



# SZABO SCANDIC

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## Produktinformation



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### Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

### SZABO-SCANDIC HandelsgmbH

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# PRODUCT INFORMATION



**RO4987655**

Item No. 19488

**CAS Registry No.:** 874101-00-5

**Formal Name:** 3,4-difluoro-2-[(2-fluoro-4-iodophenyl)amino]-N-(2-hydroxyethoxy)-5-[(tetrahydro-3-oxo-2H-1,2-oxazin-2-yl)methyl]-benzamide

**Synonym:** CH4987655

**MF:** C<sub>20</sub>H<sub>19</sub>F<sub>3</sub>IN<sub>3</sub>O<sub>5</sub>

**FW:** 565.3

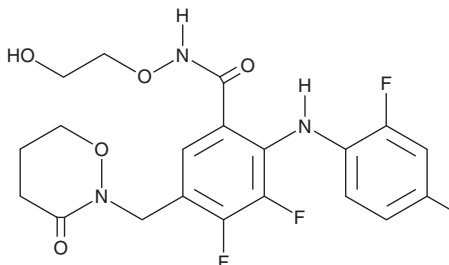
**Purity:** ≥98%

**UV/Vis.:** λ<sub>max</sub>: 280 nm

**Supplied as:** A crystalline solid

**Storage:** -20°C

**Stability:** ≥2 years



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

## Laboratory Procedures

RO4987655 is supplied as a crystalline solid. A stock solution may be made by dissolving the RO4987655 in the solvent of choice. RO4987655 is soluble in organic solvents such as DMSO and dimethyl formamide (DMF), which should be purged with an inert gas. The solubility of RO4987655 in these solvents is approximately 10 and 15 mg/ml, respectively.

RO4987655 is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, RO4987655 should first be dissolved in DMF and then diluted with the aqueous buffer of choice. RO4987655 has a solubility of approximately 0.16 mg/ml in a 1:5 solution of DMF:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

## Description

RO4987655 is a MEK inhibitor (IC<sub>50</sub> = 5 nM).<sup>1</sup> It is selective for MEK over 400 kinases at 10 μM. It inhibits proliferation of COLO 205, HT-29, QG56, MIA PaCa-2, and C32 cells with IC<sub>50</sub> values of 0.86, 1.7, 9.5, 3.3, and 8.4 nM, respectively. RO4987655 reduces tumor growth in a variety of mouse xenograft models and inhibits the phosphorylation of ERK in tumor tissue in an HT-29 mouse xenograft model when administered at a dose of 6.25 mg/kg per day. It acts synergistically with the mTOR inhibitor everolimus (Item No. 11597) to reduce tumor volume in an HCT116 mouse xenograft model.

## Reference

1. Isshiki, Y., Kohchi, Y., Iikura, H., *et al.* Design and synthesis of novel allosteric MEK inhibitor CH4987655 as an orally available anticancer agent. *Bioorg. Med. Chem. Lett.* **21**(6), 1795-1801 (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.

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