

Product Name: RWJ 56110

Catalog No.: 2614

Batch No.: 2

IUPAC Name: (*α,S*)-*N*-[*(1S)*-3-Amino-1-[[*(phenylmethyl)*amino]carbonyl]propyl]-*α*-[[[1-[(2,6-dichlorophenyl)methyl]-3-(1-pyrrolidinylmethyl)-1*H*-indol-6-yl]amino]carbonyl]amino]-3,4-difluoro-benzenepropanamide

1. PHYSICAL AND CHEMICAL PROPERTIES

Batch Molecular Formula: C₄₁H₄₃Cl₂F₂N₇O₃·2HCl·1½H₂O

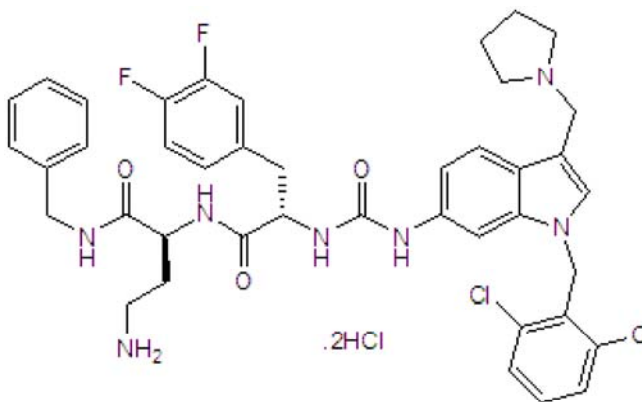
Batch Molecular Weight: 890.67

Physical Appearance: Grey solid

Solubility: water to 25 mM
DMSO to 100 mM

Storage: Store at -20°C

Batch Molecular Structure:



2. ANALYTICAL DATA

TLC: R_f = 0.19 (Chloroform:Methanol:Ammonia soln. [9:1:0.1])

HPLC: Shows 98.1% purity

¹H NMR: Consistent with structure

Mass Spectrum: Consistent with structure

Microanalysis:

	Carbon	Hydrogen	Nitrogen
Theoretical	55.29	5.43	11.01
Found	55.11	5.15	10.85

Caution - Not Fully Tested • Research Use Only • Not For Human or Veterinary Use

Product Name: RWJ 56110

Catalog No.: 2614

Batch No.: 2

IUPAC Name: (α S)-N-[(1S)-3-Amino-1-[[[(phenylmethyl)amino]carbonyl]propyl]- α -[[[1-[(2,6-dichlorophenyl)methyl]-3-(1-pyrrolidinylmethyl)-1H-indol-6-yl]amino]carbonyl]amino]-3,4-difluoro-benzenepropanamide

Description:

Selective protease-activated receptor-1 (PAR₁) antagonist; displays no activity at PAR₂, PAR₃, or PAR₄ subtypes. Blocks thrombin-induced platelet aggregation and activation of MAPK in HUVECs. Also inhibits angiogenesis in a chick embryo angiogenesis model in vivo.

Physical and Chemical Properties:

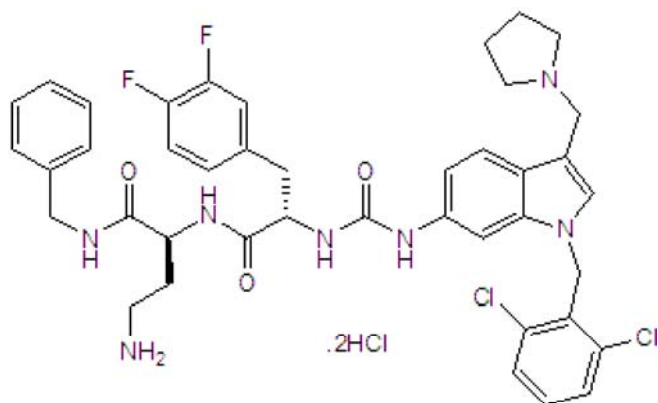
Batch Molecular Formula: C₄₁H₄₃Cl₂F₂N₇O₃.2HCl.1½H₂O

Batch Molecular Weight: 890.67

Physical Appearance: Grey solid

Minimum Purity: >96%

Batch Molecular Structure:



Storage: Store at -20°C

Solubility & Usage Info:

water to 25 mM

DMSO to 100 mM

This product is supplied as a lyophilized solid and may be very hard to visualize. Solutions should be made by adding solvent directly to the vial. The vial should then be vortexed vigorously to ensure the product has completely dissolved.

Stability and Solubility Advice:

Some solutions can be difficult to obtain and can be encouraged by rapid stirring, sonication or gentle warming (in a 45-60°C water bath).

Information concerning product stability, particularly in solution, has rarely been reported and in most cases we can only offer a general guide. Our standard recommendations are:

SOLIDS: Provided storage is as stated on the product label and the vial is kept tightly sealed, the product can be stored for up to 6 months from date of receipt.

SOLUTIONS: We recommend that stock solutions, once prepared, are stored aliquoted in tightly sealed vials at -20°C or below and used within 1 month. Wherever possible solutions should be made up and used on the same day.

Other Information:

INFORMATION FOR CUSTOMERS IN THE UK ONLY

This product is a Schedule 1 Home Office controlled substance and customers in the UK are required to hold the relevant licence or be exempt from restrictions in order to purchase and possess this material.

References:

Zania et al (2006) Blockade of angiogenesis by small molecule antagonists to protease-activated receptor-1: association with endothelial cell growth suppression and induction of apoptosis. *J.Pharmacol.Exp.Ther.* **318** 246. PMID: 16595737.

Maryanoff et al (2003) Discovery of potent peptide-mimetic antagonists for the human thrombin receptor, protease-activated receptor-1 (PAR-1). *Curr.Med.Chem.Cardiovasc.Hematol.Agents* **1** 13. PMID: 15317288.

Andrade-Gordon et al (1999) Design, synthesis and biological characterization of a peptide-mimetic antagonist for a tethered-ligand receptor. *Proc.Natl.Acad.Sci.USA* **96** 12257. PMID: 10535908.

Caution - Not Fully Tested • Research Use Only • Not For Human or Veterinary Use

bio-techne.com

info@bio-techne.com

techsupport@bio-techne.com

North America

Tel: (800) 343 7475

China

info.cn@bio-techne.com

Tel: +86 (21) 52380373

Europe Middle East Africa

Tel: +44 (0)1235 529449

Rest of World

www.tocris.com/distributors

Tel:+1 612 379 2956