



# 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) 

## Narnatumab

Cat. No.:	HY-P99375
CAS No.:	1188275-92-4
Target:	Others
Pathway:	Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.

### BIOLOGICAL ACTIVITY

<b>Description</b>	Narnatumab (IMC-RON8) is a neutralizing human monoclonal antibody that blocks RON binding to its ligand, macrophage-stimulating protein (MSP), with a $K_d$ of 32 pM. Narnatumab can be used for the research of cancer <sup>[1]</sup> .								
<b>IC<sub>50</sub> &amp; Target</b>	RON(MSPR) <sup>[1]</sup>								
<b>In Vitro</b>	<p>Narnatumab (100 nM; 24 h) inhibits MSP-induced migration of human lung and breast cancer cell lines<sup>[1]</sup>.</p> <p>Narnatumab inhibits the MSP-induced mitogenic response of a pancreatic cancer cell line<sup>[1]</sup>.</p> <p>Narnatumab (0.01-100 nM; pretreatment for 1 h) blocks ligand-induced receptor phosphorylation and downstream signaling molecules phosphorylation in RON-expressing tumor cells and in a RON-transfected cell line<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p>								
<b>In Vivo</b>	<p>Narnatumab (2-60 mg/kg; i.p. every 3 days) shows antitumor activity in non-small cell lung cancer (NSCLC) and bladder cancer models in athymic mice<sup>[1]</sup>.</p> <p>Narnatumab exhibits terminal half-life (<math>t_{1/2}</math>=5.2 d) and achieves antitumor effects at a steady-state plasma trough level of approximately 140 µg/mL in mice<sup>[1]</sup>.</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <table border="1"> <tr> <td>Animal Model:</td> <td>Female athymic nu/nu mice (6-8 weeks) were injected NCI-H292 and BFTC-905 cells<sup>[1]</sup></td> </tr> <tr> <td>Dosage:</td> <td>2, 20, 60 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>I.p. every 3 days for 36 and 18 days</td> </tr> <tr> <td>Result:</td> <td>Inhibited tumor growth in a dose-dependent manner.</td> </tr> </table>	Animal Model:	Female athymic nu/nu mice (6-8 weeks) were injected NCI-H292 and BFTC-905 cells <sup>[1]</sup>	Dosage:	2, 20, 60 mg/kg	Administration:	I.p. every 3 days for 36 and 18 days	Result:	Inhibited tumor growth in a dose-dependent manner.
Animal Model:	Female athymic nu/nu mice (6-8 weeks) were injected NCI-H292 and BFTC-905 cells <sup>[1]</sup>								
Dosage:	2, 20, 60 mg/kg								
Administration:	I.p. every 3 days for 36 and 18 days								
Result:	Inhibited tumor growth in a dose-dependent manner.								

### REFERENCES

[1]. LoRusso PM, et, al. Phase 1 study of narnatumab, an anti-RON receptor monoclonal antibody, in patients with advanced solid tumors. Invest New Drugs. 2017 Aug;35(4):442-450.

---

**Caution: Product has not been fully validated for medical applications. For research use only.**

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: [tech@MedChemExpress.com](mailto:tech@MedChemExpress.com)

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA