MedChemExpress

## Lornoxicam-d4

| Cat. No.: | $\mathrm{HY}-\mathrm{B0} 0367 \mathrm{~S}$ |
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| CAS No.: | $1216527-48-8$ |
| Molecular Formula: | $\mathrm{C}_{13} \mathrm{H}_{6} \mathrm{D}_{4} \mathrm{ClN}_{3} \mathrm{O}_{4} \mathrm{~S}_{2}$ |
| Molecular Weight: | 375.84 |
| Target: | COX ; Endogenous Metabolite; Isotope-Labeled Compounds |
| Pathway: | Immunology/Inflammation; Metabolic Enzyme/Protease; Others |
| Storage: | Please store the product under the recommended conditions in the Certificate of |
|  | Analysis. |



## BIOLOGICAL ACTIVITY

| Description | Lornoxicam $-\mathrm{d}_{4}$ is the deuterium labeled Lornoxicam. Lornoxicam (Chlortenoxicam), a COX-1 and COX-2 inhibitor, is a new <br> nonsteroidal anti-inflammatory agent (NSAID). |
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| In Vitro | Stable heavy isotopes of hydrogen, carbon, and other elements have been incorporated into drug molecules, largely as <br> tracers for quantitation during the drug development process. Deuteration has gained attention because of its potential to <br> affect the pharmacokinetic and metabolic profiles of drugs ${ }^{[1]}$. <br> MCE has not independently confirmed the accuracy of these methods. They are for reference only. |

## REFERENCES

[1]. Russak EM, et al. Impact of Deuterium Substitution on the Pharmacokinetics of Pharmaceuticals. Ann Pharmacother. 2019;53(2):211-216
[2]. Spyra S, et al. COX-2-selective inhibitors celecoxib and deracoxib modulate transient receptor potential vanilloid 3 channels. Br J Pharmacol. 2017 Aug;174(16):26962705.
[3]. Rose, P. and C. Steinhauser, Comparison of Lornoxicam and Rofecoxib in Patients with Activated Osteoarthritis (COLOR Study). Clin Drug Investig, 2004. 24(4): p. 227-36.
[4]. Bianchi, M. and A.E. Panerai, Effects of lornoxicam, piroxicam, and meloxicam in a model of thermal hindpaw hyperalgesia induced by formalin injection in rat tail.
Pharmacol Res, 2002. 45(2): p. 101-5.

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