

Product Data Sheet

Tripamide

Cat. No.: HY-106570 **CAS No.:** 73803-48-2

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Molecular Weight: 369.87

Target: Potassium Channel

Pathway: Membrane Transporter/Ion Channel

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

BIOLOGICAL ACTIVITY

| Description | Tripamide is an orally active sulfonamide-derived diuretic antihypertensive agent $^{[1]}$. |
|-------------|--|
| In Vitro | Tripamide (less than $10 \mu\text{g/ml}$) does not modify the membrane potential and resistance, but does suppress the spike evoked by outward current pulses in the presence of 3-5 mM TEA ^[1] . In the mesenteric artery, Tripamide suppresses the amplitude of e.j.ps evoked by perivascular nerve stimulation. However, the facilitation process produced by repetitive stimulation is less affected by Tripamide ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. |
| In Vivo | In rats loaded orally with 25 ml/kg of normal saline, Tripamide (0.6-160 mg/kg) increases urine volume and sodium and chloride excretion in a dose-dependent fashion. Only at a dose of 160 mg/kg, there an increase in potassium excretion in rats [2]. Tripamide has anti-hypertensive effects, during administration to spontaneously hypertensive rats at a dose of 10 mg/kg/day for 4 weeks, tripamide doubled urine volume and sodium excretion, while potassium excretion is increased by <50% in rats ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. |

REFERENCES

[1]. H Asada, et al. Effects of N-(4-azo-endo-tricyclo[5.2.1.0.(2.6)]-decan-4-yl)-4-chloro-3-sulfamoylbenzamide (E614; tripamide) on vascular smooth muscles. Gen Pharmaco. 1982;13(3):215-23.

[2]. Philip Hampel, et al. Azosemide is more potent than bumetanide and various other loop diuretics to inhibit the sodium-potassium-chloride-cotransporter human variants hNKCC1A and hNKCC1B. Sci Rep. 2018 Jun 29;8(1):9877.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$

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