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DATASHEET

Tranlycypromine hydrochloride

Product overview

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|--------------------------|--|
| Name | Tranlycypromine hydrochloride |
| Cat No | HB1412 |
| Alternative names | trans-2-phenylcyclopropylamine; 2-PCPA; Parnate |
| Biological action | Inhibitor |
| Purity | >98% |
| Description | LSD1 / MAO inhibitor. Enables reprogramming of mouse embryonic fibroblasts into iPS cells. |

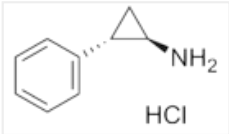
Biological Data

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|-------------------------------|--|
| Biological description | Lysine-specific demethylase 1 (LSD1) and monoamine oxidase (MAO) inhibitor ($IC_{50} = < 2 \mu M$ for LSD1). Irreversible inhibition of LSD1, inhibits H3K4 demethylation. Enables reprogramming of mouse embryonic fibroblasts into iPS cells. Shows antidepressant actions. |
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Solubility & Handling

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| Solubility overview | Soluble in water (100mM) or DMSO (100mM) |
| Storage instructions | room temperature (desiccate) |
| Storage of solutions | Prepare and use solutions on the same day if possible. Store solutions at $-20^{\circ}C$ for up to one month if storage is required. Equilibrate to RT and ensure the solution is precipitate free before use. |
| Shipping Conditions | Stable for ambient temperature shipping. Follow storage instructions on receipt. |
| Important | This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use. |

Chemical Data

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|---------------------------|---|
| Chemical name | (±)-trans-2-Phenylcyclopropylamine hydrochloride |
| Molecular Weight | 169.65 |
| Chemical structure |  <p>The chemical structure shows a cyclopropyl ring with a phenyl group attached to one carbon and an amino group (-NH₂) attached to the adjacent carbon. The stereochemistry is trans. Below the structure is the label "HCl".</p> |
| Molecular Formula | C ₉ H ₁₁ N.HCl |
| CAS Number | 1986-47-6 |
| PubChem identifier | 2723716 |
| SMILES | <chem>N[C@H]1[C@H]([C@]2=CC=CC=C2)C1.Cl</chem> |
| InChiKey | ZPEFMSTTZXJOTM-OULXEKPRSA-N |

References

trans-2-Phenylcyclopropylamine is a mechanism-based inactivator of the histone demethylase LSD1.

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Histone H3 lysine 4 demethylation is a target of nonselective antidepressive medications.

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Tranylcypromine: new perspectives on an 'old' drug.

Frieling H *et al* (2006) *Eur Arch Psychiatry Clin Neurosci* 256(5)

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