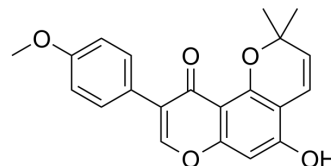


Nrf2/HO-1 activator 1

| | |
|---------------------------|---|
| Cat. No.: | HY-151430 |
| Molecular Formula: | C ₂₁ H ₁₈ O ₅ |
| Molecular Weight: | 350.36 |
| Target: | Keap1-Nrf2; ERK; Akt; JNK; Reactive Oxygen Species |
| Pathway: | NF-κB; MAPK/ERK Pathway; Stem Cell/Wnt; PI3K/Akt/mTOR; Immunology/Inflammation; Metabolic Enzyme/Protease |
| Storage: | Please store the product under the recommended conditions in the Certificate of Analysis. |



BIOLOGICAL ACTIVITY

| Description | Nrf2/HO-1 activator 1 (Compound 24) is a potent Nrf2/HO-1 activator, neuroprotective agent. Nrf2/HO-1 activator 1 shows neuroprotective and antioxidant activities. Nrf2/HO-1 activator 1 can be used in Parkinson's disease (PD) research ^[1] . | |
|--------------------------------------|---|--|
| In Vitro | Nrf2/HO-1 activator 1 (0.3-30 μM; 24 h) exhibits potent neuroprotection in both 6-OHDA- and rotenone-induced cell death models ^[1] . | |
| | Nrf2/HO-1 activator 1 (1-30 μM; 6 h) increases HO-1 expression in PC12 cells ^[1] . | |
| | Nrf2/HO-1 activator 1 (12 μM; 1-12 h) increases phosphorylation of ERK1/2, JNK, and Akt in PC12 cells ^[1] . | |
| | MCE has not independently confirmed the accuracy of these methods. They are for reference only. | |
| | Cell Viability Assay ^[1] | |
| | Cell Line: | PC12 cells |
| | Concentration: | 0.3, 1, 3, 10, and 30 μM |
| | Incubation Time: | 4 hours |
| | Result: | Inhibited 6-OHDA-induced toxicity with IC ₅₀ of 3.9 μM. Inhibited rotenone-induced toxicity with IC ₅₀ of 4.8 μM. |
| | Western Blot Analysis ^[1] | |
| Cell Line: | PC12 cells | |
| Concentration: | 10 μM | |
| Incubation Time: | 1, 3, 6, and 12 hours | |
| Result: | Induced phosphorylation of ERK1/2, JNK, and Akt. | |
| Western Blot Analysis ^[1] | | |
| Cell Line: | PC12 cells | |
| Concentration: | 1, 3, 10, and 30 μM | |
| Incubation Time: | 6 hours | |

| | | |
|----------------|--|---|
| | Result: | Increased HO-1 expression in a dose-dependent manner, with 2-fold increases at 30 μ M concentrations. |
| In Vivo | Nrf2/HO-1 activator 1 inhibits the production of lipid peroxide in rat brain homogenates by 50.5% at 100 μ M ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. | |

REFERENCES

[1]. Qili Lu, et al. Novel cudraisoiflavone J derivatives as potent neuroprotective agents for the treatment of Parkinson's disease via the activation of Nrf2/HO-1 signaling. Eur J Med Chem. 2022 Nov 15;242:114692.

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