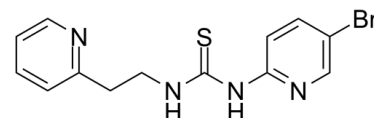


## Data Sheet

<b>Product Name:</b>	Trovirdine
<b>Cat. No.:</b>	CS-5538
<b>CAS No.:</b>	149488-17-5
<b>Molecular Formula:</b>	C <sub>13</sub> H <sub>13</sub> BrN <sub>4</sub> S
<b>Molecular Weight:</b>	337.24
<b>Target:</b>	HIV
<b>Pathway:</b>	Anti-infection
<b>Solubility:</b>	10 mM in DMSO



### BIOLOGICAL ACTIVITY:

Trovirdine inhibits HIV-1 RT with an IC<sub>50</sub> of 7 nM when employing heteropolymeric primer/template (oligo-DNA/ribosomal RNA) and dGTP as substrate. IC<sub>50</sub> value: 7 nM Target: HIV-1 Trovirdine is currently in phase I clinical trials for potential use in the treatment of AIDS.

### References:

- [1]. Zhang, H. et al. Inhibition of human immunodeficiency virus type 1 wild-type and mutant reverse transcriptases by the phenyl ethylthiazolyl thiourea derivatives trovirdine and MSC-127.
- [2]. Cantrell, A.S. et al. Phenethylthiazolylthiourea (PETT) compounds as a new class of HIV-1 reverse transcriptase inhibitors. 2. Synthesis and further structure-activity relationship studies of PETT analogs.

### CAIndexNames:

Thiourea, N-(5-bromo-2-pyridinyl)-N'-[2-(2-pyridinyl)ethyl]-

### SMILES:

S=C(NCCC1=NC=CC=C1)NC2=NC=C(Br)C=C2

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

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