



## CERTIFICATE OF ANALYSIS

<b>PRODUCT:</b>	<b>Atpenin A5</b> Mitochondrial complex II inhibitor; [3-[(2 <i>S</i> ,4 <i>S</i> ,5 <i>R</i> )-5,6-Dichloro-2,4-dimethyl-1-oxohexyl]-4-hydroxy-5,6-dimethoxy-2(1 <i>H</i> )-pyridinone], C <sub>15</sub> H <sub>21</sub> NO <sub>5</sub> Cl <sub>2</sub> , M.W. 366.2, CAS# [119509-24-9]
<b>PRODUCT NUMBER:</b>	A-1438
<b>LOT NUMBER:</b>	C1207
<b>APPEARANCE:</b>	White powder
<b>SOLUBILITY:</b>	Soluble in acetone, acetonitrile, chloroform, ethyl acetate, DMSO, methanol or ethanol. Insoluble in water or hexane.
<b>PURITY (HPLC):</b>	96.1%
<b>SOURCE/HOST:</b>	Synthetic. Originally isolated from <i>Penicillium</i> sp. strain FO-125.
<b>DESCRIPTION:</b>	The IC <sub>50</sub> value against bovine heart complex II is 3.6nM (which is ~300-fold lower than the IC <sub>50</sub> value of carboxin (1.1μM)). It also inhibits fumarate reductase of <i>Ascaris suum</i> (IC <sub>50</sub> = 12nM). Inhibition of <i>E. coli</i> succinate dehydrogenase is less potent (IC <sub>50</sub> = 5μM). By co-crystallization studies of atpenin A5 and complex II, the binding site of atpenin A5 was found to be the quinone-binding site of complex II. Additionally, atpenin A5 has been shown to have a protective action against ischemia-reperfusion via the activation of mitochondrial K <sub>ATP</sub> channels.
<b>STORAGE &amp; HANDLING:</b>	Store short term at +4°C and store long term at -20°C. <b><u>WARNING: TOXIC!</u></b> CAUTION: For laboratory research & scientific manufacturing use only. Not for human or drug use. The pharmacological and toxicological properties of this product have not been fully investigated. Use caution when handling. Do not use this compound if you are not fully trained or are unaware of the hazards involved.

Verified: DD