# PRODUCT INFORMATION



# Tamoxifen-d<sub>5</sub> Item No. 18262

CAS Registry No.: 157698-32-3

Formal Name: 2-[4-[(1Z)-1,2-diphenyl-1-buten-1-yl-3,3,4,4,4-d<sub>5</sub>]

phenoxy]-N,N-dimethyl-ethanamine

MF:  $C_{26}H_{24}D_5NO$ FW: 376.6

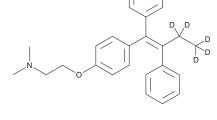
≥98% (Tamoxifen) **Chemical Purity:** 

Deuterium

Incorporation:  $\geq$ 99% deuterated forms (d<sub>1</sub>-d<sub>5</sub>);  $\leq$ 1% d<sub>0</sub>

Supplied as: A 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-d<sub>5</sub> is intended for use as an internal standard for the quantification of tamoxifen (Item No. 13258) by GC- or LC-MS. The accuracy of the sample weight in this vial is between 5% over and 2% under the amount shown on the vial. If better precision is required, the deuterated standard should be quantitated against a more precisely weighed unlabeled standard by constructing a standard curve of peak intensity ratios (deuterated versus unlabeled).

Tamoxifen- $d_{\epsilon}$  is supplied as a solid. A stock solution may be made by dissolving the tamoxifen- $d_{\epsilon}$  in the solvent of choice, which should be purged with an inert gas. Tamoxifen- $d_s$  is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of tamoxifen-d<sub>5</sub> in ethanol and DMF is approximately 20 mg/ml and approximately 2 mg/ml in DMSO.

#### Description

Tamoxifen-d<sub>5</sub> is intended for use as an internal standard for the quantification of tamoxifen (Item No. 13258) by GC- or LC-MS. Tamoxifen is an estrogen receptor antagonist (IC<sub>50</sub> = 45 nM for the rabbit receptor).<sup>1</sup> It reduces the proliferation of MCF-7 breast cancer cells when used at a concentration of 10 μM.<sup>2</sup> Tamoxifen is active against S. cerevisiae, C. neoformans, and five Candida species (MICs = 12, 64, and 8-64 μg/ml, respectively).<sup>3</sup> 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.<sup>4</sup> Tamoxifen (0.8 mg/kg every two weeks) also reduces proteinuria and increases survival in an NZBWF1 mouse model of systemic lupus erythematosus (SLE).<sup>5</sup> It can be used as a regulator for Cre-recombinase inducible gene expression in vivo. 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).
- Wu, W.-M., Lin, B.-F., Su, Y.-C., et al. Scand. J. Immunol. 52(4), 393-400 (2000).
- 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.

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.

## WARRANTY AND LIMITATION OF REMEDY

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