

Product Data Sheet

Zolertine hydrochloride

 Cat. No.:
 HY-123368

 CAS No.:
 7241-94-3

 Molecular Formula:
 C₁₃H₁₉ClN₆

 Molecular Weight:
 294.78

Target: Adrenergic Receptor

Pathway: GPCR/G Protein; Neuronal Signaling

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

BIOLOGICAL ACTIVITY

Description	Zolertine hydrochloride is an α -adrenoceptor antagonist with a pK _i of 6.81 in rat liver (α_{1B} -adrenoceptors) and 6.35 in rabbit liver (α_{1A} -adrenoceptors) membranes ^[1] .
IC ₅₀ & Target	α1A-adrenergic receptor α1B-adrenergic receptor 6.35 (pKi, in rabbit liver membrane) 6.81 (pKi, in rat liver membrane)
In Vitro	The contractile responses induced by noradrenaline are competitively antagonized by Zolertine hydrochloride in rat carotid and aorta arteries, yielding pA $_2$ values of WKY, 7.48 \pm 0.18; SHR, 7.43 \pm 0.13 and WKY, 7.57 \pm 0.24; SHR, 7.40 \pm 0.08, respectively. Zolertine hydrochloride is a non-competitive antagonist in some blood vessels as Schild plot slopes are lower than unity. The pK $_b$ estimates for Zolertine hydrochloride are WKY, 6.98 \pm 0.16; SHR, 6.81 \pm 0.18 in the mesenteric artery, WKY, 5.73 \pm 0.11; SHR, 5.87 \pm 0.25 in the caudal artery and 6.65 \pm 0.09 in rabbit aorta $_{10}^{11}$. Zolertine hydrochloride shows higher affinity for α_{1D} -adrenoceptors compared to α_{1A} -adrenoceptors, while it had an intermediate affinity for α_{1B} -adrenoceptors $_{10}^{11}$. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Ibarra M, et al. The alpha-adrenoceptor antagonist, zolertine, inhibits alpha1D- and alpha1A-adrenoceptor-mediated vasoconstriction in vitro. J Auton Pharmacol. 2000 Jun;20(3):139-45.

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

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