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Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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PRODUCT INFORMATION



FRAX1036

Item No. 20711

CAS Registry No.: 1432908-05-8

Formal Name: 6-[2-chloro-4-(6-methyl-2-pyrazinyl)phenyl]-8-ethyl-2-[[2-(1-methyl-4-piperidyl)ethyl]amino]-pyrido[2,3-d]pyrimidin-7(8H)-one

MF: C₂₈H₃₂ClN₇O

FW: 518.1

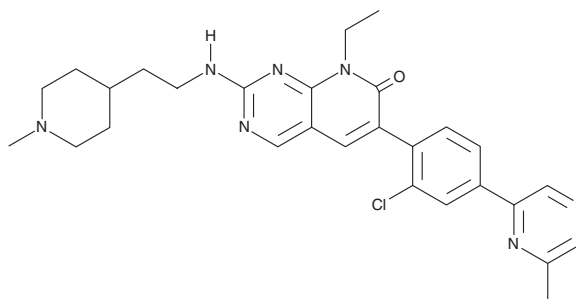
Purity: ≥98%

UV/Vis.: λ_{max}: 211, 299, 308, 355 nm

Supplied as: A crystalline solid

Storage: -20°C

Stability: As supplied, 2 years from the QC date provided on the Certificate of Analysis, when stored properly



Laboratory Procedures

FRAX1036 is supplied as a crystalline solid. A stock solution may be made by dissolving the FRAX1036 in the solvent of choice. FRAX1036 is soluble in organic solvents such as ethanol and dimethyl formamide (DMF), which should be purged with an inert gas. The solubility of FRAX1036 in these solvents is approximately 0.33 and 1 mg/ml, respectively.

FRAX1036 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, FRAX1036 should first be dissolved in DMF and then diluted with the aqueous buffer of choice. FRAX1036 has a solubility of approximately 0.25 mg/ml in a 1:3 solution of DMF:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

FRAX1036 is a selective inhibitor of p21-activated kinase 1 (PAK1), a serine/threonine kinase downstream of Rac1 and Cdc42 that is involved in tumorigenesis. It can induce apoptosis in breast cancer cells and has been shown to potentiate the activity of the microtubule stabilizing agent, docetaxel (Item No. 11637).¹ Disruption of PAK1 via FRAX1036 has been used to inhibit oncogenic KRAS signaling in non-small cell lung cancer cells.²

References

1. Ong, C.C., Gierke, S., Pitt, C., *et al.* Small molecule inhibition of group I p21-activated kinases in breast cancer induces apoptosis and potentiates the activity of microtubule stabilizing agents. *Breast Cancer Res.* **17**(59) (2015).
2. Mortazavi, F., Lu, J., Phan, R., *et al.* Significance of KRAS/PAK1/Crk pathway in non-small cell lung cancer oncogenesis. *BMC Cancer* **15**(381) (2015).

WARNING

THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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