

Adebrelimab

Cat. No.:	HY-P99422
CAS No.:	2247114-85-6
Target:	PD-1/PD-L1; Apoptosis
Pathway:	Immunology/Inflammation; Apoptosis
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.

BIOLOGICAL ACTIVITY

Description	Adebrelimab (SHR-1316) is a humanized IgG4 monoclonal PD-L1 (PD-1/PD-L1) antibody. Adebrelimab has promising antitumor activity in solid tumors including extensive-stage small-cell lung cancer (SCLC) ^{[1][2]} .																						
In Vitro	<p>Adebrelimab (SHR-1316; 0.06- 1mg/mL; 48 hours) inhibits effects on the cell proliferation, migration, invasion of SK-BR-3 and AU565 cells^[1].</p> <p>Adebrelimab (SHR-1316; 0.1-1mg/mL; 48 hours) downregulated the expression of PD-L1, p-PI3K, p-AKT and upregulates the expression FOXO1^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Cell Viability Assay^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>SK-BR-3 and AU565 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.06 mg/mL, 0.13 mg/mL, 0.25 mg/mL, 0.5 mg/mL, 1 mg/mL</td> </tr> <tr> <td>Incubation Time:</td> <td>48 hours</td> </tr> <tr> <td>Result:</td> <td>The viability of SK-BR-3 and AU565 cells decreased gradually.</td> </tr> </table> <p>Apoptosis Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>SK-BR-3 and AU565 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.1 mg/mL, 1 mg/mL</td> </tr> <tr> <td>Incubation Time:</td> <td>48 hours</td> </tr> <tr> <td>Result:</td> <td>Promoted cell apoptosis.</td> </tr> </table> <p>Cell Migration Assay ^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>SK-BR-3 and AU565 cells</td> </tr> <tr> <td>Concentration:</td> <td>0.1 mg/mL, 1 mg/mL</td> </tr> <tr> <td>Incubation Time:</td> <td>48 hours</td> </tr> </table>	Cell Line:	SK-BR-3 and AU565 cells	Concentration:	0.06 mg/mL, 0.13 mg/mL, 0.25 mg/mL, 0.5 mg/mL, 1 mg/mL	Incubation Time:	48 hours	Result:	The viability of SK-BR-3 and AU565 cells decreased gradually.	Cell Line:	SK-BR-3 and AU565 cells	Concentration:	0.1 mg/mL, 1 mg/mL	Incubation Time:	48 hours	Result:	Promoted cell apoptosis.	Cell Line:	SK-BR-3 and AU565 cells	Concentration:	0.1 mg/mL, 1 mg/mL	Incubation Time:	48 hours
Cell Line:	SK-BR-3 and AU565 cells																						
Concentration:	0.06 mg/mL, 0.13 mg/mL, 0.25 mg/mL, 0.5 mg/mL, 1 mg/mL																						
Incubation Time:	48 hours																						
Result:	The viability of SK-BR-3 and AU565 cells decreased gradually.																						
Cell Line:	SK-BR-3 and AU565 cells																						
Concentration:	0.1 mg/mL, 1 mg/mL																						
Incubation Time:	48 hours																						
Result:	Promoted cell apoptosis.																						
Cell Line:	SK-BR-3 and AU565 cells																						
Concentration:	0.1 mg/mL, 1 mg/mL																						
Incubation Time:	48 hours																						

	Result:	Decreased the migration rate of SK-BR -3 and AU565 cells.
	Western Blot Analysis ^[1]	
	Cell Line:	SK-BR-3 and AU565 cells
	Concentration:	0.1 mg/mL, 1 mg/mL
	Incubation Time:	48 hours
	Result:	Downregulated the expression of PD-L1, p-PI3K, p-AKT and upregulated the expression FOXO1.
In Vivo	Adebrelimab (SHR-1316; 200 µg; three times a week; for 21 days) shows inhibition effects on tumor growth ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	
	Animal Model:	Pathogen free (SPF) BALB/c nude female mice (aged 5-6 weeks) injected with SK-BR-3 cells [1]
	Dosage:	200 µg
	Administration:	Three times a week; for 21 days
	Result:	Significantly inhibited tumor growth.

REFERENCES

[1]. Zhiwei Liu, et al. The Synergistic Effects of Pyrotinib and SHR-1316 on HER2-positive breast cancer. Research Square. May 19th, 2022.

[2]. Wanpu Yan, et al. Adebrelimab (SHR-1316) in combination with chemotherapy as perioperative treatment in patients with resectable stage II-III NSCLC: an open-label, multicenter, phase 1b trial. J Thorac Oncol. 2022 Sep 30;S1556-0864(22)01822-6.

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