

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

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

customercare-usa@hellobio.com



DATASHEET

Tamoxifen Citrate

Product overview

Name	Tamoxifen Citrate
Cat No	HB0602
Alternative names	TAM
Biological action	Antagonist
Purity	>98%
Description	Estrogen receptor antagonist/ partial agonist. May be used in genome engineering (e.g. CreER/CRISPR-Cas9). Citrate salt.

Images



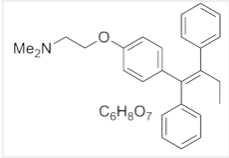
Biological Data

Biological description	Citrate salt of Tamoxifen, which is an estrogen receptor antagonist and partial agonist. It is also a potent chloride channel HSV-1 inhibitor and is blood brain barrier permeable. Tamoxifen inhibits tumor growth and induces apoptosis in breast cancer cells. It also selectively inhibits sterol biosynthesis ($IC_{50} = 1000 \text{ nM}$) and shows neuroprotective potential. It is used in gene editing in which tamoxifen (Tm)-inducible Cre recombinases are widely used to perform gene inactivation and lineage tracing studies in mice. It can also switch on the inducible CRISPR-Cas9 system (iCas).
-------------------------------	---

Solubility & Handling

Solubility overview	Soluble in Dimethylsulfoxide (100mM), and in ethanol (5 mM with warming)
Storage instructions	+4 °C
Storage of solutions	Prepare and use solutions on the same day if possible. Store solutions at -20 °C for up to one month if storage is required. Equilibrate to RT and ensure the solution is precipitate free before use.
Shipping Conditions	Stable for ambient temperature shipping. Follow storage instructions on receipt.
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	(Z)-2-[4-(1,2-Diphenyl-1-butenyl)phenoxy]-N,N-dimethylethanamine citrate
Molecular Weight	563.7
Chemical structure	
Molecular Formula	C ₃₂ H ₃₇ NO ₈
CAS Number	54965-24-1
PubChem identifier	2733525
SMILES	<chem>CCC(=C(C1=CC=CC=C1)C2=CC=C(C(C=C2)OCCN(C)C)C3=CC=CC=C3.C(C(=O)O)C(CC(=O)O)(C(=O)O)O</chem>
InChi	InChI=1S/C26H29NO.C6H8O7/c1-4-25(21-11-7-5-8-12-21)26(22-13-9-6-10-14-22)23-15-17-24(18-16-23)28-20-19-27(2)3;7-3(8)1-6(13,5(11)12)2-4(9)10/h5-18H,4,19-20H2,1-3H3;13H,1-2H2,(H,7,8)(H,9,10)(H,11,12)/b26-25-;
InChiKey	FQZTYTYWMLGAPFJ-OQKDUQJOSA-N
MDL number	MFCD00058321
Appearance	White solid

References

Tamoxifen inhibits inward rectifier K⁺ 2.x family of inward rectifier channels by interfering with phosphatidylinositol 4,5-bisphosphate-channel interactions.

Ponce-Balbuena D *et al* (2009) J Pharmacol Exp Ther 331(2)

PubMedID [19654266](#)

Both the immunosuppressant SR31747 and the antiestrogen tamoxifen bind to an emopamil-insensitive site of mammalian Delta8-Delta7 sterol isomerase.

Paul R *et al* (1998) J Pharmacol Exp Ther 285(3)

PubMedID [9618436](#)

Growth inhibition of estrogen receptor-positive and aromatase-positive human breast cancer cells in monolayer and spheroid cultures by letrozole, anastrozole, and tamoxifen.

Kijima I *et al* (2005) J Steroid Biochem Mol Biol 97(4)

PubMedID [16263272](#)

Inhibition of herpes simplex virus type 1 entry by chloride channel inhibitors tamoxifen and NPPB.

Zheng K *et al* (2014) Biochem Biophys Res Commun 446(4)

PubMedID [24657267](#)

A Chemical-Inducible CRISPR-Cas9 System for Rapid Control of Genome Editing

Liu et al (2016) Nat Chem Biol. 12(11)

PubMedID [27618190](#)