

# Small Molecules

## (+)-Bay K8644

L-type calcium channel inhibitor

Catalog # 72362  
72364

1 mg  
10 mg



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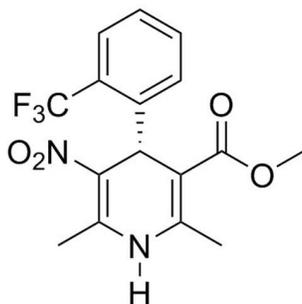
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## Product Description

Bay K8644, originally described as a modulator of potential operated calcium channels, exists as two enantiomers that have opposite actions (Hess et al.; Nowycky et al.; O'Neill et al.; Ravens et al.; Yamamoto et al.). (+)-Bay K8644 is an L-type channel blocker that has negative inotropic and vasodilatory effects (Artigas et al.; Franckowiak et al.; Ravens et al.).

Alternative Names:	(+)-Bay-K 8644; (+)-Bay-R 4407; (R)-(+)-Bay K 8644; NI 105; R 4407
CAS Number:	98791-67-4
Chemical Formula:	C <sub>16</sub> H <sub>15</sub> F <sub>3</sub> N <sub>2</sub> O <sub>4</sub>
Molecular Weight:	356.3 g/mol
Purity:	≥ 95%
Chemical Name:	R-(+)-1,4-Dihydro-2,6-dimethyl-5-nitro-4-[2-(trifluoromethyl)phenyl]-3-pyridinecarboxylic acid methyl ester
Structure:	



## Properties

Physical Appearance:	A crystalline solid
Storage:	Product stable at -20°C as supplied. Protect from prolonged exposure to light. Stable as supplied for 12 months from date of receipt.
Solubility:	· DMSO ≤ 140 mM · Absolute ethanol ≤ 140 mM For example, to prepare a 10 mM stock solution in DMSO, resuspend 1 mg in 281 µL of fresh DMSO.

Prepare stock solution fresh before use. Information regarding stability of small molecules in solution has rarely been reported; however, as a general guide we recommend storage in DMSO at -20°C. Aliquot into working volumes to avoid repeated freeze-thaw cycles. The effect of storage of stock solution on compound performance should be tested for each application.

Compound has low solubility in aqueous media. For use as a cell culture supplement, stock solution should be diluted into culture medium immediately before use. Avoid final DMSO concentration above 0.1% due to potential cell toxicity.

## Published Applications

### REPROGRAMMING

· When combined with BIX01294 (Catalog #72042), enables reprogramming of mouse embryonic fibroblasts after transduction with OCT4 and KLF4 only (Shi et al.).

### DIFFERENTIATION

· Increases neuronal differentiation from neural stem and progenitor cells (NSCs) derived from the brains of postnatal mice (D'Ascenzo et al.).

## References

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Hess P et al. (1984) Different modes of Ca channel gating behaviour favoured by dihydropyridine Ca agonists and antagonists. *Nature* 311(5986): 538–44.

Nowycky MC et al. (1985) Long-opening mode of gating of neuronal calcium channels and its promotion by the dihydropyridine calcium agonist Bay K 8644. *Proc Natl Acad Sci USA* 82(7): 2178–82.

O'Neill SK & Bolger GT. (1988) Enantiomer selectivity and the development of tolerance to the behavioral effects of the calcium channel activator BAY K 8644. *Brain Res Bull* 21(6): 865–72.

Ravens U & Schöpper H-P. (1990) Opposite cardiac actions of the enantiomers of Bay K 8644 at different membrane potentials in guinea-pig papillary muscles. *Naunyn Schmiedebergs Arch Pharmacol* 341(3).

Shi Y et al. (2008) Induction of pluripotent stem cells from mouse embryonic fibroblasts by Oct4 and Klf4 with small-molecule compounds. *Cell Stem Cell* 3(5): 568–74.

Yamamoto H et al. (1984) Bay K8644 differentiates between potential and receptor operated Ca<sup>2+</sup> channels. *Eur J Pharmacol* 102(3-4): 555–7.

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