PRODUCT INFORMATION



LY2606368

Item No. 21490

CAS Registry No.: 1234015-52-1

Formal Name: 5-[[5-[2-(3-aminopropoxy)-6-

methoxyphenyl]-1H-pyrazol-3-yl]

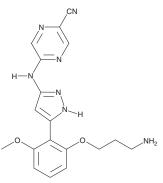
amino]-2-pyrazinecarbonitrile

Synonym: Prexasertib MF: $C_{18}H_{19}N_7O_2$ FW: 365.4 **Purity:**

UV/Vis.: λ_{max} : 214, 295 nm A crystalline solid Supplied as:

-20°C Storage: Stability: ≥4 years

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.



Description

LY2606368 is a checkpoint kinase 1 (Chk1) inhibitor with a K_i value of 0.9 nM against the purified target and IC_{50} values of <13 nM in a viability study of multiple colorectal cancer cell lines. ^{1,2} Inhibition of Chk1 causes double-strand DNA breakage in cells leading to an excessive cell damage burden and subsequent cell death. 1,2 In vitro, LY2606368 inhibits the doxorubicin-activated G_2/M checkpoint in p53-deficient HeLa cells with an EC_{50} value of 9 nM. LY2606368, at 25 nM, also significantly induces apoptosis and inhibits colony formation in AGS and MKN1 gastric cancer cells.3

References

- 1. King, C., Diaz, H.B., McNeely, S., et al. LY2606368 causes replication catastrophe and antitumor effects through CHK1-dependent mechanisms. Mol. Cancer Ther. 14(9), 2004-2013 (2015).
- 2. Manic, G., Signore, M., Sistigu, A., et al. CHK1-targeted therapy to deplete DNA replication-stressed, p53-deficient, hyperdiploid colorectal cancer stem cells. Gut 67(5), 903-917 (2017).
- 3. Yin, Y., Shen, Q., Zhang, P., et al. Chk1 inhibition potentiates the therapeutic efficacy of PARP inhibitor BMN673 in gastric cancer. Am. J. Cancer Res. 7(3), 473-483 (2017).

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

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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