



# 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

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

[mail@szabo-scandic.com](mailto:mail@szabo-scandic.com)

[www.szabo-scandic.com](http://www.szabo-scandic.com)

[linkedin.com/company/szaboscandic](https://www.linkedin.com/company/szaboscandic) 

# PRODUCT INFORMATION

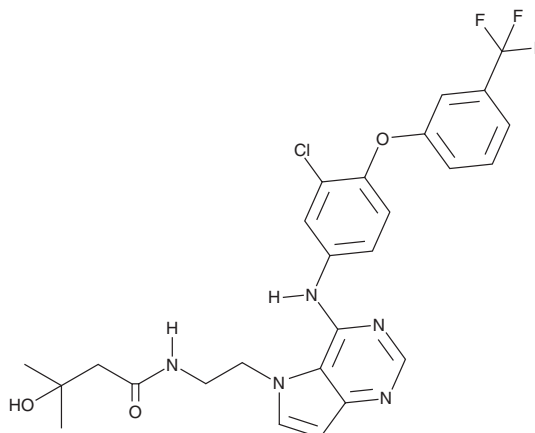


## TAK-285

Item No. 29157

**CAS Registry No.:** 871026-44-7  
**Formal Name:** N-[2-[4-[[3-chloro-4-[3-(trifluoromethyl)phenoxy]phenyl]amino]-5H-pyrrolo[3,2-d]pyrimidin-5-yl]ethyl]-3-hydroxy-3-methyl-butanamide

**MF:** C<sub>26</sub>H<sub>25</sub>ClF<sub>3</sub>N<sub>5</sub>O<sub>3</sub>  
**FW:** 548.0  
**Purity:** ≥95%  
**UV/Vis.:** λ<sub>max</sub>: 215, 306 nm  
**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

TAK-285 is supplied as a solid. A stock solution may be made by dissolving the TAK-285 in the solvent of choice, which should be purged with an inert gas. TAK-285 is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of TAK-285 in ethanol is approximately 10 mg/ml and approximately 30 mg/ml in DMSO and DMF.

TAK-285 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, TAK-285 should first be dissolved in DMSO and then diluted with the aqueous buffer of choice. TAK-285 has a solubility of approximately 0.25 mg/ml in a 1:3 solution of DMSO:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

### Description

TAK-285 is a dual inhibitor of EGFR and HER2 (IC<sub>50</sub>s = 23 and 17 nM, respectively).<sup>1</sup> It is selective for EGFR and HER2 over a panel of 36 kinases (IC<sub>50</sub>s = 250->10,000 nM). TAK-285 inhibits the proliferation of BT474 breast cancer cells (GI<sub>50</sub> = 17 nM) as well as induces tumor regression in a 4-1st gastric adenocarcinoma rat xenograft model when administered at doses of 25 and 50 mg/kg.

### Reference

1. Ishikawa, T., Seto, M., Banno, H., *et al.* Design and synthesis of novel human epidermal growth factor receptor 2 (HER2)/epidermal growth factor receptor (EGFR) dual inhibitors bearing a pyrrolo[3,2-d]pyrimidine scaffold. *J. Med. Chem.* **54**(23), 8030-8050 (2011).

#### 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.

#### WARRANTY AND LIMITATION OF REMEDY

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#### CAYMAN CHEMICAL

1180 EAST ELLSWORTH RD  
ANN ARBOR, MI 48108 · USA

**PHONE:** [800] 364-9897  
[734] 971-3335

**FAX:** [734] 971-3640

CUSTSERV@CAYMANCHEM.COM  
WWW.CAYMANCHEM.COM