PRODUCT INFORMATION



Tamoxifen (citrate)

Item No. 11629

CAS Registry No.:	54965-24-1		
Formal Name:	2-[4-[(1Z)-1,2-diphenyl-1-buten-1-yl]		
	phenoxy]-N,N-dimethyl-ethanamine,		
	2-hydroxy-1,2,3-propanetricarboxylate		
Synonyms:	ICI 46474, TMX	\uparrow	ОН
MF:	$C_{26}H_{29}NO \bullet C_{6}H_{8}O_{7}$		• HOOC
FW:	563.6		
Purity:	≥98%		COOH
UV/Vis.:	λ _{max} : 237, 276 nm		
Supplied as:	A crystalline solid		
Storage:	-20°C		
Stability:	≥4 years		

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Laboratory Procedures

Tamoxifen (citrate) is supplied as a crystalline solid. A stock solution may be made by dissolving the Tamoxifen (citrate) in the solvent of choice, which should be purged with an inert gas. Tamoxifen (citrate) is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide. The solubility of Tamoxifen (citrate) in these solvents is approximately 10, 2, and 20 mg/ml, respectively.

Tamoxifen (citrate) is sparingly soluble in aqueous buffers. For maximum solubility in aqueous buffers, Tamoxifen (citrate) should first be dissolved in ethanol and then diluted with the aqueous buffer of choice. Tamoxifen (citrate) has a solubility of approximately 0.3 mg/ml in a 1:2 solution of ethanol:PBS (pH 7.2) using this method. We do not recommend storing the aqueous solution for more than one day.

Description

Tamoxifen is an estrogen receptor antagonist (IC₅₀ = 45 nM for the rabbit receptor).¹ It reduces the proliferation of MCF-7 breast cancer cells when used at a concentration of 10 µM.² Tamoxifen is active against S. cerevisiae, C. neoformans, and five Candida species (MICs = 12, 64, and 8-64 μ g/ml, respectively).³ It decreases tumor proliferation, weight, and volume in an MCF-7 mouse xenograft model when administered at a dose of 100 µg/animal per day.⁴ Tamoxifen (0.8 mg/kg every two weeks) also reduces proteinuria and increases survival in an NZBWF1 mouse model of systemic lupus erythematosus (SLE).⁵ It can be used as a regulator for Cre-recombinase inducible gene expression in vivo.⁶ Formulations containing tamoxifen have been used in the treatment of estrogen receptor-positive breast cancer.

References

- 1. Löser, R., Seibel, K., Roos, W., et al. Eur. J. Cancer Clin. Oncol. 21(8), 985-990 (1985).
- 2. Lippman, M., Bolan, G., and Huff, K. Cancer Res. 36(12), 4595-4601 (1976).
- 3. Dolan, K., Montgomery, S., Buchheit, B., et al. Antimicrob. Agents Chemother. 53(8), 3337-3346 (2009).
- 4. Long, B.J., Jelovac, D., Handratta, V., et al. J. Natl. Cancer Inst. 96(6), 456-465 (2004).
- 5. Wu, W.-M., Lin, B.-F., Su, Y.-C., et al. Scand. J. Immunol. 52(4), 393-400 (2000).
- 6. Chen, M., Lichtler, A.C., Sheu, T.-J., et al. Genesis 45(1), 44-50 (2007).

WARNING THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFFTY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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