



# SZABO SCANDIC

Part of Europa Biosite

## Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

Weitere Information auf den folgenden Seiten!  
See the following pages for more information!



### Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

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# PRODUCT INFORMATION

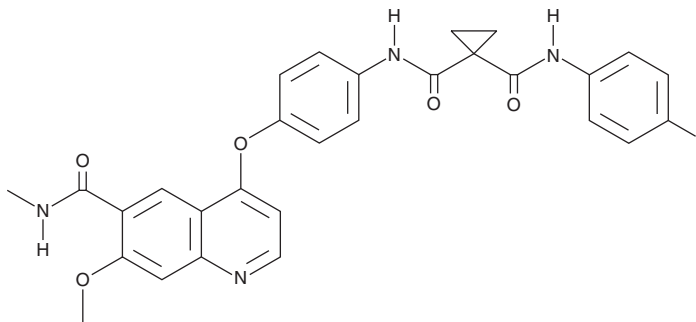


## XL092

Item No. 37749

**CAS Registry No.:** 2367004-54-2  
**Formal Name:** N-(4-fluorophenyl)-N'-[4-[[7-methoxy-6-[(methylamino)carbonyl]-4-quinolinyl]oxy]phenyl]-1,1-cyclopropanedicarboxamide

**Synonym:** Zanzalintinib  
**MF:** C<sub>29</sub>H<sub>25</sub>FN<sub>4</sub>O<sub>5</sub>  
**FW:** 528.5  
**Purity:** ≥98%  
**Supplied as:** A solid  
**Storage:** -20°C  
**Stability:** ≥4 years



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

### Laboratory Procedures

XL092 is supplied as a solid. A stock solution may be made by dissolving the XL092 in the solvent of choice, which should be purged with an inert gas. XL092 is soluble in DMSO.

### Description

XL092 is a multi-kinase inhibitor.<sup>1</sup> It selectively inhibits the receptor tyrosine kinases (RTKs) MET, VEGFR2, Axl, and Mer (IC<sub>50</sub>s = 3, 15, 5.8, and 0.6 nM, respectively) over serine/threonine kinases for which it has no activity, but does inhibit 28 other RTKs with IC<sub>50</sub> values ranging from 3 to 54 nM, as well as additional kinases by greater than 70% at 1 μM. XL092 inhibits the proliferation of SNU-5 human gastric carcinoma cells (IC<sub>50</sub> = 98.9 nM), which highly express MET, and human umbilical vein endothelial cells (HUVECs; IC<sub>50</sub> = 10.4 nM). It reduces tumor volume and intratumoral MET phosphorylation in NCI H441 human lung cancer and SNU-5 mouse xenograft models when administered at doses of 3 and 10 mg/kg. XL092 (30 mg/kg) increases the number of peripheral B cells and CD4<sup>+</sup> T cells, and decreases the number of peripheral myeloid cells, in an MC-38 syngeneic mouse model of colon carcinoma, indicating a conversion to an immune-permissive tumor microenvironment.

### Reference

1. Hsu, J., Chong, C., Serrill, J., *et al.* Preclinical characterization of XL092, a novel receptor tyrosine kinase inhibitor of MET, VEGFR2, AXL, and MER. *Mol. Cancer Ther.* **22(2)**, 179-191 (2023).

#### 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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