

Vernakalant (Hydrochloride)

Catalog No: tcsc0799

Available Sizes

Size: 2mg		
Size: 5mg		
Size: 10mg		
Size: 50mg		
Size: 100mg		
Specifications		
CAS No: 748810-28-8		
Formula:		

 $C_{20}H_{32}CINO_4$

Pathway: Membrane Transporter/Ion Channel

Target: Potassium Channel

Purity / Grade:

>98%

Solubility:

10 mM in DMSO

Alternative Names:

RSD1235 hydrochloride

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Observed Molecular Weight:

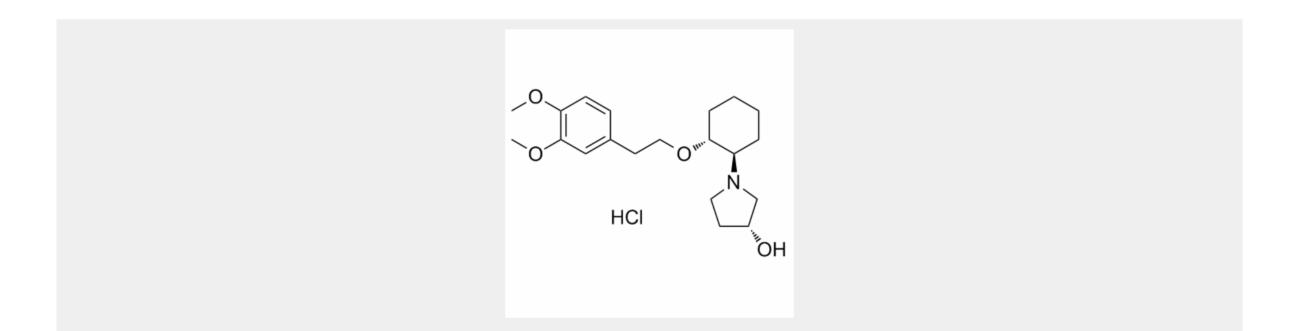
385.93

Product Description

Vernakalant hydrochloride is a mixed voltage- and frequency-dependent Na⁺ and atria-preferred K⁺ channel blocker. IC₅₀ for block by Vernakalant of wild-type and mutant Kv1.5 channels Fractional block is $13.35\pm0.93 \mu$ M, $0.61\pm0.03 \mu$ M, and $1.63\pm0.09 \mu$ M for Kv1.5 channel^{wt}, Kv1.5 channel^{I508F}, Kv1.5 channel^{T479A}, respectively.

IC50 & Target: IC50: 13.35±0.93 μ M (Kv1.5 channel^{wt}), 0.61±0.03 μ M (^{I508F}), 1.63±0.09 μ M (Kv1.5 channel ^{T479A})^[1]

In Vitro: Block of Kv1.5 by Vernakalant Hydrochloride is mediated after channel activation, because Vernakalant causes a relatively rapid onset of block of channel current upon depolarization but little evidence of resting or "tonic" block of the channel. In the presence of 10 μ M Vernakalant, rapid block is apparent after channel opening, and a steady-state current level is rapidly reached. The most important effect is the reduction in potency for Vernakalant centered at I502A, which had an IC₅₀ of 329±19 μ M (n=4-10), compared with a control IC₅₀ of 13.4±0.9 μ M (n=5-23), which is a 25-fold decrease in potency. V505A, I508A, T480A, and C500A showed lesser reductions in potency on Kv1.5, of between 3- and 4-fold. I508Y in our experiments increased the IC₅₀ for Vernakalant on Kv1.5 to 24.7 μ M, again similar to the reported value for hERG^[1].



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