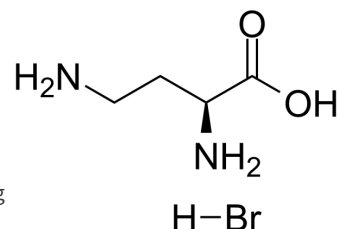


L-DABA hydrobromide

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|---------------------------|--|
| Cat. No.: | HY-101414A |
| CAS No.: | 73143-97-2 |
| Molecular Formula: | C ₄ H ₁₁ BrN ₂ O ₂ |
| Molecular Weight: | 199.05 |
| Target: | Endogenous Metabolite; GABA Receptor |
| Pathway: | Metabolic Enzyme/Protease; Membrane Transporter/Ion Channel; Neuronal Signaling |
| Storage: | 4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture) |



BIOLOGICAL ACTIVITY

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|-------------------------------------|--|
| Description | L-DABA (L-2,4-Diaminobutyric acid) hydrobromide is a weak GABA transaminase inhibitor with an IC ₅₀ of larger than 500 μM; exhibits antitumor activity in vivo and in vitro. |
| IC₅₀ & Target | IC ₅₀ : larger than 500 μM (GABA transaminase) ^[1] |
| In Vitro | The tumor cells are irreversibly and totally damaged by incubation with 10 mM L-2,4-Diaminobutyric acid for 24 h at 37°C. The cell-destructive effect by L-DABA hydrobromide is probably due to an osmotic lysis induced by the non-saturated intracellular accumulation of L-DABA hydrobromide. The harmful effect of L-DABA hydrobromide could be abolished by concomitant incubation with L-alanine and L-methionine ^[1] . Kinetic studies indicates that L-DABA hydrobromide is a non-linear, non-competitive inhibitor of GABA transaminase activity. The L-DABA hydrobromide-induced elevation of GABA levels parallels the inhibition of GABA transaminase activity ^[2] . L-2,4-Diaminobutyric acid, an amino acid analogue, produces a cytolytic effect with a human glioma cell line, SKMG-1, and normal human fibroblasts. The concentrations of L-DABA hydrobromide necessary to reduce the cell count to 50% of control following a 24-h incubation at 37°C are 12.5 mM for the human fibroblasts and 20 mM for the glioma cell line ^[3] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. |
| In Vivo | Treatment with L-DABA hydrobromide results in 43.4% reduction of tumor growth ^[1] . L-DABA hydrobromide is a more effective inhibitor of GABA transaminase in vivo than in vitro ^[2] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. |

REFERENCES

- [1]. Ronquist G, et al. Antitumor activity of L-2,4 diaminobutyric acid against mouse fibrosarcoma cells in vitro and in vivo. *J Cancer Res Clin Oncol.* 1980;96(3):259-68.
- [2]. Beart PM, et al. L-2,4-Diaminobutyric acid and the GABA system. *Neurosci Lett.* 1977 Jul;5(3-4):193-8.
- [3]. Panasci L, et al. The cytolytic effect of L-2,4 diaminobutyric acid with malignant glioma cells and fibroblasts. *Cancer Chemother Pharmacol.* 1988;21(2):143-4.

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

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