warm the solution to 37°C for 2 to 5 minutes. For cell culture, the media should be prewarmed prior to adding the reconstituted compound. Note: for most cells, the maximum tolerance to DMSO is greater than 0.5%. This molecule is soluble in DMSO at 100 mM.
<b>Storage and Stability</b>	Store powder at 4°C protected from light. Following reconstitution, store aliquots at -20°C. Stock solutions are stable for 6 months when stored as directed.
<b>Quality Control</b>	The purity of SB431542 was determined by HPLC analysis. The accurate mass was determined by mass spectrometry. Cellular toxicity of SB431542 was tested on mouse embryonic stem cells.
<b>References</b>	<ol style="list-style-type: none"> <li>Laping, N.J., Grygielko, E., Mathur, A., Butter, S., Bomberger, J., Tweed, C., Martin, W., Fornwald, J., Lehr, R., Harling, J., Gaster, L., Callahan, J.F., and Olson, B.A. (2002) Inhibition of transforming growth factor (TGF)-beta1-induced extracellular matrix with a novel inhibitor of the TGF-beta type I receptor kinase activity: SB-431542. <i>Mol Pharmacol</i> 62: 58-64.</li> <li>Inman, G.J., Nicolás, F.J., Callahan, J.F., Harling, J.D., Gaster, L.M., Reith, A.D., Laping, N.J., and Hill, C.S. (2002) SB-431542 is a potent and specific inhibitor of transforming growth factor-beta superfamily type I activin receptor-like kinase (ALK) receptors ALK4, ALK5, and ALK7. <i>Mol Pharmacol</i> 62: 65-74.</li> <li>Watabe, T., Nishihara, A., Mishima, K., Yamashita, J., Shimizu, K., Miyazawa, K., Nishikawa, S., and Miyazono, K. (2003) TGF-beta receptor kinase inhibitor enhances growth and integrity of embryonic stem cell-derived endothelial cells. <i>J Cell Biol.</i> 163: 1303-1311.</li> </ol>

For research use only. Not for use in diagnostic procedures.

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