

Data Sheet

Product Information

Catalog Number	BP13799
Product Name	Delcasertib
Description	Delcasertib is a potent and selective inhibitor of δ -protein kinase C (δ PKC).
In vitro	Delcasertib is composed of a selective δ -protein kinase C (δ PKC) inhibitor peptide derived from the δ V1-1 portion of δ PKC (termed "cargo peptide"), conjugated reversibly to the cell-penetrating peptide 11-amino acid, arginine-rich sequence of the HIV type 1 transactivator protein (TAT47-57; termed "carrier peptide") via a disulfide bond.
In vivo	KAI-9803 ameliorates pathological conditions in acute myocardial infarction and reduce pain via specific modulation of membrane-translocation of PKC delta or epsilon. Delcasertib has an acceptable safety and tolerability profile when delivered via intracoronary injection during primary percutaneous coronary intervention for ST-segment elevation myocardial infarction. Delcasertib administration at the end of ischemia has been found to reduce cardiac damage caused by ischemia-reperfusion in a rat model of acute myocardial infarction. 14 C-KAI-9803 is rapidly delivered to many tissues, including the heart ($1.21 \mu\text{g eq/g}$ tissue), while being quickly cleared from the systemic circulation. The distribution of Delcasertib to tissues such as the liver, kidney, and heart is facilitated by the reversible conjugation to TAT47-57.
Synonyms	BMS-875944, KAI-9803
CAS No.	949100-39-4
Chemical Formula	C ₁₂₀ H ₁₉₉ N ₄₅ O ₃₄ S ₂
Molecular Weight	2880.31

Solubility	DMSO: 99 mg/mL(34.37 mM)
Storage	Powder: -20°C for 2 years In solvent: -80°C for 1 year
Chemical Structure OR Tested Image	<p>Sequence 1:Cys-Tyr-Gly-Arg-Lys-Lys-Arg-Arg-Gln-Arg-Arg-Arg; Sequence 1':Ser-Phe-Asn-Ser-Tyr-Glu-Leu-Gly-Ser-Leu (Disulfide bridge:Cys₁-Cys₁)</p>

Purdue Bioscience Inc.

750 50th St, Brooklyn, NY 11220, USA

<https://www.purduebio.com>

1-877.618.7311

info@purduebio.com

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