Proteins

Ned 19

Cat. No.: HY-103316A CAS No.: 874374-25-1 Molecular Formula: $C_{30}H_{31}FN_{4}O_{3}$ Molecular Weight: 514.59

Calcium Channel Target:

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling

Storage: Powder -20°C 3 years

In solvent

4°C 2 years -80°C 6 months

-20°C 1 month

Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: 125 mg/mL (242.91 mM; Need ultrasonic)

| | Solvent Mass Concentration | 1 mg | 5 mg | 10 mg |
|------------------------------|-------------------------------|-----------|-----------|------------|
| Preparing Stock Solutions | 1 mM | 1.9433 mL | 9.7165 mL | 19.4329 mL |
| | 5 mM | 0.3887 mL | 1.9433 mL | 3.8866 mL |
| | 10 mM | 0.1943 mL | 0.9716 mL | 1.9433 mL |

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

| Description | Ned 19 is a selective membrane-permeant non competitive NAADP antagonist and inhibits NAADP-mediated Ca^{2+} signaling, with an IC_{50} of 65 nM ^[1] . Ned 19 strongly inhibits tumor growth and vascularization as well as lung metastases in mice ^[2] . |
|---------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| IC ₅₀ & Target | NAADP, $Ca_{2+}^{[1]}$ |

Ned 19 (25-100 μ M; 24-72 hours) reduces cell proliferation^[2]. In Vitro

Ned 19 (25-100 μ M; 24-72 hours) reduces markedly the cell number^[2].

Ned 19 (25-100 μ M; 24-72 hours) reduces the S phase percentage and increases of the G0/G1 phase percentage evaluated by cell cycle analysis^[2].

Ned 19 (25-100 μ M; 24-72 hours) induces cell apoptosis a time-dependent manner [2].

Ned 19 (25-100 μM; 24-72 hours) reduces expression of N-cadherin and increases expression of E-cadherin, affecting the cell migratory behavior^[2].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Proliferation Assay^[2]

| | B16 cells | |
|--------------------------------------|-----------------------------------------------------------------------------|--|
| Concentration: | 25, 50, 100 μΜ | |
| Incubation Time: | 24, 48, 72 hours | |
| Result: | Reduced cell proliferation. | |
| Cell Viability Assay ^[2] | | |
| Cell Line: | B16 cells | |
| Concentration: | 25, 50, 100 μΜ | |
| Incubation Time: | 24, 48, 72 hours | |
| Result: | Reduced markedly the cell number. | |
| Cell Cycle Analysis ^[2] | | |
| Cell Line: | B16 cells | |
| Concentration: | 25, 50, 100 μΜ | |
| Incubation Time: | 24, 48, 72 hours | |
| Result: | Reduced the S phase percentage and increased of the G0/G1 phase percentage. | |
| Apoptosis Analysis ^[2] | | |
| Cell Line: | B16 cells | |
| Concentration: | 25, 50, 100 μΜ | |
| Incubation Time: | 24, 48, 72 hours | |
| Result: | Induced cell apoptosis a time-dependent manner. | |
| Western Blot Analysis ^[2] | | |
| Cell Line: | B16 cells | |
| Concentration: | 25, 50, 100 μM | |
| | 24, 48, 72 hours | |
| Incubation Time: | Reduced expression of N-cadherin and increased expression of E-cadherin. | |

In Vivo

| Animal Model: | Adult male C57BL/6 mice ^[2] | |
|-----------------|----------------------------------------|--|
| Dosage: | 5 mg/Kg | |
| Administration: | I.p.; every second day; 4 week | |
| Result: | Impaired severely tumor growth. | |

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CUSTOMER VALIDATION

- Nat Commun. 2023 Jan 14;14(1):226.
- Cell Rep. 2023 Dec 26;42(12):113573.

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REFERENCES

[1]. Rosen D, et al. Analogues of the nicotinic acid adenine dinucleotide phosphate (NAADP) antagonistNed-19 indicate two binding sites on the NAADP receptor. J Biol Chem. 2009 Dec 11;284(50):34930-4.

[2]. Annarita Favia, et al. NAADP-Dependent Ca2+ Signaling Controls Melanoma Progression, Metastatic Dissemination and Neoangiogenesis. Sci Rep. 2016; 6: 18925.

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

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