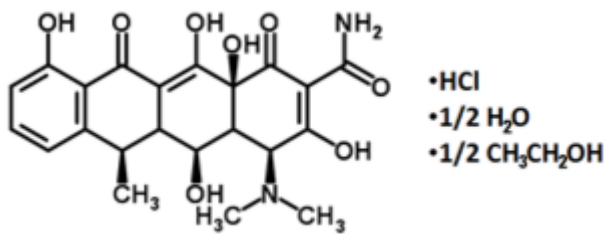


# PRODUCT SPECIFICATIONS SHEET:

## Stemolecule™ Doxycycline hyclate

<b>Product Name</b>	Stemolecule™ Doxycycline Hyclate
<b>Description</b>	Stemolecule Doxycycline hyclate (dox) is a broad spectrum antibiotic derivative of tetracycline and an inhibitor of matrix metalloproteinases <sup>1</sup> . Tetracycline-controlled transcriptional activation is a method of inducible expression whereby transcription is reversibly turned on or off in the presence of tetracycline or one of its derivatives such as dox <sup>2</sup> . For this type of reprogramming method, dox-inducible lentiviral reagents are used to induce the expression of virally transduced genes and generate induced pluripotent stem (iPS) cells from somatic cells <sup>3-7</sup> .
<b>Catalog Number</b>	04-0016
<b>Size</b>	10 mg
<b>Alternate Names</b>	(2Z,4S,5S,6R,12aS)-2-[amino(hydroxy)methylidene]-4-(dimethylamino)-5,10,11,12a-tetrahydroxy-6-methyl-4a,5,5a,6-tetrahydro-4H-tetracene-1,3,12-trione; ethanol; hydrochloride
<b>Chemical Formula</b>	$C_{22}H_{24}N_2O_8 \cdot HCl \cdot \frac{1}{2}(H_2O) \cdot \frac{1}{2}(C_2H_6O)$
<b>Structure</b>	
<b>Molecular Weight</b>	512.94
<b>CAS Number</b>	24390-14-5
<b>Purity</b>	>98% pure by TLC
<b>Formulation</b>	Yellow to yellow with a green cast powder



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<b>Solubility</b>	For a 10 mM concentrated stock solution of Doxycycline hyclate, reconstitute the compound by adding 1.95 mL of water to the entire contents of the vial.
<b>Storage and Stability</b>	Store powder at 4 °C or -20 °C, protected from light.  Information about the stability of Stemolecules in solution is largely not available. As a general guideline, we recommend that stock solution be freshly made and stored in aliquots at -20 °C. The effect of storage of stock solutions should be verified for each application.
<b>Quality Control</b>	The purity of Doxycycline hyclate was determined by TLC analysis. The accurate mass was determined by mass spectrometry. The structure was verified by NMR spectroscopy. Cellular toxicity was tested on mouse embryonic stem cells.
<b>Regulatory Disclaimer</b>	For research use only. Not for use in therapeutic or diagnostic procedures.
<b>References</b>	<ol style="list-style-type: none"><li>1. Burgraff, D., Trinkl, A., Dichgans, M., and Hamann, G.F. (2007) Doxycycline inhibits MMPs via modulation of plasminogen activators in focal cerebral ischemia. <i>Neurobiol Dis</i> 25: 506-513.</li><li>2. Bujard, H., and Gossen, M. (1992) Tight control of gene expression in mammalian cells by tetracycline-responsive promoters. <i>Proc Natl Acad Sci</i> 89: 5547-5551.</li><li>3. Brambrink, T., Foreman, R., Welstead, G.G., Lengner, C.J., Wernig, M., Suh, H., and Jaenisch, R. (2008) Sequential expression of pluripotency markers during direct reprogramming of mouse somatic cells. <i>Cell Stem Cell</i> 2: 151- 159.</li><li>4. Wernig, M., Lengner, C.J., Hanna, J., Lodato, M.A., Steine, E., Foreman, R., Staerk, J., Markoulaki, S., and Jaenisch, R. (2008) A drug-inducible transgenic system for direct reprogramming of multiple somatic cell types. <i>Nat Biotechnol</i> 26: 916-924.</li><li>5. Hockemeyer, D., Soldner, F., Cook, E.G., Gao, Q., Mitalipova, M., and Jaenisch R. (2008) A drug-inducible system for direct reprogramming of human somatic cells to pluripotency. <i>Cell Stem Cell</i> 3: 346-353.</li><li>6. Welstead, G.G., Brambrink, T., and Jaenisch, R. (2008) Generating iPS cells from MEFS through forced expression of Sox-2, Oct-4, c-Myc, and Klf4. <i>J Vis Exp</i> 7: 734.</li><li>7. Markoulaki, S., Hanna, J., Beard, C., Carey, B.W., Cheng, A.W., Lengner, C.J., Dausman, J.A., Fu, D., Gao, Q., Wu, S., Cassady, J.P., and Jaenisch, R. (2009) Transgenic mice with defined combinations of drug-inducible reprogramming factors. <i>Nat Biotechnol</i> 27: 169-171</li></ol>

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