



KB-R7943 (mesylate)

Catalog No: tcsc0848



Available Sizes

Size: 10mg

Size: 50mg



Specifications

CAS No:

182004-65-5

Formula:

 $C_{17}H_{21}N_3O_6S_2$

Pathway:

Membrane Transporter/Ion Channel

Target:

Na+/Ca2+ Exchanger

Purity / Grade:

>98%

Solubility:

DMSO : \geq 27 mg/mL (63.16 mM); H2O : 4.3 mg/mL (10.06 mM; Need warming)

Observed Molecular Weight:

427.5

Product Description

KB-R7943 mesylate is a widely used inhibitor of the reverse Na^+/Ca^{2+} exchanger (NCX_{rev}) with IC_{50} of 5.7±2.1 μ M.

IC50 & Target: IC50: $5.7\pm2.1~\mu M~(Na^+/Ca^{2+}~exchanger)^{[1]}$

In Vitro: KB-R7943 mesylate blocks NMDAR-mediated ion currents, and inhibits NMDA-induced increase in cytosolic Ca $^{2+}$ with IC $_{50}$ =13.4±3.6 μ M but accelerates calcium deregulation and mitochondrial depolarization in glutamate-treated neurons. KB-R7943



depolarizes mitochondria in a Ca²⁺-independent manner. KB-R7943 inhibits 2,4-dinitrophenol-stimulated respiration of cultured neurons with IC₅₀=11.4±2.4 μ M. In addition to NCX_{rev}, KB-R7943 dose-dependently and reversibly blocked ion currents elicited by NMDA. KB-R7943 dose-dependently inhibits NMDA-induced increases in [Ca²⁺]_c with IC₅₀=13.4±3.6 μ M confirming the inhibition of NMDA receptors observed in electrophysiological experiments^[1]. wtRyR1-HEK 293 pretreated with KB-R7943 (10 μ M, 10 min) dissolved in the bulk perfusion exhibited significantly attenuated responses to caffeine. In this regard, KB-R7943 produced more pronounced inhibition of caffeine-induced Ca²⁺ release elicited by 1 mM compared with 0.5 and 0.75 mM (60 versus 58 versus 37%, p[2]. KB-R7943 inhibits both I_{hERG} and native I_{Kr} rapidly on membrane depolarization with IC₅₀ values of ~89 and ~120 nM, respectively, for current tails at -40 mV following depolarizing voltage commands to +20 mV. I_{hERG} inhibition by KB-R7943 exhibits both time- and voltage-dependence but shows no preference for inactivated over activated channels^[3].

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