

L-Cystine, from non-animal source Cell culture tested, meets EP testing specifications

Product Number **C7602**
Store at Room Temperature

Product Description

Molecular Formula: $C_6H_{12}N_2O_4S_2$
Molecular Weight: 240.3
CAS Number: 56-89-3
Synonyms: [*R*-(*R**, *R**)]-3,3'-dithiobis[2-aminopropanoic acid], dicysteine, β , β' -dithiodialanine¹

This product is tested for endotoxin levels and suitability for cell culture experiments.

Cystine is a derived amino acid that is formed from the oxidative linkage of two cysteine residues to give a disulfide covalent bond. Cystines form in many proteins after incorporation of free cysteines into the primary structure to stabilize their folded conformation. Cystine is the form in which cysteine exists in blood and urine.²

The two cystine-related clinical conditions are cystinuria, which involves the defective membrane transport of cystine, and cystinosis, the accumulation of cystine in lysosomes.^{2,3,4} A review of cystine transport into rat brain cells has been published.⁵ An investigation into cysteine and cystine levels in normal and malignant cells with a relationship to γ -cystathionase levels and tumor sensitivity to L-cysteine and cystine depletion has been reported.⁶

Mass spectrometry (GC-MS) methods for the analysis of cystine from granulocytes of cystinosis patients⁷ and in the urine of homocystinuria patients⁸ have been published.

Precautions and Disclaimer

For Laboratory Use Only. Not for drug, household or other uses.

Preparation Instructions

This product is soluble in 1 M HCl (100 mg/ml), with heat as needed. The solubility of cystine in water is 0.112 mg/ml at 25 °C; cystine is more soluble in aqueous solutions with pH < 2 or pH > 8.¹

References

1. The Merck Index, 12th ed., Entry# 2851.
2. Textbook of Biochemistry with Clinical Correlations, Devlin, T. M., ed., Wiley-Liss (New York, NY: 1992), pp. 33, 503.
3. Palacin, M., et al., The molecular bases of cystinuria and lysinuric protein intolerance. *Curr. Opin. Genet. Dev.*, **11(3)**, 328-335 (2001).
4. Gahl, W. A., et al., Cystinosis. *N. Engl. J. Med.*, **347(2)**, 111-121 (2002).
5. McBean, G. J., and Flynn, J., Molecular mechanisms of cystine transport. *Biochem. Soc. Trans.*, **29(Pt 6)**, 717-722 (2001).
6. Uren, J. R., and Lazarus, H., L-cyst(e)ine requirements of malignant cells and progress toward depletion therapy. *Cancer Treat. Rep.*, **63(6)**, 1073-1079 (1979).
7. Schierbeek, H., et al., Stable isotope dilution analysis of cystine in granulocyte suspensions as cysteine: a powerful method for the diagnosis, the follow-up, and treatment of patients with cystinosis. *Clin. Chim. Acta*, **191(1-2)**, 39-47 (1990).
8. Kuhara, T., et al., Differential diagnosis of homocystinuria by urease treatment, isotope dilution and gas chromatography-mass spectrometry. *J. Chromatogr. B Biomed. Sci. Appl.*, **742(1)**, 59-70 (2000).

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