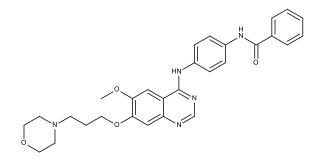


Catalog # 10-1037 ZM-447439

N-[4-[[6-Methoxy-7-[3-(4-morpholinyl)propoxy]-4-quinazolinyl]amino]phenyl]benzamide CAS# 331771-20-1 Lot # X101911



ZM-447439 is a selective ATP-competitive inhibitor of Aurora kinase B¹ (IC₅₀ = 50 nM, Aur A IC₅₀ = 1 μ M, Aur C IC₅₀ = 250 nM at physiological ATP concentrations², Ref.1 lists Aur A IC₅₀ = 110nM and Aur B IC₅₀ = 113nM). It did not significantly inhibit a panel of 14 other kinases including mitotic kinases CDK1 and PLK1. It prevents chromosome alignment, spindle checkpoint function and cytokinesis.^{1,2}

- 1) Ditchfield et al (2003) Aurora B couples chromosome alignment with anaphase by targeting BubR1, Mad2, and Cenp-E to kinetochores. J.Cell Biol. **161** 267
- 2) Girdler et al (2006) Validating Aurora B as an anti-cancer drug target. J.Cell Sci. 119 3664

PHYSICAL DATA

Molecular Weight:	513.59
Molecular Formula:	C ₂₉ H ₃₁ N ₅ O ₄
Purity:	98% (TLC: 10% Methanol/methylene chloride; Rf = 0.30)
Solubility:	DMSO (at least 50 mg/ml); ethanol (at least 25 mg/mL)
Physical Description:	White solid
Storage and Stability:	Store as supplied at room temperature for up to one year from the date of purchase. Solutions in DMSO or ethanol may be stored at -20°C for up to 3 months

Materials provided by Focus Biomolecules are for laboratory research use only and are not intended for human or veterinary applications.

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