



# 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) 

## Cemiplimab

Cat. No.:	HY-P99203
CAS No.:	1801342-60-8
Target:	PD-1/PD-L1
Pathway:	Immunology/Inflammation
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.

### BIOLOGICAL ACTIVITY

<b>Description</b>	Cemiplimab (Anti-Human PD-1) is a high-affinity programmed death receptor-1 (PD-1) monoclonal IgG4 antibody that blocks PD-1/PD-L1-mediated T-cell suppression. Cemiplimab is commonly used in squamous cell skin cancer research <sup>[1][2]</sup> .								
<b>In Vivo</b>	<p>Cemiplimab (i.p., 10 mg/kg, five injections within 2 weeks) can increased the proportion of effector T cells in tumors and dLNs to reduce tumor growth by combined immunotherapy with REGN3767 in MC38.Ova tumor mice model<sup>[2]</sup>. 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 human PD-1×LAG-3 knockin mice (8-10 weeks) with MC38.Ova cells<sup>[2]</sup></td> </tr> <tr> <td>Dosage:</td> <td>10 mg/kg</td> </tr> <tr> <td>Administration:</td> <td>Intraperitoneal injection; five injections within 2 weeks</td> </tr> <tr> <td>Result:</td> <td>Partially inhibited the growth of MC38.Ova tumors and prolonged the survival of mice by administering alone. Significantly increased the proportion of CD4<sup>+</sup> TILs in IFN<math>\gamma</math> and TNF<math>\alpha</math> by combined immunotherapy with REGN3767.</td> </tr> </table>	Animal Model:	Female human PD-1×LAG-3 knockin mice (8-10 weeks) with MC38.Ova cells <sup>[2]</sup>	Dosage:	10 mg/kg	Administration:	Intraperitoneal injection; five injections within 2 weeks	Result:	Partially inhibited the growth of MC38.Ova tumors and prolonged the survival of mice by administering alone. Significantly increased the proportion of CD4 <sup>+</sup> TILs in IFN $\gamma$ and TNF $\alpha$ by combined immunotherapy with REGN3767.
Animal Model:	Female human PD-1×LAG-3 knockin mice (8-10 weeks) with MC38.Ova cells <sup>[2]</sup>								
Dosage:	10 mg/kg								
Administration:	Intraperitoneal injection; five injections within 2 weeks								
Result:	Partially inhibited the growth of MC38.Ova tumors and prolonged the survival of mice by administering alone. Significantly increased the proportion of CD4 <sup>+</sup> TILs in IFN $\gamma$ and TNF $\alpha$ by combined immunotherapy with REGN3767.								

### REFERENCES

- [1]. Michael R Migden, et al. PD-1 Blockade with Cemiplimab in Advanced Cutaneous Squamous-Cell Carcinoma. *N Engl J Med*. 2018 Jul 26;379(4):341-351.
- [2]. Elena Burova, et al. Preclinical Development of the Anti-LAG-3 Antibody REGN3767: Characterization and Activity in Combination with the Anti-PD-1 Antibody Cemiplimab in Human PD-1xLAG-3-Knockin Mice. *Mol Cancer Ther*. 2019 Nov;18(11):2051-2062.

**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

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