

Hello Bio, Inc.  
304 Wall St., Princeton, NJ 08540 USA

T. 609-683-7500  
F. 609-228-4994

customercare-usa@hellobio.com



## DATASHEET

### Tiagabine hydrochloride

## Product overview

Name	Tiagabine hydrochloride
Cat No	HB0978
Biological action	Inhibitor
Purity	>99%
Description	Selective GAT-1 GABA uptake inhibitor

## Images



## Biological Data

Biological description	Selective GAT-1 GABA uptake inhibitor ( $IC_{50} = 67 \text{ nM}$ ). Selective for GAT-1 over mGAT2, mGAT3 and mGAT4 ( $IC_{50}$ values are 0.8, 300, >300 and 800 $\mu\text{M}$ respectively). Shows anticonvulsive, antinociceptive, anxiolytic and antidepressant actions. Blood-brain barrier permeable.
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## Solubility & Handling

Storage instructions	Room temperature (desiccate)
Solubility overview	Soluble in water (50mM) and in DMSO (100mM)
Important	This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use.

## Chemical Data

Chemical name	(3R)-1-[4,4-Bis(3-methyl-2-thienyl)-3-butenyl]-3-piperidinecarboxylic acid hydrochloride
Molecular Weight	412.01
Chemical structure	The chemical structure shows a complex molecule with a piperidine ring substituted at the 3-position with a 4-(3-methyl-2-thienyl)-3-butenoate group. The thienyl ring has a methyl group at the 3-position. The butenoate side chain is attached to the thienyl ring at the 4-position. The entire molecule is shown with its hydrochloride salt form indicated by the HCl label below the structure.
Molecular Formula	$C_{20}H_{25}NO_2S_2\cdot HCl$
CAS Number	145821-59-6
PubChem identifier	91274
SMILES	<chem>CC1=C(SC=C1)C(=CCN2CCC[C@H](C2)C(=O)O)C3=C(C=CS3)C.Cl</chem>

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InChi	InChI=1S/C20H25NO2S2.CIH/c1-14-7-11-24-18(14)17(19-15(2)8-12-25-19)6-4-10-21-9-3-5-16(13-21)20(22)23;h6-8,11-12,16H,3-5,9-10,13H2,1-2H3,(H,22,23);1H/t16-;/m1./s1
InChiKey	YUKARLAABCGMCN-PKLMIRHRSA-N
MDL number	MFCD07369025

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## References

**Long-term anxiolytic and antidepressant-like behavioural effects of tiagabine, a selective GABA transporter-1 (GAT-1) inhibitor, coincide with a decrease in HPA system activity in C57BL/6 mice.**

Thoeringer CK *et al* (2010) J Psychopharmacol 24(5)

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**Correlation between anticonvulsant activity and inhibitory action on glial gamma-aminobutyric acid uptake of the highly selective mouse gamma-aminobutyric acid transporter 1 inhibitor 3-hydroxy-4-amino-4,5,6,7-tetrahydro-1,2-benzisoxazole and its N-alkyls**

White HS *et al* (2002) J Pharmacol Exp Ther 302(2)

PubMedID [12130726](#)

**Comparison of antiepileptic drugs tiagabine, lamotrigine, and gabapentin in mouse models of acute, prolonged, and chronic nociception.**

Laughlin TM *et al* (2002) J Pharmacol Exp Ther 302(3)

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