L-796449

MedChemExpress

| Cat. No.: | HY-123654 | | Screen |
|--------------------|---|---------|---------|
| CAS No.: | 194608-80-5 | | ning |
| Molecular Formula: | $C_{28}H_{27}CIO_{4}S$ | ζ | Libr |
| Molecular Weight: | 495.03 | | raries |
| Target: | PPAR | СІСІОН | es |
| Pathway: | Cell Cycle/DNA Damage; Vitamin D Related/Nuclear Receptor | | • |
| Storage: | Please store the product under the recommended conditions in the Certificate of | | Pr |
| | Analysis. | | roteins |
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| BIOLOGICAL ACTIVITY | | | | |
|---------------------------|--|--|--|--|
| Description | L-796449 is a potent PPARγ agonist. L-796449 shows neuroprotective. L-796449 has the potential for the research of stroke ^[1] . | | | |
| IC ₅₀ & Target | PPARy ^[1] | | | |
| In Vivo | L-796449 (1 mg/kg; i.p.) shows neuroprotective in middle cerebral artery occlusion (MCAO) in the rat brain ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. | | | |
| | Animal Model: | 250 g adult male Fischer rats ^[1] | | |
| | Dosage: | 1 mg/kg (MCAO 10 minutes before IP injection) | | |
| | Administration: | l.p. | | |
| | Result: | Decreased MCAO-induced infarct size determined 2 and 7 days after the ischemic injury, inhibited MCAO-induced expression of iNOS and MMP-9 but did not affect COX-2 levels at the time examined; increased the expression of heme oxygenase-1 (HO-1) and inhibited NF-ĸB signaling. | | |

REFERENCES

[1]. Pereira MP, et al. The nonthiazolidinedione PPARgamma agonist L-796,449 is neuroprotective in experimental stroke. J Neuropathol Exp Neurol. 2005 Sep;64(9):797-805.

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

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Product Data Sheet

Inhibitors

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