



LKT Laboratories, Inc.

LY-2603618

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## Product Information

**Product ID** L9704

**CAS No.** 911222-45-2

**Chemical Name** 1-[5-bromo-4-methyl-2-[[[(2S)-morpholin-2-yl]methoxy]phenyl]-3-(5-methylpyrazin-2-yl)urea

**Synonym** Rabusertib

**Formula** C<sub>18</sub>H<sub>22</sub>BrN<sub>5</sub>O<sub>3</sub>

**Formula Wt.** 436.31

**Melting Point**

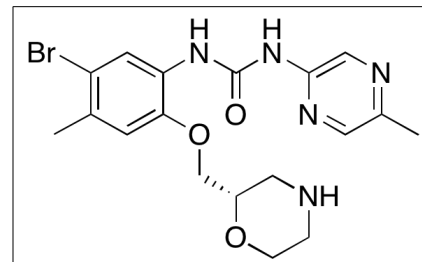
**Purity** ≥98%

**Solubility** 13mg/mL in DMSO, insoluble in water and ethanol

**Store Temp** -20°C

**Ship Temp** Ambient

**Description** LY-2603618 is an inhibitor of checkpoint kinase (CHK) 1 that exhibits anticancer chemotherapeutic activity. In various animal models with tumor xenografts, this compound inhibits the S phase DNA damage checkpoint and increases DNA damage. In lung cancer cells, LY-2603618 induces G2/M phase cell cycle arrest and autophagy. LY-2603618 also enhances activity of other co-administered chemotherapeutics such as gemcitabine.



## Pricing and Availability

**Bulk quantities available upon request**

Product ID	Size	List Price
L9704	1 mg	\$65.00
L9704	5 mg	\$95.00
L9704	10 mg	\$175.00
L9704	25 mg	\$305.00

**References** Barnard D, Diaz HB, Burke T, et al. LY2603618, a selective CHK1 inhibitor, enhances the anti-tumor effect of gemcitabine in xenograft tumor models. *Invest New Drugs*. 2015 Nov 27. [Epub ahead of print]. PMID: 26612134.

Wang FZ, Fei HR, Cui YJ, et al. The checkpoint 1 kinase inhibitor LY2603618 induces cell cycle arrest, DNA damage response and autophagy in cancer cells. *Apoptosis*. 2014 Sep;19(9):1389-98. PMID: 24928205.

King C, Diaz H, Barnard D, et al. Characterization and preclinical development of LY2603618: a selective and potent Chk1 inhibitor. *Invest New Drugs*. 2014 Apr;32(2):213-26. PMID: 24114124.

**Caution:** This product is intended for laboratory and research use only. It is not for human or drug use.